

SYNTHESIS OF HALICHONDRINS

RELATED APPLICATIONS

[0001] This application claims priority under 35 U.S.C. § 119(e) to U.S. provisional patent applications, U.S.S.N. 62/529,333, filed July 6, 2017; and U.S.S.N. 62/529,310, filed July 6, 2017; the entire contents of each of which is incorporated herein by reference.

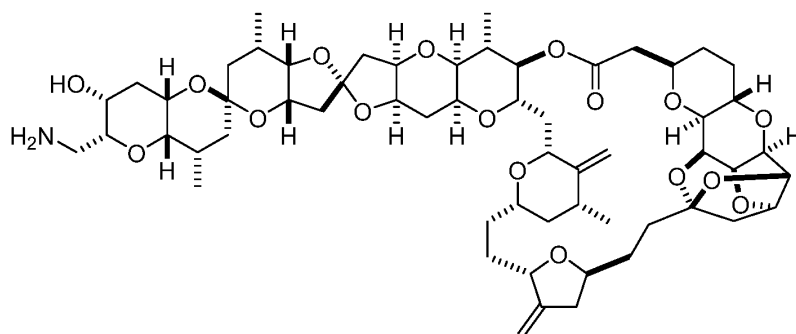
BACKGROUND OF THE INVENTION

[0002] Halichondrins are polyether natural products, originally isolated from the marine scavenger *Halichondria okadai* by Uemura, Hirata, and coworkers. See, *e.g.*, Uemura, D.; Takahashi, K.; Yamamoto, T.; Katayama, C.; Tanaka, J.; Okumura, Y.; Hirata, Y. *J. Am. Chem. Soc.* **1985**, *107*, 4796; Hirata, Y.; Uemura, D. *Pure Appl. Chem.* **1986**, *58*, 701. Several additional members, including halistatin, were isolated from various marine scavengers. This class of natural products displays interesting structural diversity, such as the oxidation state of the carbons of the C8-C14 polycycle, and the length of the carbon backbone. Thus, this class of natural products is sub-grouped into the norhalichondrin series (*e.g.*, norhalichondrin A, B, and C), the halichondrin series (*e.g.*, halichondrin A, B, C), and the homohalichondrin series (*e.g.*, homohalichondrin A, B, C) (see *Figure 1*). Except halichondrin A, all the members have been isolated from natural sources. Due to their intriguing structural architecture and extraordinary antitumor activity, halichondrins have received much attention from the scientific community.

SUMMARY OF THE INVENTION

[0003] The present invention provides new synthetic methods useful in the synthesis of halichondrin natural products and related molecules. As described herein, a novel nickel/zirconium-mediated coupling reaction has been developed as a key step in the synthesis. In addition to synthetic methods, the present invention also provides compounds which are useful synthetic intermediates in the synthesis of halichondrin natural products and analogs thereof.

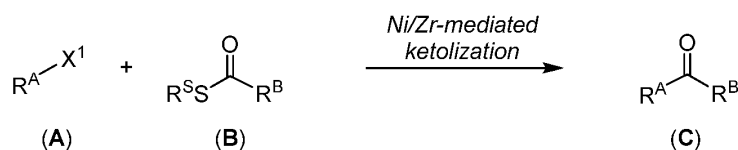
[0004] For example, in certain embodiments, provided herein are compounds and methods useful in the synthesis of Compound (1):



Compound (1).

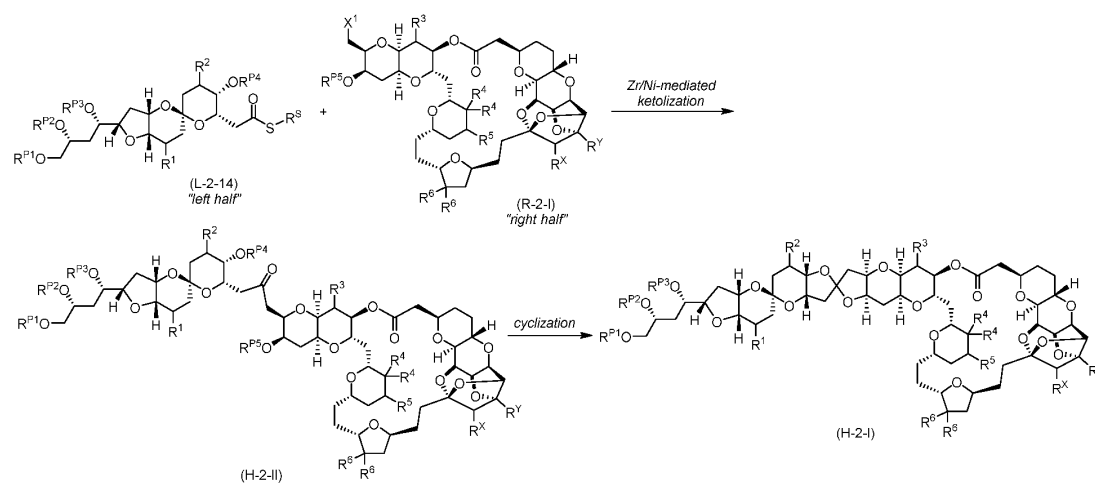
[0005] In one aspect, the present invention provides methods for preparing ketones using a Ni/Zr-mediated coupling reaction, as outlined in *Scheme 1A*. These coupling reactions can be applied to the synthesis of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C; norhalichondrin A, B, C), and analogs thereof.

Scheme 1A



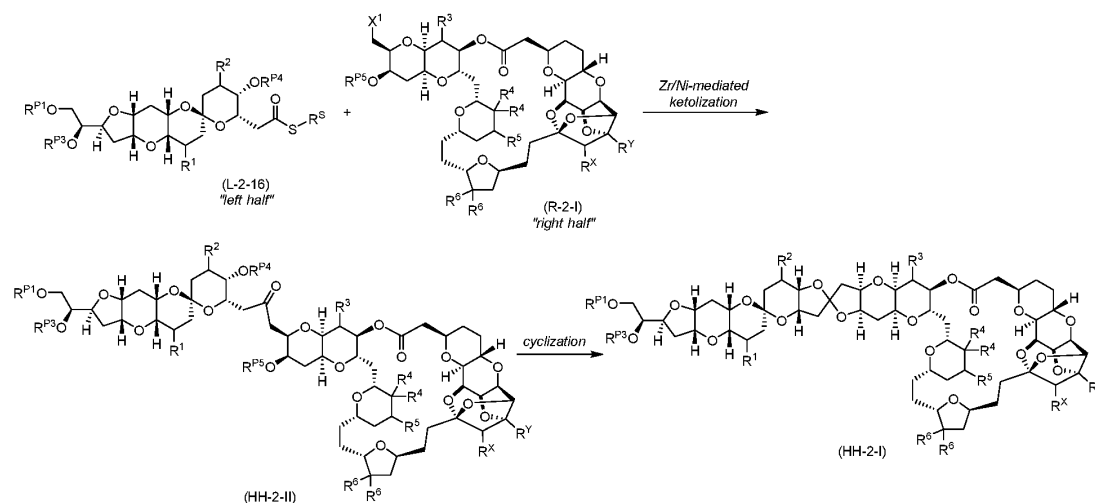
[0006] Application of Ni/Zr-mediated coupling reactions provided herein to the preparation of compounds in the halichondrin series (*e.g.*, halichondrin A, B, C, and analogs thereof) is outlined in *Scheme 2A*, for example. This strategy involves a coupling of a “left half” building block with a “right half” building block via a Ni/Zr-mediated ketolization reaction described herein.

Scheme 2A



[0007] Application of Ni/Zr-mediated coupling reactions provided herein to the preparation of compounds in the homohalichondrin series (*e.g.*, homohalichondrin A, B, C, and analogs thereof) is outlined in *Scheme 2B*, for example. This strategy involves a coupling of a "left half" building block with a "right half" building block via a Ni/Zr-mediated ketolization reaction described herein.

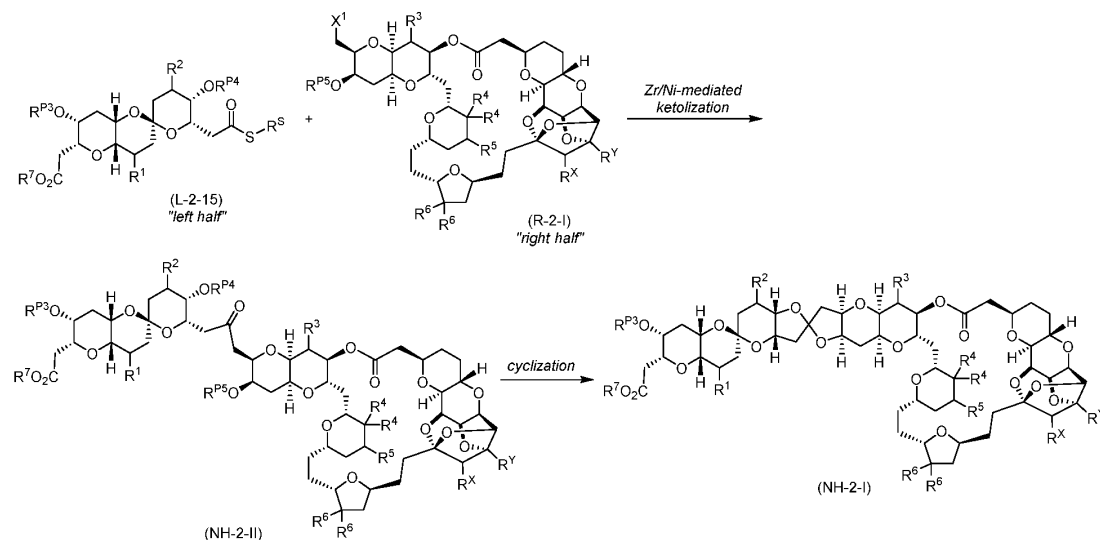
Scheme 2B



[0008] Application of Ni/Zr-mediated coupling reactions provided herein to the preparation of compounds in the norhalichondrin series (*e.g.*, norhalichondrin A, B, C, and analogs thereof) is outlined in *Scheme 2C*, for example. This strategy involves coupling of a "left

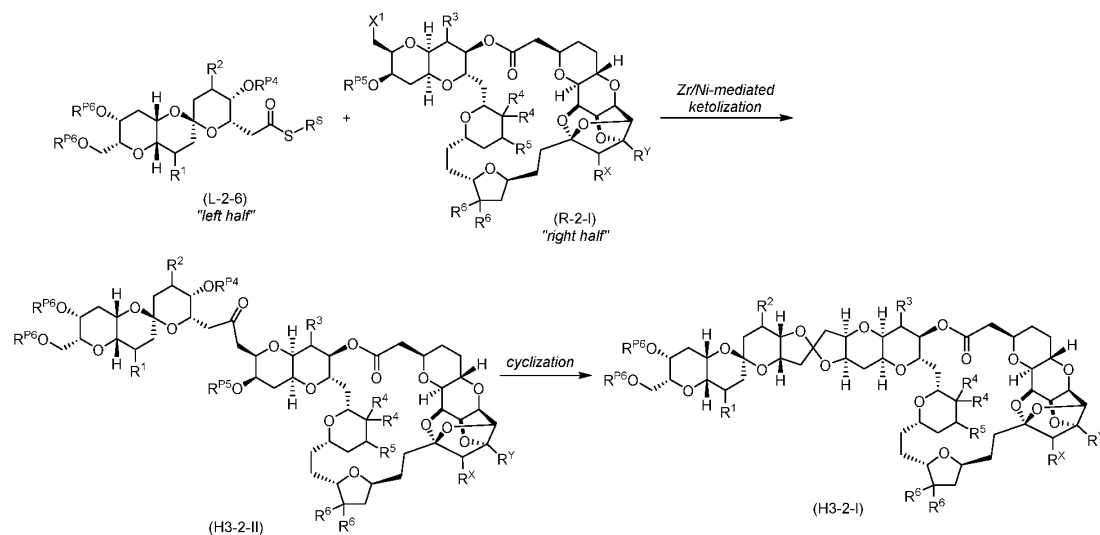
half” building block with a “right half” building block via a Ni/Zr-mediated ketolization reaction described herein.

Scheme 2C



[0009] Application of Ni/Zr-mediated coupling reactions provided herein to the preparation of additional halichondrin analogs is outlined in *Scheme 2D*, for example. This strategy involves a coupling of a “left half” building block with a “right half” building block via a Ni/Zr-mediated ketolization reaction described herein.

Scheme 2D



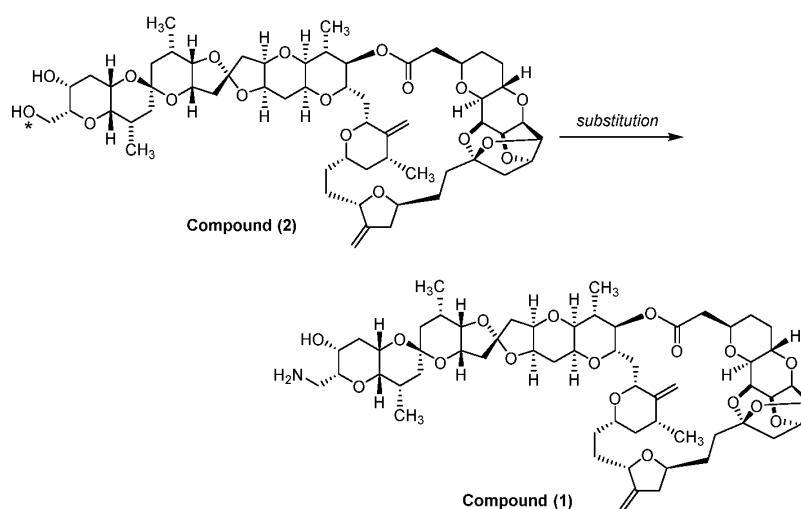
[0010] In general, the provided methods for the preparation of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C; norhalichondrin A, B, C), and analogs thereof, involve the coupling of a “left half” fragment with a “right half” fragment. In another aspect, the present invention provides methods useful in the preparation of said “right half” and “left half” building blocks.

[0011] In another aspect, the present invention provides compounds which are useful intermediates *en route* to halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C; norhalichondrin A, B, C), and analogs thereof. For example, in one aspect, the present invention provides novel “left half” and “right half” building blocks of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C; norhalichondrin A, B, C), and analogs thereof, and intermediates useful in the preparation of said building blocks.

[0012] In yet another aspect, the present invention provides methods useful in the preparation of halichondrin analogs; in particular, the preparation of Compound (1). The present invention also provides compounds (*i.e.*, synthetic intermediates) useful in the synthesis of Compound (1).

[0013] In one aspect, the present invention provides methods for preparing Compound (1) that involve substituting the primary hydroxyl group of Compound (2) ($-\text{OH}$; denoted by * in *Scheme 1*) with an amino group ($-\text{NH}_2$). The substitution may be carried out in one or more steps. For example, the substitution may be carried out by converting the primary hydroxyl group of Compound (2) to a leaving group (*e.g.*, $-\text{OR}^1$), followed by substitution of the leaving group with an amine or amine precursor (*e.g.*, azide).

Scheme 1



[0014] Current methods for the synthesis of halichondrins can be found, for example, in international PCT publications, WO 2016/176560, published November 3, 2016, and WO 2016/003975, published January 7, 2016; the entire contents of each of which is incorporated herein by reference.

[0015] Other current methods for the synthesis of halichondrins can be found, for example, in U.S. Patent No. 9,938,288, issued April 10, 2018; U.S. Provisional Patent Application, U.S.S.N. 62/586,416, filed November 15, 2017; International Application No. PCT/US2018/031765, filed May 9, 2018; U.S. Patent Application Publication No. US 2018/0155361, published June 7, 2018; the entire contents of each of which is incorporated herein by reference.

[0016] The details of certain embodiments of the invention are set forth in the Detailed Description of Certain Embodiments, as described below. Other features, objects, and advantages of the invention will be apparent from the Definitions, Examples, Figures, and Claims.

DEFINITIONS

[0017] Definitions of specific functional groups and chemical terms are described in more detail below. The chemical elements are identified in accordance with the Periodic Table of the Elements, CAS version, *Handbook of Chemistry and Physics*, 75th Ed., inside cover, and specific functional groups are generally defined as described therein. Additionally, general principles of organic chemistry, as well as specific functional moieties and reactivity, are described in *Organic Chemistry*, Thomas Sorrell, University Science Books, Sausalito, 1999; Smith and March, *March's Advanced Organic Chemistry*, 5th Edition, John Wiley & Sons, Inc., New York, 2001; Larock, *Comprehensive Organic Transformations*, VCH Publishers, Inc., New York, 1989; and Carruthers, *Some Modern Methods of Organic Synthesis*, 3rd Edition, Cambridge University Press, Cambridge, 1987.

[0018] Compounds described herein can comprise one or more asymmetric centers, and thus can exist in various stereoisomeric forms, *e.g.*, enantiomers and/or diastereomers. For example, the compounds described herein can be in the form of an individual enantiomer, diastereomer or geometric isomer, or can be in the form of a mixture of stereoisomers, including racemic mixtures and mixtures enriched in one or more stereoisomer. Isomers can be isolated from mixtures by methods known to those skilled in the art, including chiral high pressure liquid chromatography (HPLC) and the formation and crystallization of chiral salts; or preferred isomers can be prepared by asymmetric syntheses. See, for example, Jacques *et*

al., *Enantiomers, Racemates and Resolutions* (Wiley Interscience, New York, 1981); Wilen *et al.*, *Tetrahedron* 33:2725 (1977); Eliel, E.L. *Stereochemistry of Carbon Compounds* (McGraw-Hill, NY, 1962); and Wilen, S.H., *Tables of Resolving Agents and Optical Resolutions* p. 268 (E.L. Eliel, Ed., Univ. of Notre Dame Press, Notre Dame, IN 1972). The invention additionally encompasses compounds as individual isomers substantially free of other isomers, and alternatively, as mixtures of various isomers.

[0019] Unless otherwise stated, structures depicted herein are also meant to include compounds that differ only in the presence of one or more isotopically enriched atoms. For example, compounds having the present structures except for the replacement of hydrogen by deuterium or tritium, replacement of ^{19}F with ^{18}F , or the replacement of ^{12}C with ^{13}C or ^{14}C are within the scope of the disclosure. Such compounds are useful, for example, as analytical tools or probes in biological assays.

[0020] When a range of values is listed, it is intended to encompass each value and sub-range within the range. For example “C₁₋₆ alkyl” is intended to encompass, C₁, C₂, C₃, C₄, C₅, C₆, C₁₋₆, C₁₋₅, C₁₋₄, C₁₋₃, C₁₋₂, C₂₋₆, C₂₋₅, C₂₋₄, C₂₋₃, C₃₋₆, C₃₋₅, C₃₋₄, C₄₋₆, C₄₋₅, and C₅₋₆ alkyl.

[0021] The term “aliphatic” refers to alkyl, alkenyl, alkynyl, and carbocyclic groups. Likewise, the term “heteroaliphatic” refers to heteroalkyl, heteroalkenyl, heteroalkynyl, and heterocyclic groups.

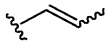
[0022] The term “alkyl” refers to a radical of a straight-chain or branched saturated hydrocarbon group having from 1 to 10 carbon atoms (“C₁₋₁₀ alkyl”). In some embodiments, an alkyl group has 1 to 9 carbon atoms (“C₁₋₉ alkyl”). In some embodiments, an alkyl group has 1 to 8 carbon atoms (“C₁₋₈ alkyl”). In some embodiments, an alkyl group has 1 to 7 carbon atoms (“C₁₋₇ alkyl”). In some embodiments, an alkyl group has 1 to 6 carbon atoms (“C₁₋₆ alkyl”). In some embodiments, an alkyl group has 1 to 5 carbon atoms (“C₁₋₅ alkyl”). In some embodiments, an alkyl group has 1 to 4 carbon atoms (“C₁₋₄ alkyl”). In some embodiments, an alkyl group has 1 to 3 carbon atoms (“C₁₋₃ alkyl”). In some embodiments, an alkyl group has 1 to 2 carbon atoms (“C₁₋₂ alkyl”). In some embodiments, an alkyl group has 1 carbon atom (“C₁ alkyl”). In some embodiments, an alkyl group has 2 to 6 carbon atoms (“C₂₋₆ alkyl”). Examples of C₁₋₆ alkyl groups include methyl (C₁), ethyl (C₂), propyl (C₃) (*e.g.*, n-propyl, isopropyl), butyl (C₄) (*e.g.*, n-butyl, tert-butyl, sec-butyl, iso-butyl), pentyl (C₅) (*e.g.*, n-pentyl, 3-pentanyl, amyl, neopentyl, 3-methyl-2-butanyl, tertiary amyl), and hexyl (C₆) (*e.g.*, n-hexyl). Additional examples of alkyl groups include n-heptyl (C₇), n-octyl (C₈), and the like. Unless otherwise specified, each instance of an alkyl group is independently unsubstituted (an “unsubstituted alkyl”) or substituted (a “substituted alkyl”)

with one or more substituents (*e.g.*, halogen, such as F). In certain embodiments, the alkyl group is an unsubstituted C₁₋₁₀ alkyl (such as unsubstituted C₁₋₆ alkyl, *e.g.*, –CH₃ (Me), unsubstituted ethyl (Et), unsubstituted propyl (Pr, *e.g.*, unsubstituted n-propyl (n-Pr), unsubstituted isopropyl (i-Pr)), unsubstituted butyl (Bu, *e.g.*, unsubstituted n-butyl (n-Bu), unsubstituted tert-butyl (tert-Bu or t-Bu), unsubstituted sec-butyl (sec-Bu), unsubstituted isobutyl (i-Bu)). In certain embodiments, the alkyl group is a substituted C₁₋₁₀ alkyl (such as substituted C₁₋₆ alkyl, *e.g.*, –CF₃, Bn).

[0023] The term “haloalkyl” is a substituted alkyl group, wherein one or more of the hydrogen atoms are independently replaced by a halogen, *e.g.*, fluoro, bromo, chloro, or iodo. In some embodiments, the haloalkyl moiety has 1 to 8 carbon atoms (“C₁₋₈ haloalkyl”). In some embodiments, the haloalkyl moiety has 1 to 6 carbon atoms (“C₁₋₆ haloalkyl”). In some embodiments, the haloalkyl moiety has 1 to 4 carbon atoms (“C₁₋₄ haloalkyl”). In some embodiments, the haloalkyl moiety has 1 to 3 carbon atoms (“C₁₋₃ haloalkyl”). In some embodiments, the haloalkyl moiety has 1 to 2 carbon atoms (“C₁₋₂ haloalkyl”). Examples of haloalkyl groups include –CHF₂, –CH₂F, –CF₃, –CH₂CF₃, –CF₂CF₃, –CF₂CF₂CF₃, –CCl₃, –CFCl₂, –CF₂Cl, and the like.

[0024] The term “heteroalkyl” refers to an alkyl group, which further includes at least one heteroatom (*e.g.*, 1, 2, 3, or 4 heteroatoms) selected from oxygen, nitrogen, or sulfur within (*i.e.*, inserted between adjacent carbon atoms of) and/or placed at one or more terminal position(s) of the parent chain. In certain embodiments, a heteroalkyl group refers to a saturated group having from 1 to 10 carbon atoms and 1 or more heteroatoms within the parent chain (“heteroC₁₋₁₀ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 9 carbon atoms and 1 or more heteroatoms within the parent chain (“heteroC₁₋₉ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 8 carbon atoms and 1 or more heteroatoms within the parent chain (“heteroC₁₋₈ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 7 carbon atoms and 1 or more heteroatoms within the parent chain (“heteroC₁₋₇ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 6 carbon atoms and 1 or more heteroatoms within the parent chain (“heteroC₁₋₆ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 5 carbon atoms and 1 or 2 heteroatoms within the parent chain (“heteroC₁₋₅ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 4 carbon atoms and 1 or 2 heteroatoms within the parent chain (“heteroC₁₋₄ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 to 3 carbon atoms and 1 heteroatom within the parent chain (“heteroC₁₋₃ alkyl”). In some embodiments, a

heteroalkyl group is a saturated group having 1 to 2 carbon atoms and 1 heteroatom within the parent chain (“heteroC₁₋₂ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 1 carbon atom and 1 heteroatom (“heteroC₁ alkyl”). In some embodiments, a heteroalkyl group is a saturated group having 2 to 6 carbon atoms and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₆ alkyl”). Unless otherwise specified, each instance of a heteroalkyl group is independently unsubstituted (an “unsubstituted heteroalkyl”) or substituted (a “substituted heteroalkyl”) with one or more substituents. In certain embodiments, the heteroalkyl group is an unsubstituted heteroC₁₋₁₀ alkyl. In certain embodiments, the heteroalkyl group is a substituted heteroC₁₋₁₀ alkyl.

[0025] The term “alkenyl” refers to a radical of a straight-chain or branched hydrocarbon group having from 2 to 10 carbon atoms and one or more carbon-carbon double bonds (*e.g.*, 1, 2, 3, or 4 double bonds). In some embodiments, an alkenyl group has 2 to 9 carbon atoms (“C₂₋₉ alkenyl”). In some embodiments, an alkenyl group has 2 to 8 carbon atoms (“C₂₋₈ alkenyl”). In some embodiments, an alkenyl group has 2 to 7 carbon atoms (“C₂₋₇ alkenyl”). In some embodiments, an alkenyl group has 2 to 6 carbon atoms (“C₂₋₆ alkenyl”). In some embodiments, an alkenyl group has 2 to 5 carbon atoms (“C₂₋₅ alkenyl”). In some embodiments, an alkenyl group has 2 to 4 carbon atoms (“C₂₋₄ alkenyl”). In some embodiments, an alkenyl group has 2 to 3 carbon atoms (“C₂₋₃ alkenyl”). In some embodiments, an alkenyl group has 2 carbon atoms (“C₂ alkenyl”). The one or more carbon-carbon double bonds can be internal (such as in 2-butenyl) or terminal (such as in 1-butenyl). Examples of C₂₋₄ alkenyl groups include ethenyl (C₂), 1-propenyl (C₃), 2-propenyl (C₃), 1-butenyl (C₄), 2-butenyl (C₄), butadienyl (C₄), and the like. Examples of C₂₋₆ alkenyl groups include the aforementioned C₂₋₄ alkenyl groups as well as pentenyl (C₅), pentadienyl (C₅), hexenyl (C₆), and the like. Additional examples of alkenyl include heptenyl (C₇), octenyl (C₈), octatrienyl (C₈), and the like. Unless otherwise specified, each instance of an alkenyl group is independently unsubstituted (an “unsubstituted alkenyl”) or substituted (a “substituted alkenyl”) with one or more substituents. In certain embodiments, the alkenyl group is an unsubstituted C₂₋₁₀ alkenyl. In certain embodiments, the alkenyl group is a substituted C₂₋₁₀ alkenyl. In an alkenyl group, a C=C double bond for which the stereochemistry is not specified (*e.g.*, $-\text{CH}=\text{CHCH}_3$ or ) may be an (*E*)- or (*Z*)-double bond.

[0026] The term “heteroalkenyl” refers to an alkenyl group, which further includes at least one heteroatom (*e.g.*, 1, 2, 3, or 4 heteroatoms) selected from oxygen, nitrogen, or sulfur

within (*i.e.*, inserted between adjacent carbon atoms of) and/or placed at one or more terminal position(s) of the parent chain. In certain embodiments, a heteroalkenyl group refers to a group having from 2 to 10 carbon atoms, at least one double bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₁₀ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 9 carbon atoms at least one double bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₉ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 8 carbon atoms, at least one double bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₈ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 7 carbon atoms, at least one double bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₇ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 6 carbon atoms, at least one double bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₆ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 5 carbon atoms, at least one double bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₅ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 4 carbon atoms, at least one double bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₄ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 3 carbon atoms, at least one double bond, and 1 heteroatom within the parent chain (“heteroC₂₋₃ alkenyl”). In some embodiments, a heteroalkenyl group has 2 to 6 carbon atoms, at least one double bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₆ alkenyl”). Unless otherwise specified, each instance of a heteroalkenyl group is independently unsubstituted (an “unsubstituted heteroalkenyl”) or substituted (a “substituted heteroalkenyl”) with one or more substituents. In certain embodiments, the heteroalkenyl group is an unsubstituted heteroC₂₋₁₀ alkenyl. In certain embodiments, the heteroalkenyl group is a substituted heteroC₂₋₁₀ alkenyl.

[0027] The term “alkynyl” refers to a radical of a straight-chain or branched hydrocarbon group having from 2 to 10 carbon atoms and one or more carbon-carbon triple bonds (*e.g.*, 1, 2, 3, or 4 triple bonds) (“C₂₋₁₀ alkynyl”). In some embodiments, an alkynyl group has 2 to 9 carbon atoms (“C₂₋₉ alkynyl”). In some embodiments, an alkynyl group has 2 to 8 carbon atoms (“C₂₋₈ alkynyl”). In some embodiments, an alkynyl group has 2 to 7 carbon atoms (“C₂₋₇ alkynyl”). In some embodiments, an alkynyl group has 2 to 6 carbon atoms (“C₂₋₆ alkynyl”). In some embodiments, an alkynyl group has 2 to 5 carbon atoms (“C₂₋₅ alkynyl”). In some embodiments, an alkynyl group has 2 to 4 carbon atoms (“C₂₋₄ alkynyl”). In some embodiments, an alkynyl group has 2 to 3 carbon atoms (“C₂₋₃ alkynyl”). In some embodiments, an alkynyl group has 2 carbon atoms (“C₂ alkynyl”). The one or more carbon-carbon triple bonds can be internal (such as in 2-butyne) or terminal (such as in 1-butyne).

Examples of C₂₋₄ alkynyl groups include, without limitation, ethynyl (C₂), 1-propynyl (C₃), 2-propynyl (C₃), 1-butynyl (C₄), 2-butynyl (C₄), and the like. Examples of C₂₋₆ alkenyl groups include the aforementioned C₂₋₄ alkynyl groups as well as pentynyl (C₅), hexynyl (C₆), and the like. Additional examples of alkynyl include heptynyl (C₇), octynyl (C₈), and the like. Unless otherwise specified, each instance of an alkynyl group is independently unsubstituted (an “unsubstituted alkynyl”) or substituted (a “substituted alkynyl”) with one or more substituents. In certain embodiments, the alkynyl group is an unsubstituted C₂₋₁₀ alkynyl. In certain embodiments, the alkynyl group is a substituted C₂₋₁₀ alkynyl.

[0028] The term “heteroalkynyl” refers to an alkynyl group, which further includes at least one heteroatom (*e.g.*, 1, 2, 3, or 4 heteroatoms) selected from oxygen, nitrogen, or sulfur within (*i.e.*, inserted between adjacent carbon atoms of) and/or placed at one or more terminal position(s) of the parent chain. In certain embodiments, a heteroalkynyl group refers to a group having from 2 to 10 carbon atoms, at least one triple bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₁₀ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 9 carbon atoms, at least one triple bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₉ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 8 carbon atoms, at least one triple bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₈ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 7 carbon atoms, at least one triple bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₇ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 6 carbon atoms, at least one triple bond, and 1 or more heteroatoms within the parent chain (“heteroC₂₋₆ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 5 carbon atoms, at least one triple bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₅ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 4 carbon atoms, at least one triple bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₄ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 3 carbon atoms, at least one triple bond, and 1 heteroatom within the parent chain (“heteroC₂₋₃ alkynyl”). In some embodiments, a heteroalkynyl group has 2 to 6 carbon atoms, at least one triple bond, and 1 or 2 heteroatoms within the parent chain (“heteroC₂₋₆ alkynyl”). Unless otherwise specified, each instance of a heteroalkynyl group is independently unsubstituted (an “unsubstituted heteroalkynyl”) or substituted (a “substituted heteroalkynyl”) with one or more substituents. In certain embodiments, the heteroalkynyl group is an unsubstituted heteroC₂₋₁₀ alkynyl. In certain embodiments, the heteroalkynyl group is a substituted heteroC₂₋₁₀ alkynyl.

[0029] The term “carbocyclyl” or “carbocyclic” refers to a radical of a non-aromatic cyclic hydrocarbon group having from 3 to 14 ring carbon atoms (“C₃₋₁₄ carbocyclyl”) and zero heteroatoms in the non-aromatic ring system. In some embodiments, a carbocyclyl group has 3 to 10 ring carbon atoms (“C₃₋₁₀ carbocyclyl”). In some embodiments, a carbocyclyl group has 3 to 8 ring carbon atoms (“C₃₋₈ carbocyclyl”). In some embodiments, a carbocyclyl group has 3 to 7 ring carbon atoms (“C₃₋₇ carbocyclyl”). In some embodiments, a carbocyclyl group has 3 to 6 ring carbon atoms (“C₃₋₆ carbocyclyl”). In some embodiments, a carbocyclyl group has 4 to 6 ring carbon atoms (“C₄₋₆ carbocyclyl”). In some embodiments, a carbocyclyl group has 5 to 6 ring carbon atoms (“C₅₋₆ carbocyclyl”). In some embodiments, a carbocyclyl group has 5 to 10 ring carbon atoms (“C₅₋₁₀ carbocyclyl”). Exemplary C₃₋₆ carbocyclyl groups include, without limitation, cyclopropyl (C₃), cyclopropenyl (C₃), cyclobutyl (C₄), cyclobutenyl (C₄), cyclopentyl (C₅), cyclopentenyl (C₅), cyclohexyl (C₆), cyclohexenyl (C₆), cyclohexadienyl (C₆), and the like. Exemplary C₃₋₈ carbocyclyl groups include, without limitation, the aforementioned C₃₋₆ carbocyclyl groups as well as cycloheptyl (C₇), cycloheptenyl (C₇), cycloheptadienyl (C₇), cycloheptatrienyl (C₇), cyclooctyl (C₈), cyclooctenyl (C₈), bicyclo[2.2.1]heptanyl (C₇), bicyclo[2.2.2]octanyl (C₈), and the like. Exemplary C₃₋₁₀ carbocyclyl groups include, without limitation, the aforementioned C₃₋₈ carbocyclyl groups as well as cyclononyl (C₉), cyclononenyl (C₉), cyclodecyl (C₁₀), cyclodecenyl (C₁₀), octahydro-1H-indenyl (C₉), decahydronaphthalenyl (C₁₀), spiro[4.5]decanyl (C₁₀), and the like. As the foregoing examples illustrate, in certain embodiments, the carbocyclyl group is either monocyclic (“monocyclic carbocyclyl”) or polycyclic (*e.g.*, containing a fused, bridged or spiro ring system such as a bicyclic system (“bicyclic carbocyclyl”) or tricyclic system (“tricyclic carbocyclyl”)) and can be saturated or can contain one or more carbon-carbon double or triple bonds. “Carbocyclyl” also includes ring systems wherein the carbocyclyl ring, as defined above, is fused with one or more aryl or heteroaryl groups wherein the point of attachment is on the carbocyclyl ring, and in such instances, the number of carbons continue to designate the number of carbons in the carbocyclic ring system. Unless otherwise specified, each instance of a carbocyclyl group is independently unsubstituted (an “unsubstituted carbocyclyl”) or substituted (a “substituted carbocyclyl”) with one or more substituents. In certain embodiments, the carbocyclyl group is an unsubstituted C₃₋₁₄ carbocyclyl. In certain embodiments, the carbocyclyl group is a substituted C₃₋₁₄ carbocyclyl.

[0030] In some embodiments, “carbocyclyl” is a monocyclic, saturated carbocyclyl group having from 3 to 14 ring carbon atoms (“C₃₋₁₄ cycloalkyl”). In some embodiments, a

cycloalkyl group has 3 to 10 ring carbon atoms (“C₃₋₁₀ cycloalkyl”). In some embodiments, a cycloalkyl group has 3 to 8 ring carbon atoms (“C₃₋₈ cycloalkyl”). In some embodiments, a cycloalkyl group has 3 to 6 ring carbon atoms (“C₃₋₆ cycloalkyl”). In some embodiments, a cycloalkyl group has 4 to 6 ring carbon atoms (“C₄₋₆ cycloalkyl”). In some embodiments, a cycloalkyl group has 5 to 6 ring carbon atoms (“C₅₋₆ cycloalkyl”). In some embodiments, a cycloalkyl group has 5 to 10 ring carbon atoms (“C₅₋₁₀ cycloalkyl”). Examples of C₅₋₆ cycloalkyl groups include cyclopentyl (C₅) and cyclohexyl (C₆). Examples of C₃₋₆ cycloalkyl groups include the aforementioned C₅₋₆ cycloalkyl groups as well as cyclopropyl (C₃) and cyclobutyl (C₄). Examples of C₃₋₈ cycloalkyl groups include the aforementioned C₃₋₆ cycloalkyl groups as well as cycloheptyl (C₇) and cyclooctyl (C₈). Unless otherwise specified, each instance of a cycloalkyl group is independently unsubstituted (an “unsubstituted cycloalkyl”) or substituted (a “substituted cycloalkyl”) with one or more substituents. In certain embodiments, the cycloalkyl group is an unsubstituted C₃₋₁₄ cycloalkyl. In certain embodiments, the cycloalkyl group is a substituted C₃₋₁₄ cycloalkyl.

[0031] The term “heterocyclyl” or “heterocyclic” refers to a radical of a 3- to 14-membered non-aromatic ring system having ring carbon atoms and 1 to 4 ring heteroatoms, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur (“3-14 membered heterocyclyl”). In heterocyclyl groups that contain one or more nitrogen atoms, the point of attachment can be a carbon or nitrogen atom, as valency permits. A heterocyclyl group can either be monocyclic (“monocyclic heterocyclyl”) or polycyclic (*e.g.*, a fused, bridged or spiro ring system such as a bicyclic system (“bicyclic heterocyclyl”) or tricyclic system (“tricyclic heterocyclyl”)), and can be saturated or can contain one or more carbon-carbon double or triple bonds. Heterocyclyl polycyclic ring systems can include one or more heteroatoms in one or both rings. “Heterocyclyl” also includes ring systems wherein the heterocyclyl ring, as defined above, is fused with one or more carbocyclyl groups wherein the point of attachment is either on the carbocyclyl or heterocyclyl ring, or ring systems wherein the heterocyclyl ring, as defined above, is fused with one or more aryl or heteroaryl groups, wherein the point of attachment is on the heterocyclyl ring, and in such instances, the number of ring members continue to designate the number of ring members in the heterocyclyl ring system. Unless otherwise specified, each instance of heterocyclyl is independently unsubstituted (an “unsubstituted heterocyclyl”) or substituted (a “substituted heterocyclyl”) with one or more substituents. In certain embodiments, the heterocyclyl group is an unsubstituted 3-14 membered heterocyclyl. In certain embodiments, the heterocyclyl group is a substituted 3-14 membered heterocyclyl.

[0032] In some embodiments, a heterocyclyl group is a 5-10 membered non-aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur ("5-10 membered heterocyclyl"). In some embodiments, a heterocyclyl group is a 5-8 membered non-aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur ("5-8 membered heterocyclyl"). In some embodiments, a heterocyclyl group is a 5-6 membered non-aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur ("5-6 membered heterocyclyl"). In some embodiments, the 5-6 membered heterocyclyl has 1-3 ring heteroatoms selected from nitrogen, oxygen, and sulfur. In some embodiments, the 5-6 membered heterocyclyl has 1-2 ring heteroatoms selected from nitrogen, oxygen, and sulfur. In some embodiments, the 5-6 membered heterocyclyl has 1 ring heteroatom selected from nitrogen, oxygen, and sulfur.

[0033] Exemplary 3-membered heterocyclyl groups containing 1 heteroatom include, without limitation, azirdinyl, oxiranyl, and thiiranyl. Exemplary 4-membered heterocyclyl groups containing 1 heteroatom include, without limitation, azetidiny, oxetanyl, and thietanyl. Exemplary 5-membered heterocyclyl groups containing 1 heteroatom include, without limitation, tetrahydrofuranyl, dihydrofuranyl, tetrahydrothiophenyl, dihydrothiophenyl, pyrrolidinyl, dihydropyrrolyl, and pyrrolyl-2,5-dione. Exemplary 5-membered heterocyclyl groups containing 2 heteroatoms include, without limitation, dioxolanyl, oxathiolanyl and dithiolanyl. Exemplary 5-membered heterocyclyl groups containing 3 heteroatoms include, without limitation, triazoliny, oxadiazoliny, and thiadiazoliny. Exemplary 6-membered heterocyclyl groups containing 1 heteroatom include, without limitation, piperidinyl, tetrahydropyranyl, dihydropyridiny, and thianyl. Exemplary 6-membered heterocyclyl groups containing 2 heteroatoms include, without limitation, piperaziny, morpholiny, dithianyl, and dioxanyl. Exemplary 6-membered heterocyclyl groups containing 3 heteroatoms include, without limitation, triazinyl. Exemplary 7-membered heterocyclyl groups containing 1 heteroatom include, without limitation, azepanyl, oxepanyl and thiepanyl. Exemplary 8-membered heterocyclyl groups containing 1 heteroatom include, without limitation, azocanyl, oxecanyl and thiocanyl. Exemplary bicyclic heterocyclyl groups include, without limitation, indoliny, isoindoliny, dihydrobenzofuranyl, dihydrobenzothienyl, tetrahydrobenzothienyl, tetrahydrobenzofuranyl, tetrahydroindolyl, tetrahydroquinoliny, tetrahydroisoquinoliny, decahydroquinoliny, decahydroisoquinoliny, octahydrochromenyl, octahydroisochromenyl, decahydronaphthyridiny, decahydro-1,8-

naphthyridinyl, octahydropyrrolo[3,2-b]pyrrole, indolinyl, phthalimidyl, naphthalimidyl, chromanyl, chromenyl, 1H-benzo[e][1,4]diazepinyl, 1,4,5,7-tetrahydropyrano[3,4-b]pyrrolyl, 5,6-dihydro-4H-furo[3,2-b]pyrrolyl, 6,7-dihydro-5H-furo[3,2-b]pyranyl, 5,7-dihydro-4H-thieno[2,3-c]pyranyl, 2,3-dihydro-1H-pyrrolo[2,3-b]pyridinyl, 2,3-dihydrofuro[2,3-b]pyridinyl, 4,5,6,7-tetrahydro-1H-pyrrolo[2,3-b]pyridinyl, 4,5,6,7-tetrahydrofuro[3,2-c]pyridinyl, 4,5,6,7-tetrahydrothieno[3,2-b]pyridinyl, 1,2,3,4-tetrahydro-1,6-naphthyridinyl, and the like.

[0034] The term “aryl” refers to a radical of a monocyclic or polycyclic (*e.g.*, bicyclic or tricyclic) $4n+2$ aromatic ring system (*e.g.*, having 6, 10, or 14 π electrons shared in a cyclic array) having 6-14 ring carbon atoms and zero heteroatoms provided in the aromatic ring system (“C₆₋₁₄ aryl”). In some embodiments, an aryl group has 6 ring carbon atoms (“C₆ aryl”; *e.g.*, phenyl). In some embodiments, an aryl group has 10 ring carbon atoms (“C₁₀ aryl”; *e.g.*, naphthyl such as 1-naphthyl and 2-naphthyl). In some embodiments, an aryl group has 14 ring carbon atoms (“C₁₄ aryl”; *e.g.*, anthracyl). “Aryl” also includes ring systems wherein the aryl ring, as defined above, is fused with one or more carbocyclyl or heterocyclyl groups wherein the radical or point of attachment is on the aryl ring, and in such instances, the number of carbon atoms continue to designate the number of carbon atoms in the aryl ring system. Unless otherwise specified, each instance of an aryl group is independently unsubstituted (an “unsubstituted aryl”) or substituted (a “substituted aryl”) with one or more substituents. In certain embodiments, the aryl group is an unsubstituted C₆₋₁₄ aryl. In certain embodiments, the aryl group is a substituted C₆₋₁₄ aryl.

[0035] The term “heteroaryl” refers to a radical of a 5-14 membered monocyclic or polycyclic (*e.g.*, bicyclic, tricyclic) $4n+2$ aromatic ring system (*e.g.*, having 6, 10, or 14 π electrons shared in a cyclic array) having ring carbon atoms and 1-4 ring heteroatoms provided in the aromatic ring system, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur (“5-14 membered heteroaryl”). In heteroaryl groups that contain one or more nitrogen atoms, the point of attachment can be a carbon or nitrogen atom, as valency permits. Heteroaryl polycyclic ring systems can include one or more heteroatoms in one or both rings. “Heteroaryl” includes ring systems wherein the heteroaryl ring, as defined above, is fused with one or more carbocyclyl or heterocyclyl groups wherein the point of attachment is on the heteroaryl ring, and in such instances, the number of ring members continue to designate the number of ring members in the heteroaryl ring system. “Heteroaryl” also includes ring systems wherein the heteroaryl ring, as defined above, is

fused with one or more aryl groups wherein the point of attachment is either on the aryl or heteroaryl ring, and in such instances, the number of ring members designates the number of ring members in the fused polycyclic (aryl/heteroaryl) ring system. Polycyclic heteroaryl groups wherein one ring does not contain a heteroatom (*e.g.*, indolyl, quinolinyl, carbazolyl, and the like) the point of attachment can be on either ring, *i.e.*, either the ring bearing a heteroatom (*e.g.*, 2-indolyl) or the ring that does not contain a heteroatom (*e.g.*, 5-indolyl).

[0036] In some embodiments, a heteroaryl group is a 5-10 membered aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms provided in the aromatic ring system, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur (“5-10 membered heteroaryl”). In some embodiments, a heteroaryl group is a 5-8 membered aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms provided in the aromatic ring system, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur (“5-8 membered heteroaryl”). In some embodiments, a heteroaryl group is a 5-6 membered aromatic ring system having ring carbon atoms and 1-4 ring heteroatoms provided in the aromatic ring system, wherein each heteroatom is independently selected from nitrogen, oxygen, and sulfur (“5-6 membered heteroaryl”). In some embodiments, the 5-6 membered heteroaryl has 1-3 ring heteroatoms selected from nitrogen, oxygen, and sulfur. In some embodiments, the 5-6 membered heteroaryl has 1-2 ring heteroatoms selected from nitrogen, oxygen, and sulfur. In some embodiments, the 5-6 membered heteroaryl has 1 ring heteroatom selected from nitrogen, oxygen, and sulfur. Unless otherwise specified, each instance of a heteroaryl group is independently unsubstituted (an “unsubstituted heteroaryl”) or substituted (a “substituted heteroaryl”) with one or more substituents. In certain embodiments, the heteroaryl group is an unsubstituted 5-14 membered heteroaryl. In certain embodiments, the heteroaryl group is a substituted 5-14 membered heteroaryl.

[0037] Exemplary 5-membered heteroaryl groups containing 1 heteroatom include, without limitation, pyrrolyl, furanyl, and thiophenyl. Exemplary 5-membered heteroaryl groups containing 2 heteroatoms include, without limitation, imidazolyl, pyrazolyl, oxazolyl, isoxazolyl, thiazolyl, and isothiazolyl. Exemplary 5-membered heteroaryl groups containing 3 heteroatoms include, without limitation, triazolyl, oxadiazolyl, and thiadiazolyl. Exemplary 5-membered heteroaryl groups containing 4 heteroatoms include, without limitation, tetrazolyl. Exemplary 6-membered heteroaryl groups containing 1 heteroatom include, without limitation, pyridinyl. Exemplary 6-membered heteroaryl groups containing 2 heteroatoms include, without limitation, pyridazinyl, pyrimidinyl, and pyrazinyl. Exemplary 6-membered heteroaryl groups containing 3 or 4 heteroatoms include, without limitation,

triazinyl and tetrazinyl, respectively. Exemplary 7-membered heteroaryl groups containing 1 heteroatom include, without limitation, azepinyl, oxepinyl, and thiepinyl. Exemplary 5,6-bicyclic heteroaryl groups include, without limitation, indolyl, isoindolyl, indazolyl, benzotriazolyl, benzothiophenyl, isobenzothiophenyl, benzofuranyl, benzoisofuranyl, benzimidazolyl, benzoxazolyl, benzisoxazolyl, benzoxadiazolyl, benzthiazolyl, benzisothiazolyl, benzthiadiazolyl, indolizinyl, and purinyl. Exemplary 6,6-bicyclic heteroaryl groups include, without limitation, naphthyridinyl, pteridinyl, quinolinyl, isoquinolinyl, cinnolinyl, quinoxalinyl, phthalazinyl, and quinazolinyl. Exemplary tricyclic heteroaryl groups include, without limitation, phenanthridinyl, dibenzofuranyl, carbazolyl, acridinyl, phenothiazinyl, phenoxazinyl, and phenazinyl.

[0038] The term “unsaturated bond” refers to a double or triple bond.

[0039] The term “unsaturated” or “partially unsaturated” refers to a moiety that includes at least one double or triple bond.

[0040] The term “saturated” refers to a moiety that does not contain a double or triple bond, *i.e.*, the moiety only contains single bonds.

[0041] Affixing the suffix “-ene” to a group indicates the group is a divalent moiety, *e.g.*, alkylene is the divalent moiety of alkyl, alkenylene is the divalent moiety of alkenyl, alkynylene is the divalent moiety of alkynyl, heteroalkylene is the divalent moiety of heteroalkyl, heteroalkenylene is the divalent moiety of heteroalkenyl, heteroalkynylene is the divalent moiety of heteroalkynyl, carbocyclylene is the divalent moiety of carbocyclyl, heterocyclylene is the divalent moiety of heterocyclyl, arylene is the divalent moiety of aryl, and heteroarylene is the divalent moiety of heteroaryl.

[0042] A group is optionally substituted unless expressly provided otherwise. The term “optionally substituted” refers to being substituted or unsubstituted. In certain embodiments, alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl groups are optionally substituted. “Optionally substituted” refers to a group which may be substituted or unsubstituted (*e.g.*, “substituted” or “unsubstituted” alkyl, “substituted” or “unsubstituted” alkenyl, “substituted” or “unsubstituted” alkynyl, “substituted” or “unsubstituted” heteroalkyl, “substituted” or “unsubstituted” heteroalkenyl, “substituted” or “unsubstituted” heteroalkynyl, “substituted” or “unsubstituted” carbocyclyl, “substituted” or “unsubstituted” heterocyclyl, “substituted” or “unsubstituted” aryl or “substituted” or “unsubstituted” heteroaryl group). In general, the term “substituted” means that at least one hydrogen present on a group is replaced with a permissible substituent, *e.g.*, a substituent which upon substitution results in a stable compound, *e.g.*, a compound which

does not spontaneously undergo transformation such as by rearrangement, cyclization, elimination, or other reaction. Unless otherwise indicated, a “substituted” group has a substituent at one or more substitutable positions of the group, and when more than one position in any given structure is substituted, the substituent is either the same or different at each position. The term “substituted” is contemplated to include substitution with all permissible substituents of organic compounds, and includes any of the substituents described herein that results in the formation of a stable compound. The present invention contemplates any and all such combinations in order to arrive at a stable compound. For purposes of this invention, heteroatoms such as nitrogen may have hydrogen substituents and/or any suitable substituent as described herein which satisfy the valencies of the heteroatoms and results in the formation of a stable moiety. The invention is not intended to be limited in any manner by the exemplary substituents described herein.

[0043] Exemplary carbon atom substituents include, but are not limited to, halogen, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}_3$, $-\text{SO}_2\text{H}$, $-\text{SO}_3\text{H}$, $-\text{OH}$, $-\text{OR}^{\text{aa}}$, $-\text{ON}(\text{R}^{\text{bb}})_2$, $-\text{N}(\text{R}^{\text{bb}})_2$, $-\text{N}(\text{R}^{\text{bb}})_3^+\text{X}^-$, $-\text{N}(\text{OR}^{\text{cc}})\text{R}^{\text{bb}}$, $-\text{SH}$, $-\text{SR}^{\text{aa}}$, $-\text{SSR}^{\text{cc}}$, $-\text{C}(=\text{O})\text{R}^{\text{aa}}$, $-\text{CO}_2\text{H}$, $-\text{CHO}$, $-\text{C}(\text{OR}^{\text{cc}})_3$, $-\text{CO}_2\text{R}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OCO}_2\text{R}^{\text{aa}}$, $-\text{C}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OC}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{NR}^{\text{bb}}\text{C}(=\text{O})\text{R}^{\text{aa}}$, $-\text{NR}^{\text{bb}}\text{CO}_2\text{R}^{\text{aa}}$, $-\text{NR}^{\text{bb}}\text{C}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{C}(=\text{NR}^{\text{bb}})\text{R}^{\text{aa}}$, $-\text{C}(=\text{NR}^{\text{bb}})\text{OR}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{R}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{OR}^{\text{aa}}$, $-\text{C}(=\text{NR}^{\text{bb}})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{NR}^{\text{bb}}\text{C}(=\text{NR}^{\text{bb}})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{C}(=\text{O})\text{NR}^{\text{bb}}\text{SO}_2\text{R}^{\text{aa}}$, $-\text{NR}^{\text{bb}}\text{SO}_2\text{R}^{\text{aa}}$, $-\text{SO}_2\text{N}(\text{R}^{\text{bb}})_2$, $-\text{SO}_2\text{R}^{\text{aa}}$, $-\text{SO}_2\text{OR}^{\text{aa}}$, $-\text{OSO}_2\text{R}^{\text{aa}}$, $-\text{S}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OS}(=\text{O})\text{R}^{\text{aa}}$, $-\text{Si}(\text{R}^{\text{aa}})_3$, $-\text{OSi}(\text{R}^{\text{aa}})_3$, $-\text{C}(=\text{S})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{C}(=\text{O})\text{SR}^{\text{aa}}$, $-\text{C}(=\text{S})\text{SR}^{\text{aa}}$, $-\text{SC}(=\text{S})\text{SR}^{\text{aa}}$, $-\text{SC}(=\text{O})\text{SR}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{SR}^{\text{aa}}$, $-\text{SC}(=\text{O})\text{OR}^{\text{aa}}$, $-\text{SC}(=\text{O})\text{R}^{\text{aa}}$, $-\text{P}(=\text{O})(\text{R}^{\text{aa}})_2$, $-\text{P}(=\text{O})(\text{OR}^{\text{cc}})_2$, $-\text{OP}(=\text{O})(\text{R}^{\text{aa}})_2$, $-\text{OP}(=\text{O})(\text{OR}^{\text{cc}})_2$, $-\text{P}(=\text{O})(\text{N}(\text{R}^{\text{bb}})_2)_2$, $-\text{OP}(=\text{O})(\text{N}(\text{R}^{\text{bb}})_2)_2$, $-\text{NR}^{\text{bb}}\text{P}(=\text{O})(\text{R}^{\text{aa}})_2$, $-\text{NR}^{\text{bb}}\text{P}(=\text{O})(\text{OR}^{\text{cc}})_2$, $-\text{NR}^{\text{bb}}\text{P}(=\text{O})(\text{N}(\text{R}^{\text{bb}})_2)_2$, $-\text{P}(\text{R}^{\text{cc}})_2$, $-\text{P}(\text{OR}^{\text{cc}})_2$, $-\text{P}(\text{R}^{\text{cc}})_3^+\text{X}^-$, $-\text{P}(\text{OR}^{\text{cc}})_3^+\text{X}^-$, $-\text{P}(\text{R}^{\text{cc}})_4$, $-\text{P}(\text{OR}^{\text{cc}})_4$, $-\text{OP}(\text{R}^{\text{cc}})_2$, $-\text{OP}(\text{R}^{\text{cc}})_3^+\text{X}^-$, $-\text{OP}(\text{OR}^{\text{cc}})_2$, $-\text{OP}(\text{OR}^{\text{cc}})_3^+\text{X}^-$, $-\text{OP}(\text{R}^{\text{cc}})_4$, $-\text{OP}(\text{OR}^{\text{cc}})_4$, $-\text{B}(\text{R}^{\text{aa}})_2$, $-\text{B}(\text{OR}^{\text{cc}})_2$, $-\text{BR}^{\text{aa}}(\text{OR}^{\text{cc}})$, C_{1-10} alkyl, C_{1-10} perhaloalkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, hetero C_{1-10} alkyl, hetero C_{2-10} alkenyl, hetero C_{2-10} alkynyl, C_{3-10} carbocyclyl, 3-14 membered heterocyclyl, C_{6-14} aryl, and 5-14 membered heteroaryl, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups; wherein X^- is a counterion;

or two geminal hydrogens on a carbon atom are replaced with the group $=\text{O}$, $=\text{S}$, $=\text{NN}(\text{R}^{\text{bb}})_2$, $=\text{NNR}^{\text{bb}}\text{C}(=\text{O})\text{R}^{\text{aa}}$, $=\text{NNR}^{\text{bb}}\text{C}(=\text{O})\text{OR}^{\text{aa}}$, $=\text{NNR}^{\text{bb}}\text{S}(=\text{O})_2\text{R}^{\text{aa}}$, $=\text{NR}^{\text{bb}}$, or $=\text{NOR}^{\text{cc}}$; each instance of R^{aa} is, independently, selected from C_{1-10} alkyl, C_{1-10} perhaloalkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, hetero C_{1-10} alkyl, hetero C_{2-10} alkenyl, hetero C_{2-10} alkynyl, C_{3-10}

carbocyclyl, 3-14 membered heterocyclyl, C₆₋₁₄ aryl, and 5-14 membered heteroaryl, or two R^{aa} groups are joined to form a 3-14 membered heterocyclyl or 5-14 membered heteroaryl ring, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups;

each instance of R^{bb} is, independently, selected from hydrogen, -OH, -OR^{aa}, -N(R^{cc})₂, -CN, -C(=O)R^{aa}, -C(=O)N(R^{cc})₂, -CO₂R^{aa}, -SO₂R^{aa}, -C(=NR^{cc})OR^{aa}, -C(=NR^{cc})N(R^{cc})₂, -SO₂N(R^{cc})₂, -SO₂R^{cc}, -SO₂OR^{cc}, -SOR^{aa}, -C(=S)N(R^{cc})₂, -C(=O)SR^{cc}, -C(=S)SR^{cc}, -P(=O)(R^{aa})₂, -P(=O)(OR^{cc})₂, -P(=O)(N(R^{cc})₂)₂, C₁₋₁₀ alkyl, C₁₋₁₀ perhaloalkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, heteroC₁₋₁₀ alkyl, heteroC₂₋₁₀ alkenyl, heteroC₂₋₁₀ alkynyl, C₃₋₁₀ carbocyclyl, 3-14 membered heterocyclyl, C₆₋₁₄ aryl, and 5-14 membered heteroaryl, or two R^{bb} groups are joined to form a 3-14 membered heterocyclyl or 5-14 membered heteroaryl ring, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups; wherein X⁻ is a counterion;

each instance of R^{cc} is, independently, selected from hydrogen, C₁₋₁₀ alkyl, C₁₋₁₀ perhaloalkyl, C₂₋₁₀ alkenyl, C₂₋₁₀ alkynyl, heteroC₁₋₁₀ alkyl, heteroC₂₋₁₀ alkenyl, heteroC₂₋₁₀ alkynyl, C₃₋₁₀ carbocyclyl, 3-14 membered heterocyclyl, C₆₋₁₄ aryl, and 5-14 membered heteroaryl, or two R^{cc} groups are joined to form a 3-14 membered heterocyclyl or 5-14 membered heteroaryl ring, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups;

each instance of R^{dd} is, independently, selected from halogen, -CN, -NO₂, -N₃, -SO₂H, -SO₃H, -OH, -OR^{ee}, -ON(R^{ff})₂, -N(R^{ff})₂, -N(R^{ff})₃⁺X⁻, -N(OR^{ee})R^{ff}, -SH, -SR^{ee}, -SSR^{ee}, -C(=O)R^{ee}, -CO₂H, -CO₂R^{ee}, -OC(=O)R^{ee}, -OCO₂R^{ee}, -C(=O)N(R^{ff})₂, -OC(=O)N(R^{ff})₂, -NR^{ff}C(=O)R^{ee}, -NR^{ff}CO₂R^{ee}, -NR^{ff}C(=O)N(R^{ff})₂, -C(=NR^{ff})OR^{ee}, -OC(=NR^{ff})R^{ee}, -OC(=NR^{ff})OR^{ee}, -C(=NR^{ff})N(R^{ff})₂, -OC(=NR^{ff})N(R^{ff})₂, -NR^{ff}C(=NR^{ff})N(R^{ff})₂, -NR^{ff}SO₂R^{ee}, -SO₂N(R^{ff})₂, -SO₂R^{ee}, -SO₂OR^{ee}, -OSO₂R^{ee}, -S(=O)R^{ee}, -Si(R^{ee})₃, -OSi(R^{ee})₃, -C(=S)N(R^{ff})₂, -C(=O)SR^{ee}, -C(=S)SR^{ee}, -SC(=S)SR^{ee}, -P(=O)(OR^{ee})₂, -P(=O)(R^{ee})₂, -OP(=O)(R^{ee})₂, -OP(=O)(OR^{ee})₂, C₁₋₆ alkyl, C₁₋₆ perhaloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, heteroC₁₋₆ alkyl, heteroC₂₋₆ alkenyl, heteroC₂₋₆ alkynyl, C₃₋₁₀ carbocyclyl, 3-10 membered heterocyclyl, C₆₋₁₀ aryl, 5-10 membered heteroaryl, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{gg}

groups, or two geminal R^{dd} substituents can be joined to form =O or =S; wherein X⁻ is a counterion;

each instance of R^{ee} is, independently, selected from C₁₋₆ alkyl, C₁₋₆ perhaloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, heteroC₁₋₆ alkyl, heteroC₂₋₆ alkenyl, heteroC₂₋₆ alkynyl, C₃₋₁₀ carbocyclyl, C₆₋₁₀ aryl, 3-10 membered heterocyclyl, and 3-10 membered heteroaryl, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{gg} groups;

each instance of R^{ff} is, independently, selected from hydrogen, C₁₋₆ alkyl, C₁₋₆ perhaloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, heteroC₁₋₆ alkyl, heteroC₂₋₆ alkenyl, heteroC₂₋₆ alkynyl, C₃₋₁₀ carbocyclyl, 3-10 membered heterocyclyl, C₆₋₁₀ aryl and 5-10 membered heteroaryl, or two R^{ff} groups are joined to form a 3-10 membered heterocyclyl or 5-10 membered heteroaryl ring, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{gg} groups; and

each instance of R^{gg} is, independently, halogen, -CN, -NO₂, -N₃, -SO₂H, -SO₃H, -OH, -OC₁₋₆ alkyl, -ON(C₁₋₆ alkyl)₂, -N(C₁₋₆ alkyl)₂, -N(C₁₋₆ alkyl)₃⁺X⁻, -NH(C₁₋₆ alkyl)₂⁺X⁻, -NH₂(C₁₋₆ alkyl)⁺X⁻, -NH₃⁺X⁻, -N(OC₁₋₆ alkyl)(C₁₋₆ alkyl), -N(OH)(C₁₋₆ alkyl), -NH(OH), -SH, -SC₁₋₆ alkyl, -SS(C₁₋₆ alkyl), -C(=O)(C₁₋₆ alkyl), -CO₂H, -CO₂(C₁₋₆ alkyl), -OC(=O)(C₁₋₆ alkyl), -OCO₂(C₁₋₆ alkyl), -C(=O)NH₂, -C(=O)N(C₁₋₆ alkyl)₂, -OC(=O)NH(C₁₋₆ alkyl), -NHC(=O)(C₁₋₆ alkyl), -N(C₁₋₆ alkyl)C(=O)(C₁₋₆ alkyl), -NHCO₂(C₁₋₆ alkyl), -NHC(=O)N(C₁₋₆ alkyl)₂, -NHC(=O)NH(C₁₋₆ alkyl), -NHC(=O)NH₂, -C(=NH)O(C₁₋₆ alkyl), -OC(=NH)(C₁₋₆ alkyl), -OC(=NH)OC₁₋₆ alkyl, -C(=NH)N(C₁₋₆ alkyl)₂, -C(=NH)NH(C₁₋₆ alkyl), -C(=NH)NH₂, -OC(=NH)N(C₁₋₆ alkyl)₂, -OC(=NH)NH(C₁₋₆ alkyl), -OC(=NH)NH₂, -NHC(=NH)N(C₁₋₆ alkyl)₂, -NHC(=NH)NH₂, -NHCO₂(C₁₋₆ alkyl), -SO₂N(C₁₋₆ alkyl)₂, -SO₂NH(C₁₋₆ alkyl), -SO₂NH₂, -SO₂(C₁₋₆ alkyl), -SO₂O(C₁₋₆ alkyl), -OSO₂(C₁₋₆ alkyl), -SO(C₁₋₆ alkyl), -Si(C₁₋₆ alkyl)₃, -OSi(C₁₋₆ alkyl)₃, -C(=S)N(C₁₋₆ alkyl)₂, C(=S)NH(C₁₋₆ alkyl), C(=S)NH₂, -C(=O)S(C₁₋₆ alkyl), -C(=S)SC₁₋₆ alkyl, -SC(=S)SC₁₋₆ alkyl, -P(=O)(OC₁₋₆ alkyl)₂, -P(=O)(C₁₋₆ alkyl)₂, -OP(=O)(C₁₋₆ alkyl)₂, -OP(=O)(OC₁₋₆ alkyl)₂, C₁₋₆ alkyl, C₁₋₆ perhaloalkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, heteroC₁₋₆ alkyl, heteroC₂₋₆ alkenyl, heteroC₂₋₆ alkynyl, C₃₋₁₀ carbocyclyl, C₆₋₁₀ aryl, 3-10 membered heterocyclyl, 5-10 membered heteroaryl; or two geminal R^{gg} substituents can be joined to form =O or =S; wherein X⁻ is a counterion.

[0044] In certain embodiments, carbon atom substituents include: halogen, $-\text{CN}$, $-\text{NO}_2$, $-\text{N}_3$, $-\text{SO}_2\text{H}$, $-\text{SO}_3\text{H}$, $-\text{OH}$, $-\text{OC}_{1-6}\text{ alkyl}$, $-\text{ON}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{N}(\text{C}_{1-6}\text{ alkyl})_3^+\text{X}^-$, $-\text{NH}(\text{C}_{1-6}\text{ alkyl})_2^+\text{X}^-$, $-\text{NH}_2(\text{C}_{1-6}\text{ alkyl})^+\text{X}^-$, $-\text{NH}_3^+\text{X}^-$, $-\text{N}(\text{OC}_{1-6}\text{ alkyl})(\text{C}_{1-6}\text{ alkyl})$, $-\text{N}(\text{OH})(\text{C}_{1-6}\text{ alkyl})$, $-\text{NH}(\text{OH})$, $-\text{SH}$, $-\text{SC}_{1-6}\text{ alkyl}$, $-\text{SS}(\text{C}_{1-6}\text{ alkyl})$, $-\text{C}(=\text{O})(\text{C}_{1-6}\text{ alkyl})$, $-\text{CO}_2\text{H}$, $-\text{CO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{OC}(=\text{O})(\text{C}_{1-6}\text{ alkyl})$, $-\text{OCO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{C}(=\text{O})\text{NH}_2$, $-\text{C}(=\text{O})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{OC}(=\text{O})\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $-\text{NHC}(=\text{O})(\text{C}_{1-6}\text{ alkyl})$, $-\text{N}(\text{C}_{1-6}\text{ alkyl})\text{C}(=\text{O})(\text{C}_{1-6}\text{ alkyl})$, $-\text{NHCO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{NHC}(=\text{O})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{NHC}(=\text{O})\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $-\text{NHC}(=\text{O})\text{NH}_2$, $-\text{C}(=\text{NH})\text{O}(\text{C}_{1-6}\text{ alkyl})$, $-\text{OC}(=\text{NH})(\text{C}_{1-6}\text{ alkyl})$, $-\text{OC}(=\text{NH})\text{OC}_{1-6}\text{ alkyl}$, $-\text{C}(=\text{NH})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{C}(=\text{NH})\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $-\text{C}(=\text{NH})\text{NH}_2$, $-\text{OC}(=\text{NH})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{OC}(=\text{NH})\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $-\text{OC}(=\text{NH})\text{NH}_2$, $-\text{NHC}(=\text{NH})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{NHC}(=\text{NH})\text{NH}_2$, $-\text{NHSO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{SO}_2\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{SO}_2\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $-\text{SO}_2\text{NH}_2$, $-\text{SO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{SO}_2\text{O}(\text{C}_{1-6}\text{ alkyl})$, $-\text{OSO}_2(\text{C}_{1-6}\text{ alkyl})$, $-\text{SO}(\text{C}_{1-6}\text{ alkyl})$, $-\text{Si}(\text{C}_{1-6}\text{ alkyl})_3$, $-\text{OSi}(\text{C}_{1-6}\text{ alkyl})_3$, $-\text{C}(=\text{S})\text{N}(\text{C}_{1-6}\text{ alkyl})_2$, $\text{C}(=\text{S})\text{NH}(\text{C}_{1-6}\text{ alkyl})$, $\text{C}(=\text{S})\text{NH}_2$, $-\text{C}(=\text{O})\text{S}(\text{C}_{1-6}\text{ alkyl})$, $-\text{C}(=\text{S})\text{SC}_{1-6}\text{ alkyl}$, $-\text{SC}(=\text{S})\text{SC}_{1-6}\text{ alkyl}$, $-\text{P}(=\text{O})(\text{OC}_{1-6}\text{ alkyl})_2$, $-\text{P}(=\text{O})(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{OP}(=\text{O})(\text{C}_{1-6}\text{ alkyl})_2$, $-\text{OP}(=\text{O})(\text{OC}_{1-6}\text{ alkyl})_2$, $\text{C}_{1-6}\text{ alkyl}$, $\text{C}_{1-6}\text{ perhaloalkyl}$, $\text{C}_{2-6}\text{ alkenyl}$, $\text{C}_{2-6}\text{ alkynyl}$, $\text{heteroC}_{1-6}\text{ alkyl}$, $\text{heteroC}_{2-6}\text{ alkenyl}$, $\text{heteroC}_{2-6}\text{ alkynyl}$, $\text{C}_{3-10}\text{ carbocyclyl}$, $\text{C}_{6-10}\text{ aryl}$, $3\text{-}10\text{ membered heterocyclyl}$, $5\text{-}10\text{ membered heteroaryl}$; or two geminal R^{gg} substituents can be joined to form $=\text{O}$ or $=\text{S}$; wherein X^- is a counterion.

[0045] The term “halo” or “halogen” refers to fluorine (fluoro, $-\text{F}$), chlorine (chloro, $-\text{Cl}$), bromine (bromo, $-\text{Br}$), or iodine (iodo, $-\text{I}$).

[0046] The term “hydroxyl” or “hydroxy” refers to the group $-\text{OH}$. The term “substituted hydroxyl” or “substituted hydroxyl,” by extension, refers to a hydroxyl group wherein the oxygen atom directly attached to the parent molecule is substituted with a group other than hydrogen, and includes groups selected from $-\text{OR}^{\text{aa}}$, $-\text{ON}(\text{R}^{\text{bb}})_2$, $-\text{OC}(=\text{O})\text{SR}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OCO}_2\text{R}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{R}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{OR}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OS}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OSO}_2\text{R}^{\text{aa}}$, $-\text{OSi}(\text{R}^{\text{aa}})_3$, $-\text{OP}(\text{R}^{\text{cc}})_2$, $-\text{OP}(\text{R}^{\text{cc}})_3^+\text{X}^-$, $-\text{OP}(\text{OR}^{\text{cc}})_2$, $-\text{OP}(\text{OR}^{\text{cc}})_3^+\text{X}^-$, $-\text{OP}(=\text{O})(\text{R}^{\text{aa}})_2$, $-\text{OP}(=\text{O})(\text{OR}^{\text{cc}})_2$, and $-\text{OP}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, wherein X^- , R^{aa} , R^{bb} , and R^{cc} are as defined herein.

[0047] The term “amino” refers to the group $-\text{NH}_2$. The term “substituted amino,” by extension, refers to a monosubstituted amino, a disubstituted amino, or a trisubstituted amino. In certain embodiments, the “substituted amino” is a monosubstituted amino or a disubstituted amino group.

[0048] The term “monosubstituted amino” refers to an amino group wherein the nitrogen atom directly attached to the parent molecule is substituted with one hydrogen and one group

other than hydrogen, and includes groups selected from $\text{--NH(R}^{\text{bb}}\text{)}$, $\text{--NHC(=O)R}^{\text{aa}}$, $\text{--NHCO}_2\text{R}^{\text{aa}}$, $\text{--NHC(=O)N(R}^{\text{bb}}\text{)}_2$, $\text{--NHC(=NR}^{\text{bb}}\text{)N(R}^{\text{bb}}\text{)}_2$, $\text{--NHSO}_2\text{R}^{\text{aa}}$, $\text{--NHP(=O)(OR}^{\text{cc}}\text{)}_2$, and $\text{--NHP(=O)(N(R}^{\text{bb}}\text{)}_2\text{)}_2$, wherein R^{aa} , R^{bb} and R^{cc} are as defined herein, and wherein R^{bb} of the group $\text{--NH(R}^{\text{bb}}\text{)}$ is not hydrogen.

[0049] The term “disubstituted amino” refers to an amino group wherein the nitrogen atom directly attached to the parent molecule is substituted with two groups other than hydrogen, and includes groups selected from $\text{--N(R}^{\text{bb}}\text{)}_2$, $\text{--NR}^{\text{bb}}\text{C(=O)R}^{\text{aa}}$, $\text{--NR}^{\text{bb}}\text{CO}_2\text{R}^{\text{aa}}$, $\text{--NR}^{\text{bb}}\text{C(=O)N(R}^{\text{bb}}\text{)}_2$, $\text{--NR}^{\text{bb}}\text{C(=NR}^{\text{bb}}\text{)N(R}^{\text{bb}}\text{)}_2$, $\text{--NR}^{\text{bb}}\text{SO}_2\text{R}^{\text{aa}}$, $\text{--NR}^{\text{bb}}\text{P(=O)(OR}^{\text{cc}}\text{)}_2$, and $\text{--NR}^{\text{bb}}\text{P(=O)(N(R}^{\text{bb}}\text{)}_2\text{)}_2$, wherein R^{aa} , R^{bb} , and R^{cc} are as defined herein, with the proviso that the nitrogen atom directly attached to the parent molecule is not substituted with hydrogen.

[0050] The term “trisubstituted amino” refers to an amino group wherein the nitrogen atom directly attached to the parent molecule is substituted with three groups, and includes groups selected from $\text{--N(R}^{\text{bb}}\text{)}_3$ and $\text{--N(R}^{\text{bb}}\text{)}_3^+\text{X}^-$, wherein R^{bb} and X^- are as defined herein.

[0051] The term “sulfonyl” refers to a group selected from $\text{--SO}_2\text{N(R}^{\text{bb}}\text{)}_2$, $\text{--SO}_2\text{R}^{\text{aa}}$, and $\text{--SO}_2\text{OR}^{\text{aa}}$, wherein R^{aa} and R^{bb} are as defined herein.

[0052] The term “sulfinyl” refers to the group $\text{--S(=O)R}^{\text{aa}}$, wherein R^{aa} is as defined herein.

[0053] The term “acyl” refers to a group having the general formula $\text{--C(=O)R}^{\text{X1}}$, $\text{--C(=O)OR}^{\text{X1}}$, $\text{--C(=O)--O--C(=O)R}^{\text{X1}}$, $\text{--C(=O)SR}^{\text{X1}}$, $\text{--C(=O)N(R}^{\text{X1}}\text{)}_2$, $\text{--C(=S)R}^{\text{X1}}$, $\text{--C(=S)N(R}^{\text{X1}}\text{)}_2$, $\text{--C(=S)O(R}^{\text{X1}}\text{)}$, $\text{--C(=S)S(R}^{\text{X1}}\text{)}$, $\text{--C(=NR}^{\text{X1}}\text{)R}^{\text{X1}}$, $\text{--C(=NR}^{\text{X1}}\text{)OR}^{\text{X1}}$, $\text{--C(=NR}^{\text{X1}}\text{)SR}^{\text{X1}}$, and $\text{--C(=NR}^{\text{X1}}\text{)N(R}^{\text{X1}}\text{)}_2$, wherein R^{X1} is hydrogen; halogen; substituted or unsubstituted hydroxyl; substituted or unsubstituted thiol; substituted or unsubstituted amino; substituted or unsubstituted acyl, cyclic or acyclic, substituted or unsubstituted, branched or unbranched aliphatic; cyclic or acyclic, substituted or unsubstituted, branched or unbranched heteroaliphatic; cyclic or acyclic, substituted or unsubstituted, branched or unbranched alkyl; cyclic or acyclic, substituted or unsubstituted, branched or unbranched alkenyl; substituted or unsubstituted alkynyl; substituted or unsubstituted aryl, substituted or unsubstituted heteroaryl, aliphaticoxy, heteroaliphaticoxy, alkyloxy, heteroalkyloxy, aryloxy, heteroaryloxy, aliphaticthioxy, heteroaliphaticthioxy, alkylthioxy, heteroalkylthioxy, arylthioxy, heteroarylthioxy, mono- or di- aliphaticamino, mono- or di- heteroaliphaticamino, mono- or di- alkylamino, mono- or di- heteroalkylamino, mono- or di-arylamino, or mono- or di-heteroarylamino; or two R^{X1} groups taken together form a 5- to 6-membered heterocyclic ring. Exemplary acyl groups include aldehydes (--CHO), carboxylic acids ($\text{--CO}_2\text{H}$), ketones, acyl halides, esters, amides, imines, carbonates, carbamates, and ureas. Acyl substituents include, but are not limited to, any of the substituents described herein, that result in the

formation of a stable moiety (*e.g.*, aliphatic, alkyl, alkenyl, alkynyl, heteroaliphatic, heterocyclic, aryl, heteroaryl, acyl, oxo, imino, thiooxo, cyano, isocyano, amino, azido, nitro, hydroxyl, thiol, halo, aliphaticamino, heteroaliphaticamino, alkylamino, heteroalkylamino, arylamino, heteroarylamino, alkylaryl, arylalkyl, aliphaticoxy, heteroaliphaticoxy, alkyloxy, heteroalkyloxy, aryloxy, heteroaryloxy, aliphaticthioxy, heteroaliphaticthioxy, alkylthioxy, heteroalkylthioxy, arylthioxy, heteroarylthioxy, acyloxy, and the like, each of which may or may not be further substituted).

[0054] The term “carbonyl” refers a group wherein the carbon directly attached to the parent molecule is sp^2 hybridized, and is substituted with an oxygen, nitrogen or sulfur atom, *e.g.*, a group selected from ketones (*e.g.*, $-C(=O)R^{aa}$), carboxylic acids (*e.g.*, $-CO_2H$), aldehydes ($-CHO$), esters (*e.g.*, $-CO_2R^{aa}$, $-C(=O)SR^{aa}$, $-C(=S)SR^{aa}$), amides (*e.g.*, $-C(=O)N(R^{bb})_2$, $-C(=O)NR^{bb}SO_2R^{aa}$, $-C(=S)N(R^{bb})_2$), and imines (*e.g.*, $-C(=NR^{bb})R^{aa}$, $-C(=NR^{bb})OR^{aa}$, $-C(=NR^{bb})N(R^{bb})_2$), wherein R^{aa} and R^{bb} are as defined herein.

[0055] The term “silyl” refers to the group $-Si(R^{aa})_3$, wherein R^{aa} is as defined herein.

[0056] The term “oxo” refers to the group $=O$, and the term “thiooxo” refers to the group $=S$.

[0057] Nitrogen atoms can be substituted or unsubstituted as valency permits, and include primary, secondary, tertiary, and quaternary nitrogen atoms. Exemplary nitrogen atom substituents include, but are not limited to, hydrogen, $-OH$, $-OR^{aa}$, $-N(R^{cc})_2$, $-CN$, $-C(=O)R^{aa}$, $-C(=O)N(R^{cc})_2$, $-CO_2R^{aa}$, $-SO_2R^{aa}$, $-C(=NR^{bb})R^{aa}$, $-C(=NR^{cc})OR^{aa}$, $-C(=NR^{cc})N(R^{cc})_2$, $-SO_2N(R^{cc})_2$, $-SO_2R^{cc}$, $-SO_2OR^{cc}$, $-SOR^{aa}$, $-C(=S)N(R^{cc})_2$, $-C(=O)SR^{cc}$, $-C(=S)SR^{cc}$, $-P(=O)(OR^{cc})_2$, $-P(=O)(R^{aa})_2$, $-P(=O)(N(R^{cc})_2)_2$, C_{1-10} alkyl, C_{1-10} perhaloalkyl, C_{2-10} alkenyl, C_{2-10} alkynyl, hetero C_{1-10} alkyl, hetero C_{2-10} alkenyl, hetero C_{2-10} alkynyl, C_{3-10} carbocyclyl, 3-14 membered heterocyclyl, C_{6-14} aryl, and 5-14 membered heteroaryl, or two R^{cc} groups attached to an N atom are joined to form a 3-14 membered heterocyclyl or 5-14 membered heteroaryl ring, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups, and wherein R^{aa} , R^{bb} , R^{cc} and R^{dd} are as defined above.

[0058] In certain embodiments, the substituent present on the nitrogen atom is an nitrogen protecting group (also referred to herein as an “amino protecting group”). Nitrogen protecting groups include, but are not limited to, $-OH$, $-OR^{aa}$, $-N(R^{cc})_2$, $-C(=O)R^{aa}$, $-C(=O)N(R^{cc})_2$, $-CO_2R^{aa}$, $-SO_2R^{aa}$, $-C(=NR^{cc})R^{aa}$, $-C(=NR^{cc})OR^{aa}$, $-C(=NR^{cc})N(R^{cc})_2$, $-SO_2N(R^{cc})_2$, $-SO_2R^{cc}$, $-SO_2OR^{cc}$, $-SOR^{aa}$, $-C(=S)N(R^{cc})_2$, $-C(=O)SR^{cc}$, $-C(=S)SR^{cc}$, C_{1-10} alkyl (*e.g.*, aralkyl, heteroaralkyl), C_{2-10} alkenyl, C_{2-10} alkynyl, hetero C_{1-10} alkyl, hetero C_{2-10} alkenyl, hetero C_{2-10} alkynyl, C_{3-10} carbocyclyl, 3-14 membered heterocyclyl, C_{6-14} aryl, and 5-14

membered heteroaryl groups, wherein each alkyl, alkenyl, alkynyl, heteroalkyl, heteroalkenyl, heteroalkynyl, carbocyclyl, heterocyclyl, aralkyl, aryl, and heteroaryl is independently substituted with 0, 1, 2, 3, 4, or 5 R^{dd} groups, and wherein R^{aa}, R^{bb}, R^{cc} and R^{dd} are as defined herein. Nitrogen protecting groups are well known in the art and include those described in detail in *Protecting Groups in Organic Synthesis*, T. W. Greene and P. G. M. Wuts, 3rd edition, John Wiley & Sons, 1999, incorporated herein by reference.

[0059] For example, nitrogen protecting groups such as amide groups (*e.g.*, $-\text{C}(=\text{O})\text{R}^{\text{aa}}$) include, but are not limited to, formamide, acetamide, chloroacetamide, trichloroacetamide, trifluoroacetamide, phenylacetamide, 3-phenylpropanamide, picolinamide, 3-pyridylcarboxamide, N-benzoylphenylalanyl derivative, benzamide, p-phenylbenzamide, o-nitrophenylacetamide, o-nitrophenoxycetamide, acetoacetamide, (N'-dithiobenzyloxyacylamino)acetamide, 3-(p-hydroxyphenyl)propanamide, 3-(o-nitrophenyl)propanamide, 2-methyl-2-(o-nitrophenoxycetamide), 2-methyl-2-(o-phenylazophenoxycetamide), 4-chlorobutanamide, 3-methyl-3-nitrobutanamide, o-nitrocinnamide, N-acetylmethionine derivative, o-nitrobenzamide and o-(benzyloxymethyl)benzamide.

[0060] Nitrogen protecting groups such as carbamate groups (*e.g.*, $-\text{C}(=\text{O})\text{OR}^{\text{aa}}$) include, but are not limited to, methyl carbamate, ethyl carbamate, 9-fluorenylmethyl carbamate (Fmoc), 9-(2-sulfo)fluorenylmethyl carbamate, 9-(2,7-dibromo)fluorenylmethyl carbamate, 2,7-di-t-butyl-[9-(10,10-dioxo-10,10,10,10-tetrahydrothioxanthyl)]methyl carbamate (DBD-Tmoc), 4-methoxyphenacyl carbamate (Phenoc), 2,2,2-trichloroethyl carbamate (Troc), 2-trimethylsilylethyl carbamate (Teoc), 2-phenylethyl carbamate (hZ), 1-(1-adamantyl)-1-methylethyl carbamate (Adpoc), 1,1-dimethyl-2-haloethyl carbamate, 1,1-dimethyl-2,2-dibromoethyl carbamate (DB-t-BOC), 1,1-dimethyl-2,2,2-trichloroethyl carbamate (TCBOC), 1-methyl-1-(4-biphenyl)ethyl carbamate (Bpoc), 1-(3,5-di-t-butylphenyl)-1-methylethyl carbamate (t-Bumeoc), 2-(2'- and 4'-pyridyl)ethyl carbamate (Pyoc), 2-(N,N-dicyclohexylcarboxamido)ethyl carbamate, t-butyl carbamate (BOC or Boc), 1-adamantyl carbamate (Adoc), vinyl carbamate (Voc), allyl carbamate (Alloc), 1-isopropylallyl carbamate (Ipaoc), cinnamyl carbamate (Coc), 4-nitrocinnamyl carbamate (Noc), 8-quinolyl carbamate, N-hydroxypiperidinyl carbamate, alkylthio carbamate, benzyl carbamate (Cbz), p-methoxybenzyl carbamate (Moz), p-nitrobenzyl carbamate, p-bromobenzyl carbamate, p-chlorobenzyl carbamate, 2,4-dichlorobenzyl carbamate, 4-methylsulfinylbenzyl carbamate (MsZ), 9-anthrylmethyl carbamate, diphenylmethyl carbamate, 2-methylthioethyl carbamate,

2-methylsulfonylethyl carbamate, 2-(p-toluenesulfonyl)ethyl carbamate, [2-(1,3-dithianyl)]methyl carbamate (Dmoc), 4-methylthiophenyl carbamate (Mtpc), 2,4-dimethylthiophenyl carbamate (Bmpc), 2-phosphonioethyl carbamate (Peoc), 2-triphenylphosphonioisopropyl carbamate (Ppoc), 1,1-dimethyl-2-cyanoethyl carbamate, m-chloro-p-acyloxybenzyl carbamate, p-(dihydroxyboryl)benzyl carbamate, 5-benzisoxazolylmethyl carbamate, 2-(trifluoromethyl)-6-chromonylmethyl carbamate (Tcroc), m-nitrophenyl carbamate, 3,5-dimethoxybenzyl carbamate, o-nitrobenzyl carbamate, 3,4-dimethoxy-6-nitrobenzyl carbamate, phenyl(o-nitrophenyl)methyl carbamate, t-amyl carbamate, S-benzyl thiocarbamate, p-cyanobenzyl carbamate, cyclobutyl carbamate, cyclohexyl carbamate, cyclopentyl carbamate, cyclopropylmethyl carbamate, p-decyloxybenzyl carbamate, 2,2-dimethoxyacetylvinyl carbamate, o-(N,N-dimethylcarboxamido)benzyl carbamate, 1,1-dimethyl-3-(N,N-dimethylcarboxamido)propyl carbamate, 1,1-dimethylpropynyl carbamate, di(2-pyridyl)methyl carbamate, 2-furanylmethyl carbamate, 2-iodoethyl carbamate, isoborynl carbamate, isobutyl carbamate, isonicotinyl carbamate, p-(p'-methoxyphenylazo)benzyl carbamate, 1-methylcyclobutyl carbamate, 1-methylcyclohexyl carbamate, 1-methyl-1-cyclopropylmethyl carbamate, 1-methyl-1-(3,5-dimethoxyphenyl)ethyl carbamate, 1-methyl-1-(p-phenylazophenyl)ethyl carbamate, 1-methyl-1-phenylethyl carbamate, 1-methyl-1-(4-pyridyl)ethyl carbamate, phenyl carbamate, p-(phenylazo)benzyl carbamate, 2,4,6-tri-t-butylphenyl carbamate, 4-(trimethylammonium)benzyl carbamate, and 2,4,6-trimethylbenzyl carbamate.

[0061] Nitrogen protecting groups such as sulfonamide groups (*e.g.*, $-S(=O)_2R^{aa}$) include, but are not limited to, *p*-toluenesulfonamide (Ts), benzenesulfonamide, 2,3,6-trimethyl-4-methoxybenzenesulfonamide (Mtr), 2,4,6-trimethoxybenzenesulfonamide (Mtb), 2,6-dimethyl-4-methoxybenzenesulfonamide (Pme), 2,3,5,6-tetramethyl-4-methoxybenzenesulfonamide (Mte), 4-methoxybenzenesulfonamide (Mbs), 2,4,6-trimethylbenzenesulfonamide (Mts), 2,6-dimethoxy-4-methylbenzenesulfonamide (iMDs), 2,2,5,7,8-pentamethylchroman-6-sulfonamide (Pmc), methanesulfonamide (Ms), β -trimethylsilylethanesulfonamide (SES), 9-anthracenesulfonamide, 4-(4',8'-dimethoxynaphthylmethyl)benzenesulfonamide (DNMBS), benzylsulfonamide, trifluoromethylsulfonamide, and phenacysulfonamide.

[0062] Other nitrogen protecting groups include, but are not limited to, phenothiazinyl-(10)-acyl derivative, N'-p-toluenesulfonylaminoacyl derivative, N'-phenylaminothioacyl derivative, N-benzoylphenylalanyl derivative, N-acetylmethionine derivative, 4,5-diphenyl-3-

oxazolin-2-one, N-phthalimide, N-dithiasuccinimide (Dts), N-2,3-diphenylmaleimide, N-2,5-dimethylpyrrole, N-1,1,4,4-tetramethyldisilylazacyclopentane adduct (STABASE), 5-substituted 1,3-dimethyl-1,3,5-triazacyclohexan-2-one, 5-substituted 1,3-dibenzyl-1,3,5-triazacyclohexan-2-one, 1-substituted 3,5-dinitro-4-pyridone, N-methylamine, N-allylamine, N-[2-(trimethylsilyl)ethoxy]methylamine (SEM), N-3-acetoxypyrrolamine, N-(1-isopropyl-4-nitro-2-oxo-3-pyrroline-3-yl)amine, quaternary ammonium salts, N-benzylamine, N-di(4-methoxyphenyl)methylamine, N-5-dibenzosuberylamine, N-triphenylmethylamine (Tr), N-[(4-methoxyphenyl)diphenylmethyl]amine (MMTr), N-9-phenylfluorenylamine (PhF), N-2,7-dichloro-9-fluorenylmethyleneamine, N-ferrocenylmethylamino (Fcm), N-2-picolylamino N'-oxide, N-1,1-dimethylthiomethyleneamine, N-benzylideneamine, N-p-methoxybenzylideneamine, N-diphenylmethyleneamine, N-[(2-pyridyl)mesityl]methyleneamine, N-(N',N'-dimethylaminomethylene)amine, N,N'-isopropylidenediamine, N-p-nitrobenzylideneamine, N-salicylideneamine, N-5-chlorosalicylideneamine, N-(5-chloro-2-hydroxyphenyl)phenylmethyleneamine, N-cyclohexylideneamine, N-(5,5-dimethyl-3-oxo-1-cyclohexenyl)amine, N-borane derivative, N-diphenylborinic acid derivative, N-[phenyl(pentaacylchromium- or tungsten)acyl]amine, N-copper chelate, N-zinc chelate, N-nitroamine, N-nitrosoamine, amine N-oxide, diphenylphosphinamide (Dpp), dimethylthiophosphinamide (Mpt), diphenylthiophosphinamide (Ppt), dialkyl phosphoramidates, dibenzyl phosphoramidate, diphenyl phosphoramidate, benzenesulfenamide, o-nitrobenzenesulfenamide (Nps), 2,4-dinitrobenzenesulfenamide, pentachlorobenzenesulfenamide, 2-nitro-4-methoxybenzenesulfenamide, triphenylmethylsulfenamide, and 3-nitropyridinesulfenamide (Npys). In certain embodiments, a nitrogen protecting group is benzyl (Bn), tert-butyloxycarbonyl (BOC), carbobenzyloxy (Cbz), 9-fluorenylmethyloxycarbonyl (Fmoc), trifluoroacetyl, triphenylmethyl, acetyl (Ac), benzoyl (Bz), p-methoxybenzyl (PMB), 3,4-dimethoxybenzyl (DMPM), p-methoxyphenyl (PMP), 2,2,2-trichloroethyloxycarbonyl (Troc), triphenylmethyl (Tr), tosyl (Ts), brosyl (Bs), nosyl (Ns), mesyl (Ms), triflyl (Tf), or dansyl (Ds).

[0063] In certain embodiments, the substituent present on an oxygen atom is an oxygen protecting group (also referred to herein as an "hydroxyl protecting group"). Oxygen protecting groups include, but are not limited to, $-R^{aa}$, $-N(R^{bb})_2$, $-C(=O)SR^{aa}$, $-C(=O)R^{aa}$, $-CO_2R^{aa}$, $-C(=O)N(R^{bb})_2$, $-C(=NR^{bb})R^{aa}$, $-C(=NR^{bb})OR^{aa}$, $-C(=NR^{bb})N(R^{bb})_2$, $-S(=O)R^{aa}$, $-SO_2R^{aa}$, $-Si(R^{aa})_3$, $-P(R^{cc})_2$, $-P(R^{cc})_3^+X^-$, $-P(OR^{cc})_2$, $-P(OR^{cc})_3^+X^-$, $-P(=O)(R^{aa})_2$, $-P(=O)(OR^{cc})_2$, and $-P(=O)(N(R^{bb})_2)_2$, wherein X^- , R^{aa} , R^{bb} , and R^{cc} are as defined herein.

Oxygen protecting groups are well known in the art and include those described in detail in *Protecting Groups in Organic Synthesis*, T. W. Greene and P. G. M. Wuts, 3rd edition, John Wiley & Sons, 1999, incorporated herein by reference.

[0064] Exemplary oxygen protecting groups include, but are not limited to, methyl, methoxymethyl (MOM), methylthiomethyl (MTM), t-butylthiomethyl, (phenyldimethylsilyl)methoxymethyl (SMOM), benzyloxymethyl (BOM), p-methoxybenzyloxymethyl (PMBM), (4-methoxyphenoxy)methyl (p-AOM), guaiacolmethyl (GUM), t-butoxymethyl, 4-pentenylloxymethyl (POM), siloxymethyl, 2-methoxyethoxymethyl (MEM), 2,2,2-trichloroethoxymethyl, bis(2-chloroethoxy)methyl, 2-(trimethylsilyl)ethoxymethyl (SEMOR), tetrahydropyranyl (THP), 3-bromotetrahydropyranyl, tetrahydrothiopyranyl, 1-methoxycyclohexyl, 4-methoxytetrahydropyranyl (MTHP), 4-methoxytetrahydrothiopyranyl, 4-methoxytetrahydrothiopyranyl S,S-dioxide, 1-[(2-chloro-4-methyl)phenyl]-4-methoxypiperidin-4-yl (CTMP), 1,4-dioxan-2-yl, tetrahydrofuran-2-yl, tetrahydrothiofuran-2-yl, 2,3,3a,4,5,6,7,7a-octahydro-7,8,8-trimethyl-4,7-methanobenzofuran-2-yl, 1-ethoxyethyl, 1-(2-chloroethoxy)ethyl, 1-methyl-1-methoxyethyl, 1-methyl-1-benzyloxyethyl, 1-methyl-1-benzyloxy-2-fluoroethyl, 2,2,2-trichloroethyl, 2-trimethylsilylethyl, 2-(phenylselenyl)ethyl, t-butyl, allyl, p-chlorophenyl, p-methoxyphenyl, 2,4-dinitrophenyl, benzyl (Bn), p-methoxybenzyl, 3,4-dimethoxybenzyl, o-nitrobenzyl, p-nitrobenzyl, p-halobenzyl, 2,6-dichlorobenzyl, p-cyanobenzyl, p-phenylbenzyl, 2-picolyl, 4-picolyl, 3-methyl-2-picolyl N-oxido, diphenylmethyl, p,p'-dinitrobenzhydryl, 5-dibenzosuberone, triphenylmethyl, α -naphthylmethyl, p-methoxyphenyldiphenylmethyl, di(p-methoxyphenyl)phenylmethyl, tri(p-methoxyphenyl)methyl, 4-(4'-bromophenacyloxyphenyl)diphenylmethyl, 4,4',4''-tris(4,5-dichlorophthalimidophenyl)methyl, 4,4',4''-tris(levulinoyloxyphenyl)methyl, 4,4',4''-tris(benzoyloxyphenyl)methyl, 3-(imidazol-1-yl)bis(4',4''-dimethoxyphenyl)methyl, 1,1-bis(4-methoxyphenyl)-1'-pyrenylmethyl, 9-anthryl, 9-(9-phenyl)xanthenyl, 9-(9-phenyl-10-oxo)anthryl, 1,3-benzodithiolan-2-yl, benzoisothiazolyl S,S-dioxido, trimethylsilyl (TMS), triethylsilyl (TES), triisopropylsilyl (TIPS), dimethylisopropylsilyl (IPDMS), diethylisopropylsilyl (DEIPS), dimethylhexylsilyl, t-butyltrimethylsilyl (TBDMS), t-butyltriphenylsilyl (TBDPS), tribenzylsilyl, tri-p-xylylsilyl, triphenylsilyl, diphenylmethylsilyl (DPMS), t-butylmethoxyphenylsilyl (TBMPS), formate, benzoylformate, acetate, chloroacetate, dichloroacetate, trichloroacetate, trifluoroacetate, methoxyacetate, triphenylmethoxyacetate, phenoxyacetate, p-chlorophenoxyacetate, 3-phenylpropionate, 4-

oxopentanoate (levulinate), 4,4-(ethylenedithio)pentanoate (levulinoyldithioacetal), pivaloate, adamantate, crotonate, 4-methoxycrotonate, benzoate, *p*-phenylbenzoate, 2,4,6-trimethylbenzoate (mesitoate), methyl carbonate, 9-fluorenylmethyl carbonate (Fmoc), ethyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-(trimethylsilyl)ethyl carbonate (TMSEC), 2-(phenylsulfonyl) ethyl carbonate (Psec), 2-(triphenylphosphonio) ethyl carbonate (Peoc), isobutyl carbonate, vinyl carbonate, allyl carbonate, *t*-butyl carbonate (BOC or Boc), *p*-nitrophenyl carbonate, benzyl carbonate, *p*-methoxybenzyl carbonate, 3,4-dimethoxybenzyl carbonate, *o*-nitrobenzyl carbonate, *p*-nitrobenzyl carbonate, *S*-benzyl thiocarbonate, 4-ethoxy-1-naphthyl carbonate, methyl dithiocarbonate, 2-iodobenzoate, 4-azidobutyrate, 4-nitro-4-methylpentanoate, *o*-(dibromomethyl)benzoate, 2-formylbenzenesulfonate, 2-(methylthiomethoxy)ethyl, 4-(methylthiomethoxy)butyrate, 2-(methylthiomethoxymethyl)benzoate, 2,6-dichloro-4-methylphenoxyacetate, 2,6-dichloro-4-(1,1,3,3-tetramethylbutyl)phenoxyacetate, 2,4-bis(1,1-dimethylpropyl)phenoxyacetate, chlorodiphenylacetate, isobutyrate, monosuccinoate, (E)-2-methyl-2-butenolate, *o*-(methoxyacyl)benzoate, α -naphthoate, nitrate, alkyl N,N,N',N'-tetramethylphosphorodiamidate, alkyl N-phenylcarbamate, borate, dimethylphosphinothioyl, alkyl 2,4-dinitrophenylsulfonate, sulfate, methanesulfonate (mesylate), benzyisulfonate, and tosylate (Ts). In certain embodiments, an oxygen protecting group is silyl. In certain embodiments, an oxygen protecting group is *t*-butyldiphenylsilyl (TBDPS), *t*-butyldimethylsilyl (TBDMS), triisopropylsilyl (TIPS), triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranyl (THF), *p*-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), allyl, *p*-methoxybenzyl (PMB, MPM), *t*-butyl, benzyl (Bn), allyl, or pivaloyl (Piv).

[0065] In certain embodiments, the substituent present on a sulfur atom is a sulfur protecting group (also referred to as a “thiol protecting group”). Sulfur protecting groups include, but are not limited to, $-R^{aa}$, $-N(R^{bb})_2$, $-C(=O)SR^{aa}$, $-C(=O)R^{aa}$, $-CO_2R^{aa}$, $-C(=O)N(R^{bb})_2$, $-C(=NR^{bb})R^{aa}$, $-C(=NR^{bb})OR^{aa}$, $-C(=NR^{bb})N(R^{bb})_2$, $-S(=O)R^{aa}$, $-SO_2R^{aa}$, $-Si(R^{aa})_3$, $-P(R^{cc})_2$, $-P(R^{cc})_3^+X^-$, $-P(OR^{cc})_2$, $-P(OR^{cc})_3^+X^-$, $-P(=O)(R^{aa})_2$, $-P(=O)(OR^{cc})_2$, and $-P(=O)(N(R^{bb})_2)_2$, wherein R^{aa} , R^{bb} , and R^{cc} are as defined herein. Sulfur protecting groups are well known in the art and include those described in detail in Protecting Groups in

Organic Synthesis, T. W. Greene and P. G. M. Wuts, 3rd edition, John Wiley & Sons, 1999, incorporated herein by reference. In certain embodiments, a sulfur protecting group is acetamidomethyl, *t*-butyl, 3-nitro-2-pyridine sulfenyl, 2-pyridine-sulfenyl, or triphenylmethyl.

[0066] A “counterion” or “anionic counterion” is a negatively charged group associated with a positively charged group in order to maintain electronic neutrality. An anionic counterion may be monovalent (*i.e.*, including one formal negative charge). An anionic counterion may also be multivalent (*i.e.*, including more than one formal negative charge), such as divalent or trivalent. Exemplary counterions include halide ions (*e.g.*, F[−], Cl[−], Br[−], I[−]), NO₃[−], ClO₄[−], OH[−], H₂PO₄[−], HCO₃[−], HSO₄[−], sulfonate ions (*e.g.*, methanesulfonate, trifluoromethanesulfonate, *p*-toluenesulfonate, benzenesulfonate, 10-camphor sulfonate, naphthalene-2-sulfonate, naphthalene-1-sulfonic acid-5-sulfonate, ethan-1-sulfonic acid-2-sulfonate, and the like), carboxylate ions (*e.g.*, acetate, propanoate, benzoate, glycerate, lactate, tartrate, glycolate, gluconate, and the like), BF₄[−], PF₄[−], PF₆[−], AsF₆[−], SbF₆[−], B[3,5-(CF₃)₂C₆H₃]₄[−], B(C₆F₅)₄[−], BPh₄[−], Al(OC(CF₃)₃)₄[−], and carborane anions (*e.g.*, CB₁₁H₁₂[−] or (HCB₁₁Me₅Br₆)[−]). Exemplary counterions which may be multivalent include CO₃^{2−}, HPO₄^{2−}, PO₄^{3−}, B₄O₇^{2−}, SO₄^{2−}, S₂O₃^{2−}, carboxylate anions (*e.g.*, tartrate, citrate, fumarate, maleate, malate, malonate, gluconate, succinate, glutarate, adipate, pimelate, suberate, azelate, sebacate, salicylate, phthalates, aspartate, glutamate, and the like), and carboranes.

[0067] The term “leaving group” is given its ordinary meaning in the art of synthetic organic chemistry and refers to an atom or a group capable of being displaced by a nucleophile. See, for example, Smith, *March Advanced Organic Chemistry* 6th ed. (501-502). Examples of suitable leaving groups include, but are not limited to, halogen (such as F, Cl, Br, or I (iodine)), alkoxycarbonyloxy, aryloxy carbonyloxy, alkanesulfonyloxy, arenesulfonyloxy, alkyl-carbonyloxy (*e.g.*, acetoxy), arylcarbonyloxy, aryloxy, methoxy, *N,O*-dimethylhydroxylamino, pixyl, and haloformates. In some cases, the leaving group is a sulfonic acid ester, such as toluenesulfonate (tosylate, -OTs), methanesulfonate (mesylate, -OMs), *p*-bromobenzenesulfonyloxy (brosylate, -OBs), -OS(=O)₂(CF₂)₃CF₃ (nonaflate, -ONf), or trifluoromethanesulfonate (triflate, -OTf). In some cases, the leaving group is a brosylate, such as *p*-bromobenzenesulfonyloxy. In some cases, the leaving group is a nosylate, such as 2-nitrobenzenesulfonyloxy. The leaving group may also be a phosphineoxide (*e.g.*, formed during a Mitsunobu reaction) or an internal leaving group such as an epoxide or cyclic sulfate. Other non-limiting examples of leaving groups are water, ammonia, alcohols, ether moieties, thioether moieties, zinc halides, magnesium moieties, diazonium salts, and copper

moieties. Further exemplary leaving groups include, but are not limited to, halo (*e.g.*, chloro, bromo, iodo) and activated substituted hydroxyl groups (*e.g.*, $-\text{OC}(=\text{O})\text{SR}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OCO}_2\text{R}^{\text{aa}}$, $-\text{OC}(=\text{O})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{R}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{OR}^{\text{aa}}$, $-\text{OC}(=\text{NR}^{\text{bb}})\text{N}(\text{R}^{\text{bb}})_2$, $-\text{OS}(=\text{O})\text{R}^{\text{aa}}$, $-\text{OSO}_2\text{R}^{\text{aa}}$, $-\text{OP}(\text{R}^{\text{cc}})_2$, $-\text{OP}(\text{R}^{\text{cc}})_3$, $-\text{OP}(=\text{O})_2\text{R}^{\text{aa}}$, $-\text{OP}(=\text{O})(\text{R}^{\text{aa}})_2$, $-\text{OP}(=\text{O})(\text{OR}^{\text{cc}})_2$, $-\text{OP}(=\text{O})_2\text{N}(\text{R}^{\text{bb}})_2$, and $-\text{OP}(=\text{O})(\text{NR}^{\text{bb}})_2$, wherein R^{aa} , R^{bb} , and R^{cc} are as defined herein).

[0068] As used herein, use of the phrase “at least one instance” refers to 1, 2, 3, 4, or more instances, but also encompasses a range, *e.g.*, for example, from 1 to 4, from 1 to 3, from 1 to 2, from 2 to 4, from 2 to 3, or from 3 to 4 instances, inclusive.

[0069] A “non-hydrogen group” refers to any group that is defined for a particular variable that is not hydrogen.

[0070] The following definitions are more general terms used throughout the present application.

[0071] As used herein, the term “salt” refers to any and all salts, and encompasses pharmaceutically acceptable salts. The term “pharmaceutically acceptable salt” refers to those salts which are, within the scope of sound medical judgment, suitable for use in contact with the tissues of humans and lower animals without undue toxicity, irritation, allergic response, and the like, and are commensurate with a reasonable benefit/risk ratio. Pharmaceutically acceptable salts are well known in the art. For example, Berge *et al.* describe pharmaceutically acceptable salts in detail in *J. Pharmaceutical Sciences*, 1977, 66, 1-19, incorporated herein by reference. Pharmaceutically acceptable salts of the compounds of this invention include those derived from suitable inorganic and organic acids and bases.

Examples of pharmaceutically acceptable, nontoxic acid addition salts are salts of an amino group formed with inorganic acids, such as hydrochloric acid, hydrobromic acid, phosphoric acid, sulfuric acid, and perchloric acid or with organic acids, such as acetic acid, oxalic acid, maleic acid, tartaric acid, citric acid, succinic acid, or malonic acid or by using other methods known in the art such as ion exchange. Other pharmaceutically acceptable salts include adipate, alginate, ascorbate, aspartate, benzenesulfonate, benzoate, bisulfate, borate, butyrate, camphorate, camphorsulfonate, citrate, cyclopentanepropionate, digluconate, dodecylsulfate, ethanesulfonate, formate, fumarate, glucoheptonate, glycerophosphate, gluconate, hemisulfate, heptanoate, hexanoate, hydroiodide, 2-hydroxy-ethanesulfonate, lactobionate, lactate, laurate, lauryl sulfate, malate, maleate, malonate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, nitrate, oleate, oxalate, palmitate, pamoate, pectinate, persulfate, 3-phenylpropionate, phosphate, picrate, pivalate, propionate, stearate, succinate,

sulfate, tartrate, thiocyanate, p-toluenesulfonate, undecanoate, valerate salts, and the like. Salts derived from appropriate bases include alkali metal, alkaline earth metal, ammonium, and $N^+(C_{1-4} \text{ alkyl})_4$ salts. Representative alkali or alkaline earth metal salts include sodium, lithium, potassium, calcium, magnesium, and the like. Further pharmaceutically acceptable salts include, when appropriate, nontoxic ammonium, quaternary ammonium, and amine cations formed using counterions such as halide, hydroxide, carboxylate, sulfate, phosphate, nitrate, lower alkyl sulfonate, and aryl sulfonate.

[0072] It is also to be understood that compounds that have the same molecular formula but differ in the nature or sequence of bonding of their atoms or the arrangement of their atoms in space are termed “isomers”. Isomers that differ in the arrangement of their atoms in space are termed “stereoisomers”.

[0073] Stereoisomers that are not mirror images of one another are termed “diastereomers” and those that are non-superimposable mirror images of each other are termed “enantiomers”. When a compound has an asymmetric center, for example, it is bonded to four different groups, a pair of enantiomers is possible. An enantiomer can be characterized by the absolute configuration of its asymmetric center and is described by the R- and S-sequencing rules of Cahn and Prelog, or by the manner in which the molecule rotates the plane of polarized light and designated as dextrorotatory or levorotatory (*i.e.*, as (+) or (–)-isomers respectively). A chiral compound can exist as either individual enantiomer or as a mixture thereof. A mixture containing equal proportions of the enantiomers is called a “racemic mixture”.

[0074] The term “small molecule” refers to molecules, whether naturally-occurring or artificially created (*e.g.*, via chemical synthesis) that have a relatively low molecular weight. Typically, a small molecule is an organic compound (*i.e.*, it contains carbon). The small molecule may contain multiple carbon-carbon bonds, stereocenters, and other functional groups (*e.g.*, amines, hydroxyl, carbonyls, and heterocyclic rings, *etc.*). In certain embodiments, the molecular weight of a small molecule is not more than about 1,000 g/mol, not more than about 900 g/mol, not more than about 800 g/mol, not more than about 700 g/mol, not more than about 600 g/mol, not more than about 500 g/mol, not more than about 400 g/mol, not more than about 300 g/mol, not more than about 200 g/mol, or not more than about 100 g/mol. In certain embodiments, the molecular weight of a small molecule is at least about 100 g/mol, at least about 200 g/mol, at least about 300 g/mol, at least about 400 g/mol, at least about 500 g/mol, at least about 600 g/mol, at least about 700 g/mol, at least about 800 g/mol, or at least about 900 g/mol, or at least about 1,000 g/mol. Combinations of the above ranges (*e.g.*, at least about 200 g/mol and not more than about 500 g/mol) are also possible. In

certain embodiments, the small molecule is a therapeutically active agent such as a drug (*e.g.*, a molecule approved by the U.S. Food and Drug Administration as provided in the Code of Federal Regulations (C.F.R.)).

[0075] The term “catalysis,” “catalyze,” or “catalytic” refers to the increase in rate of a chemical reaction due to the participation of a substance called a “catalyst.” In certain embodiments, the amount and nature of a catalyst remains essentially unchanged during a reaction. In certain embodiments, a catalyst is regenerated, or the nature of a catalyst is essentially restored after a reaction. A catalyst may participate in multiple chemical transformations. The effect of a catalyst may vary due to the presence of other substances known as inhibitors or poisons (which reduce the catalytic activity) or promoters (which increase the activity). Catalyzed reactions have lower activation energy (rate-limiting free energy of activation) than the corresponding uncatalyzed reaction, resulting in a higher reaction rate at the same temperature. Catalysts may affect the reaction environment favorably, bind to the reagents to polarize bonds, form specific intermediates that are not typically produced by a uncatalyzed reaction, or cause dissociation of reagents to reactive forms.

[0076] The term “solvent” refers to a substance that dissolves one or more solutes, resulting in a solution. A solvent may serve as a medium for any reaction or transformation described herein. The solvent may dissolve one or more reactants or reagents in a reaction mixture. The solvent may facilitate the mixing of one or more reagents or reactants in a reaction mixture. The solvent may also serve to increase or decrease the rate of a reaction relative to the reaction in a different solvent. Solvents can be polar or non-polar, protic or aprotic. Common organic solvents useful in the methods described herein include, but are not limited to, acetone, acetonitrile, benzene, benzonitrile, 1-butanol, 2-butanone, butyl acetate, *tert*-butyl methyl ether, carbon disulfide carbon tetrachloride, chlorobenzene, 1-chlorobutane, chloroform, cyclohexane, cyclopentane, 1,2-dichlorobenzene, 1,2-dichloroethane, dichloromethane (DCM), *N,N*-dimethylacetamide *N,N*-dimethylformamide (DMF), 1,3-dimethyl-3,4,5,6-tetrahydro-2-pyrimidinone (DMPU), 1,4-dioxane, 1,3-dioxane, diethylether, 2-ethoxyethyl ether, ethyl acetate, ethyl alcohol, ethylene glycol, dimethyl ether, heptane, *n*-hexane, hexanes, hexamethylphosphoramide (HMPA), 2-methoxyethanol, 2-methoxyethyl acetate, methyl alcohol, 2-methylbutane, 4-methyl-2-pentanone, 2-methyl-1-propanol, 2-methyl-2-propanol, 1-methyl-2-pyrrolidinone, dimethylsulfoxide (DMSO), nitromethane, 1-octanol, pentane, 3-pentanone, 1-propanol, 2-propanol, pyridine, tetrachloroethylene, tetrahyrdofuran (THF), 2-methyltetrahydrofuran, toluene, trichlorobenzene, 1,1,2-

trichlorotrifluoroethane, 2,2,4-trimethylpentane, trimethylamine, triethylamine, *N,N*-diisopropylethylamine, diisopropylamine, water, *o*-xylene, and *p*-xylene.

BRIEF DESCRIPTION OF THE DRAWINGS

[0077] The accompanying drawings, which constitute a part of this specification, illustrate several embodiments of the invention and together with the description, serve to explain the principles of the invention.

[0078] *Figure 1* shows the structures of halichondrin A, B, and C; homohalichondrin A, B, and C; and norhalichondrin A, B, and C.

[0079] *Figure 2A* shows an example of a Ni/Zr-mediated ketolization. *Figure 2B* shows an example of a Ni-catalyzed ketone coupling. *Figure 2C* shows feasibility studies under three variations of Ni-mediated one-pot ketone coupling.

[0080] *Figure 3A* shows proposed catalytic cycles for the Ni/Zr-mediated ketolization provided herein. *Figure 3B* shows exemplary coupling with common radical probes.

[0081] *Figure 4* shows one-pot ketone coupling with nucleophiles bearing a α -OR and other functional groups. Reaction conditions: **1-5** (1.0 equiv.), **1-7** (1.2 equiv.), NiBr₂•(dtbbpy) (5 mol%).

[0082] *Figure 5A* shows examples of Ni/Zr-ketolization reaction, and *Figure 5B* shows a further example.

[0083] *Figure 5C* shows results of nickel ligand screening experiments. *Figure 5D* shows NiBr₂, NiCl₂, and NiI₂ comparison experiments. *Figure 5E* shows the results of solvent screening experiments. *Figure 5F* shows the results of co-solvent screening experiments. *Figure 5G* shows additive screening experiments. *Figure 5H* shows screening of zirconium equivalents. *Figure 5I* shows studies with various electrophiles. *Figure 5J* shows reducing reagent screening experiments. *Figure 5K* shows concentration studies. *Figure 5L* shows substrate ratio experiments.

[0084] *Figure 6* shows potential routes to halichondrins and analogs thereof.

[0085] *Figure 7* shows the Ni/Zr-ketolization provided herein applied to the synthesis of a halichondrin analog. Reagents and conditions: (a) **2-5** (1.0 equiv.), **2-6** (1.3 equiv.), NiBr₂•(dtbbpy) (30 mol%), Cp₂ZrCl₂ (3 equiv.), (*t*-Bu)₂(Me)Py (4 equiv.), Zn (6 equiv.) in 5:1 DMI-EtOAc (*C* 0.1 M), rt. (b) HF•Py (20 equiv.), THF, followed by TBAF (4 equiv.), pivalic acid (2 equiv.), DMF, rt. (c) PPTS (5 equiv.), CH₂Cl₂, ~20 °C, 2 hr. Abbreviation: TES = Et₃Si-; SPy-2: 2-thiopyridine; DMI: 1,3-dimethyl-2-imidazolidinone; TBAF: tetrabutylammonium fluoride; PPTS: pyridinium *p*-toluenesulfonate.

[0086] *Figure 8A* shows exemplary right- and left-halves of halichondrins, homohalichondrins, and norhalichondrins. *Figure 8B* shows an exemplary synthesis of halichondrins. Reagents and conditions: For all the cases, step #1 was ketone coupling under the conditions specified in Scheme 3; step #2 was TBAF (10 equiv.), pivalic acid (5 equiv.), DMF, rt, 3-8 hr; step #3 was PPTS, CH₂Cl₂, ~20 °C, 2-4 hours. Epimerization of C38-*epi*-halichondrins was done with TMSOTf, CH₂Cl₂, -78 °C. For the halichondrin-A or -C series, these steps were followed by PPTS, 2,2-dimethylpropan-1,3-diol, *i*-PrOH, rt, overnight or by Pd(PPh₃)₄, dimedone, CH₂Cl₂, rt, 4-8 hours, respectively. In the norhalichondrin series, the methyl ester at C53 was hydrolyzed by treatment with aq. LiOH, THF, rt, at the end of transformation. Numbers after i and ii indicate the yield for ketone couplings and overall yield after ketone coupling, respectively.

[0087] *Figure 9A* shows an exemplary synthesis of the C27-C37 building block. Reagents and Conditions: *a.* 1. LiBH₄, Et₂O, 0 °C (~100%). 2. TES-Cl, imidazole, CH₂Cl₂, rt (~100%). 3. Swern oxidation (see, *e.g.*, Rodriguez, A.; Nomen, M.; Spur, B. W.; Godfroid, J. J. *Tetrahedron Lett.* **1999**, *40*, 5161); *b.* 1. Cr-catalyst prepared from (*S*)-**4-E** (10 mol%), (Me)₂Phen-(OMe)₂•NiCl₂ (2 mol%), LiCl (2 equiv.), Mn (excess), Cp₂ZrCl₂ (1.1 equiv.), 2,6-lutidine (1 equiv.), MeCN (C 0.4 M), rt, 1 hour (93% for 2 steps; *dr* = 19:1). 2. MPMO(=NH)CCl₃, La(OTf)₃, toluene, rt, 6 hours. 3. *p*-TsOH (cat.), MeOH-CH₂Cl₂, rt, 4 hours (88% for 2 steps). *c.* 1. K₃PO₄ (1 equiv.), 18-Crown-6 (3 equiv.), toluene (79%). 2. DIBAL, CH₂Cl₂, -78 °C, 1.5 hours (94%). Abbreviation: 18-Crown-6 = 1,4,7,10,13,16-hexaoxacyclooctadecane; DIBAL = diisobutylaluminum hydride; *p*-TsOH = *p*-toluenesulfonic acid. *Figure 9B* shows exemplary sulfonamide ligands and nickel complexes useful in the Ni/Cr coupling reactions provided herein.

[0088] *Figure 10A* shows exemplary synthesis of C20-C37 building block. Reagents and Conditions: *a.* 1. Cr-catalyst prepared from (*R*)-**4-F** (10 mol%), (Et)₂Phen•NiCl₂ (2 mol%), LiCl (2 equiv.), Mn (excess), Cp₂ZrCl₂ (1 equiv.), MeCN (C 0.3 M), rt, 3 hours. 2. TBAF (2 equiv.), AcOH (0.6 equiv.), THF, 0 °C→rt (79% for 2 steps). 3. TES-H (10 equiv.), TEOTf (5 equiv.), CH₂Cl₂, 0 °C, 3 hours (87%). 4. 2,2-dimethoxypropane (3 equiv.), acetone, 0 °C→rt. *b.* DIBAL, CH₂Cl₂, -78 °C, 1.5 hours (89% for 2 steps). Abbreviation: MPM = *p*-MeOC₆H₄CH₂-; TES = Et₃Si-. *Figure 10B* shows Analysis on stereochemical course of reductive cyclization: desired and undesired series.

[0089] *Figure 11* shows exemplary synthesis of the C1-C37 building block in the halichondrin B series. Reagents and Conditions: *a.* 1. Cr-catalyst prepared from (*S*)-**4-G** (10

mol%), (Et)₂Phen•NiCl₂ (2 mol%), LiCl (2 equiv.), Mn (excess), ZrCp₂Cl₂ (2.5 equiv.), 2,6-di-*t*-butyl-4-methylpyridine (2.5 equiv.), MeCN (C 0.x M), rt, 2 hours. 2. K₂CO₃ (10 equiv.), 60 °C, 16 hr, then add H₂O (1/10 volume of MeOH), 60 °C, 3 hours. *b.* 2-methyl-6-nitrobenzoic anhydride (6 equiv.), 4-dimethylaminopyridine (12 equiv.), *i*-Pr₂NEt (6 equiv.), toluene, 70 °C (syringe pump; 73% for 3 steps). *c.* 1. *p*-TsOH, MeOH, rt, 1 hour. 2. Tf₂O (1.2 equiv.), 2,6-lutidine (5 equiv.), CH₂Cl₂, -78 °C, 15 min, followed by addition of TESOTf (1.5 equiv.), -78 °C→0 °C, then followed by addition of NaI (5 equiv.) in DMF, rt, 2.5 hours (94% for steps). Abbreviation: TES = Et₃Si-; *p*-TsOH = *p*-toluenesulfonic acid.

[0090] Figure 12 shows the X-ray structure of C35/C37-Diol of **4-10-B**.

[0091] Figure 13 shows exemplary synthesis of the C1-C37 building block in the halichondrin A series. Reagents and conditions: *a.* 1. Ac₂O, py, rt. 2. CSA, CH₂Cl₂-MeOH, rt. 3. TBSOTf, 2,6-lutidine, CH₂Cl₂, -78 °C, 1 hour (92% for 3 steps). 4. DIBAL, CH₂Cl₂, -78 °C, 1 hour (88%). Follow the synthetic sequence under the conditions defined in Figure 11, except that (Me)₆PhenNiCl₂ (2 mol%) was used for the Ni/Cr-mediated coupling. The overall yield from bis-TBS-**4-8** to **4-12-A** was 40.8%, which was good compared with the overall yield in the halichondrin B series. Abbreviation: TBS = *t*BuMe₂Si-; CSA = camphorsulfonic acid.

[0092] Figure 14A shows an exemplary synthesis of the C1-C37 building block in the halichondrin-C series. Reagents and conditions: *a.* Follow the synthetic sequence under the conditions defined in Figure 11. The overall yield from **4-8** to **4-12-C** was 54.2%, which was good compared with the overall yield in the halichondrin B series. Figure 14B shows an X-ray structure of the product.

[0093] Figure 15 shows exemplary stereocontrolled [6,6]-spiroketal synthesis. Abbreviation: MPM = *p*-MeOC₆H₄CH₂-.

[0094] Figure 16 shows exemplary synthesis of a left half of halichondrin analogs. Reagents and conditions: *a.* 1. TBSOTf (2.5 equiv.), Et₃N (5 equiv.), CH₂Cl₂, 0 °C~rt, 3 hours. 2. NH₄Cl aq., EtOAc, THF, 50 °C, 3 hours (100% for 2 steps). *b.* 1. DIBAL (1.3 equiv.), CH₂Cl₂, -78 °C, 40 minutes. 2. MePPh₃Br (4 equiv.), *t*-BuOK (3 equiv.), THF, 0 °C~rt, 1.5 hours (96% for 2 steps). 3. 9-BBN (2.5 eq.), THF, rt, 1.5 hours then NaBO₃•H₂O aq. 4. TEMPO (10 mol%), PhI(OAc)₂ (3 equiv.), NaHCO₃ (10 equiv.), 4 °C, 15 hours (97% for 2 steps). *c.* **5** (1.4 equiv.), *t*-BuLi (2.6 equiv.), THF, -78 °C, 15 min (90%). *d.* 1. OsO₄ (10 mol%), NMMO (2 equiv.), H₂O, acetone, rt, 21 hours. 2. Pb(OAc)₄ (1.2 equiv.), K₂CO₃ (3 equiv.), CH₂Cl₂, rt, 1 hour (83% for 2 steps). 3. (MeO)₂P(=O)CH₂CO₂Bn (4 equiv.), K₃PO₄

(3 equiv.), rt, 23 hours. *e.* LiBr (10 equiv.), DBU (5 equiv.), BnOAc (10 equiv.), MeCN, rt, 12 hr, 2. DDQ (2 equiv.), CH₂Cl₂, pH 7 buffer, rt, 40 min (75% for 3 steps). 3. TESCl (2 equiv.), imidazole (4 equiv.), CH₂Cl₂, rt, 16 hours. 4. H₂ (1 atm), Pd/C, EtOAc, rt, 45 min, 5. (PyS)₂ (1.4 equiv.), PPh₃ (1.2 equiv.), CH₂Cl₂, rt, 17 hr (96% for 3 steps). Abbreviation: DIBAL = diisobutylaluminium hydride; 9-BBN = 9-borabicyclononane; TEMPO = 2,2,6,6-tetramethyl-1-piperidinyloxy; NMMO or NMO = 4-methylmorpholine *N*-oxide; DBU = 1,8-diazabicyclo[5.4.0]undec-7-ene; DDQ = 2,3-dichloro-5,6-dicyano-*p*-benzoquinone.

[0095] *Figure 17* shows an exemplary synthesis of a left hand building block of halichondrins. Reagents and conditions: *a.* **10** (1.8 equiv.), *n*-BuLi (1.75 equiv.), Li(thienylCuCN) (2.0 equiv.), BF₃•Et₂O (1.6 equiv.), Et₂O, -78 °C, 1 hour (81%). *b.* 1. VO(TMHD)₂ (5 mol%), *t*BuOOH (5.5 M in decane, 2 equiv.), toluene, rt, 5 hours. 2. TESCl (2.0 equiv.), imidazole (4.0 equiv.), CH₂Cl₂, 0 °C, 2 hr (85% for 2 steps). *c.* *t*-BuLi (2.6 equiv.), THF, -78 °C, 0.5 hours (85%). *d.* 1. (PhO)₂P(=O)OH (5 mol%), toluene (0.05M), 0 °C to rt, 12 hours. 2. TESCl (3.0 equiv.), imidazole (6.0 equiv.), CH₂Cl₂, rt, 2 hours (85% for 2 steps). *e.* 1. OsO₄ (5% mol), NMMO (2.0 equiv.), acetone/H₂O, rt, 12 hours. 2. Pb(OAc)₄ (1.5 equiv.), K₂CO₃ (10 equiv.), CH₂Cl₂, rt, 10 minutes. 3. (MeO)₂P(=O)COOBn (4 equiv.), K₃PO₄ (8 equiv.), toluene, rt, 15 hours (82% for 3 steps). 4. (PhO)₂P(=O)OH (5 mol%), THF-H₂O (4:1, 0.02M), rt, 24 hours. 5. TBSCl (1.5 equiv.), imidazole (3.0 equiv.), CH₂Cl₂, rt, 2 hours (80% for 2 steps). *f.* BnOAc (1 equiv.), and LiCl (10 equiv.), DBU (20 equiv.), MeCN (0.05M), 24 hr (86% alone with 8% **18**). or BnOAc (1 equiv.), and LiCl (10 equiv.), DBU (20 equiv.), **M** (50 mol%), MeCN (0.05M), 2 hours; then BnOAc (1 equiv.), and LiCl (10 equiv.), DBU (20 equiv.), MeCN (0.05M), 24 hours (93%). *g.* 1. DDQ (1.6 equiv.), CH₂Cl₂, phosphate buffer, 0 °C, 0.5 hours. 2. TESCl (3 equiv.), imidazole (6 equiv.), CH₂Cl₂, rt, 2 hours (90% for 2 steps). 3. Pd/C, H₂ balloon, EtOAc, rt, 1 hour. 4. (PyS)₂ (1.4 equiv.), PPh₃ (1.3 equiv.), toluene, rt, 3 hr (91% for 2 steps). Abbreviation: TMHD = tris(2,2,6,6-tetramethyl-3,5-heptanedionate).

[0096] *Figure 18* shows an exemplary synthesis of a left half building block in the homohalichondrin series. *Reagents and conditions:* *a.* 1. DIBAL (1.3 equiv.), CH₂Cl₂, -78 °C, 15 min. 2. MePPh₃Br (4 equiv.), *t*-BuOK (3 equiv.), THF, 0 °C-rt, 20 minutes. 3. TBSOTf (1.3 equiv.), 2,6-lutidine (2 equiv.), CH₂Cl₂, 0 °C-rt, 1 hour. 4. HF•py (ca. 8 equiv.), pyridine, MeCN, -10 °C-rt, 1.5 hours (96% for 4 steps). *b.* 1. Tf₂O (1.2 equiv.), 2,6-lutidine (4 equiv.), CH₂Cl₂, -78 °C, 10 minutes. 2. NaCN (10 equiv.), DMSO, rt, 1 hour. 3. TBSCl (3 equiv.), pyridine (8 equiv.), AgNO₃ (3 equiv.), DMF, 0 °C-rt, 18 hours (87% for 3 steps). *c.* 1. DIBAL (1.1 equiv.), CH₂Cl₂, hexanes, -78 °C, 30 minutes. 2. (CF₃CH₂O)₂P(O)CH₂CO₂Me

(1.5 equiv.), 18-Crown-6 (8 equiv.), KHMDS (1.5 equiv.), THF, -78 °C, 30 minutes (84% for 2 steps). 3. DIBAL (4 equiv.), THF, -78 °C~0 °C, 30 minutes (99%). *d.* 1. (+)-DET (20 mol%), Ti(OPr-*i*)₄ (15 mol%), TBHP (1.5 equiv.), MS 4A, CH₂Cl₂, -10 °C, 15 hours (86% for desired isomer, 11% for undesired isomer). 2. TBAF (6 equiv.), MS 4Å, THF (96%). *e.* 1. TBSCl (1.5 equiv.), Et₃N (4 equiv.), CH₂Cl₂, rt, 5 hours (99%). 2. TESCl (1.2 equiv.), imidazole (3 equiv.), CH₂Cl₂, 0 °C-rt, 15 minutes. 3. 9-BBN (3 equiv.), THF, 0 °C-rt, 1 hour then NaBO₃•H₂O aq. (94% for 2 steps). 4. TEMPO (20 mol%), PhI(OAc)₂ (3 equiv.), CH₂Cl₂, rt, 36 hours (95%). *f.* 1. **5** (1.3 equiv.), *t*-BuLi (2.5 equiv.), THF, -78 °C, 30 minutes. 2. OsO₄ (10 mol%), NMMO (2 equiv.), H₂O, acetone, rt, 4 hours. 3. Pb(OAc)₄ (1.5 equiv.), K₂CO₃ (10 equiv.), CH₂Cl₂, rt, 15 minutes (68% for 3 steps). 4. (MeO)₂P(=O)CH₂CO₂Bn (5 equiv.), NaH (4 equiv.), THF, 0 °C, 3 hours (88%). *g.* 1. LiBr (10 equiv.), DBU (20 equiv.), MeCN, rt, 11 hours (70%). *h.* DDQ (3 equiv.), CH₂Cl₂, *t*-BuOH, pH 7 buffer, rt, 15 minutes (86%). 2. TESCl (1.5 equiv.), imidazole (3 equiv.), CH₂Cl₂, rt, 4 hr (97%). 3. H₂ (1 atm), Pd/C, AcOEt, rt, 2 hours (89%). 4. (PyS)₂ (1.2 equiv.), PPh₃ (3 equiv.), toluene, rt, 12 hours (97%). Abbreviation: 18-Crown-6 = 1,4,7,10,13,16-hexa-oxacyclooctadecane; KHMDS = potassium bis(trimethylsilyl)amide; 9-BBN = 9-borabicyclononane; DET = diethyl tartrate; TBHP = *tert*-butyl hydroperoxide; MS = molecular sieves; TBAF = tetrabutylammonium fluoride.

[0097] Figure 19 shows an exemplary synthesis of a left hand C38-C53 building block in the norhalichondrin series. *Reagents and conditions:* *a.* 1. Tf₂O (1.2 equiv.), 2,6-lutidine (4 equiv.), CH₂Cl₂, -78 °C, 10 minutes. 2. NaCN (10 equiv.), DMSO, rt, 1 hour (87% for two steps). 3. DIBAL (4.5 equiv.), CH₂Cl₂, -78 °C, 30 minutes. 4. NaBH₄ (5 equiv.), MeOH, rt, 30 minutes. 5. TBSOTf (3 equiv.), 2,6-lutidine (3.5 equiv.), CH₂Cl₂, rt, 30 minutes (90% for 3 steps). 6. 9-BBN (2 equiv.), THF, rt, 2 hours, then NaOH, H₂O₂, H₂O, rt, 3 hr (91%). 7. TEMPO (0.5 equiv.), PhI(OAc)₂ (5.0 equiv.), CH₃CN, H₂O, THF, rt, 12 hours (90%). 8. *p*-TsOH•H₂O (1.0 equiv.), H₂O (10 equiv.), CH₂Cl₂, rt, 24 hours. 9. TESOTf (10 equiv.), 2,6-lutidine (12 equiv.), CH₂Cl₂, rt, 1 hour (76% for 2 steps). *b.* 1. **5**, *t*-BuLi (2.2 equiv.), toluene, Et₂O, -78 °C, 10 minutes (82%). 2. OsO₄ (5 mol%), NMMO (2 equiv.), H₂O, acetone, rt, 12 hours. 3. Pb(OAc)₄ (2 equiv.), K₂CO₃ (10 equiv.), rt, 30 minutes (86% for 2 steps). 4. (MeO)₂P(=O)CH₂CO₂Bn (4 equiv.), K₃PO₄ (3 equiv.), rt, 36 hours (93%). *c.* LiBr (10 equiv.), DBU (5 equiv.), BnOAc (2 equiv.), CH₃CN, rt, 12 hours (82%). *d.* 1. TBAF (1.5 equiv.), HOAc (1.0 equiv.), THF, 0 °C, 5 hours (81%). 2. Dess-Martin periodinane (2.0 equiv.), NaHCO₃ (10 equiv.), CH₂Cl₂, rt, 30 min. 3. NaClO₂ (3 equiv.), NaH₂PO₄ (4 equiv.),

2-methyl-2-butene, *t*-BuOH, H₂O, rt, 30 minutes. 4. TMSCH₂N₂ (3.0 equiv.), benzene, MeOH, rt, 5 minutes (87% for 3 steps). *e.* 1. DDQ (2.0 equiv.), CH₂Cl₂, aqueous pH7 buffer, rt, 1 hour. 2. TESOTf (2.0 equiv.), 2,6-lutidine (2.5 equiv.), CH₂Cl₂, rt, 30 minutes (83% for 2 steps). *f.* 1. Pd/C (10 wt%), H₂, EtOAc, rt, 3 hours. 2. (SPy)₂ (1.4 equiv.), PPh₃ (1.2 equiv.), toluene, rt, 12 hours (88% for 2 steps). Abbreviation: *p*-TsOH = *p*-toluenesulfonic acid.

[0098] Figure 20 shows an X-Ray Structure for Halichondrin C prepared using the methods described herein. A colorless single crystal of Halichondrin C was obtained by recrystallization from MeOH:CH₂Cl₂ = 1:1.

[0099] Figure 21 shows an exemplary synthetic scheme for the preparation of an exemplary C33-C43 fragment of halichondrins and analogs thereof.

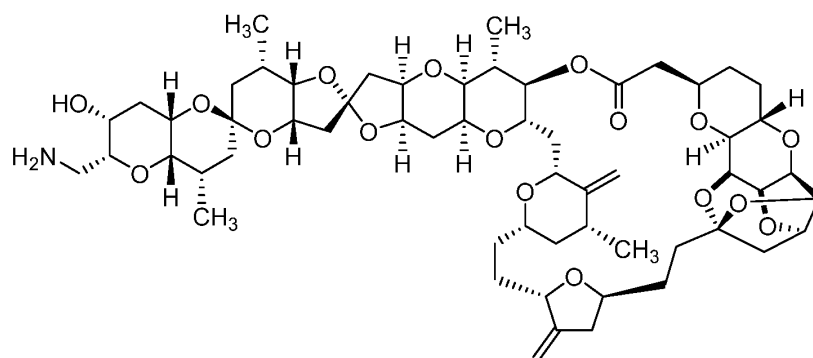
[00100] Figure 22 shows an exemplary synthetic scheme for the preparation of an exemplary C27-C37 fragment of halichondrins and analogs thereof.

[00101] Figure 23 shows an exemplary synthetic scheme for the preparation of an exemplary C39-C43 fragment of halichondrins and analogs thereof.

DETAILED DESCRIPTION OF CERTAIN EMBODIMENTS

[00102] Provided herein are Ni/Zr-mediated coupling reactions useful in the preparation of ketone-containing compounds. The Ni/Zr-mediated ketolization reactions provided herein are particularly useful in the synthesis of halichondrins and analogs thereof. Therefore, also provided herein are methods for the preparation of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C; norhalichondrin A, B, C) and analogs thereof.

[00103] In certain embodiments, provided herein are methods useful in the preparation of compounds of Formula (H3-A), including Compound (1):



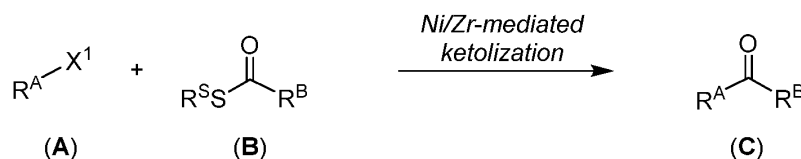
Compound (1).

[00104] The present invention also provides compounds (*i.e.*, intermediates) useful in the methods provided herein. In certain embodiments, the compounds provided herein are useful as synthetic intermediates *en route* to halichondrins and analogs thereof. Furthermore, the present invention provides reagents and catalysts useful in the methods described herein.

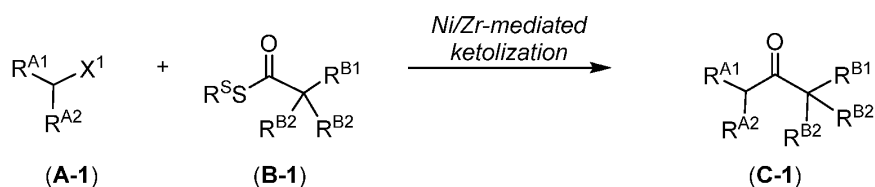
Ni/Zr-Mediated Ketolization Reactions

[00105] In one aspect, provided herein are nickel/zirconium-mediated ketolization reactions (“Ni/Zr-mediated ketolization reactions”) involving a coupling of a thioester and an alkyl halide (*e.g.*, alkyl iodide, alkyl bromide, alkyl chloride, *etc.*) or alkyl leaving group (*e.g.*, alkyl sulfonate) (*Scheme 1A*). The ketolization reactions may be intermolecular or intramolecular (*i.e.*, in *Scheme 1A*, R^A and R^B are optionally joined by a linker). In certain embodiments, the compound of Formula (A) is a primary or secondary alkyl halide (X¹ = halogen), and the compound of Formula (B) is an alkyl thioester (R^B = optionally substituted alkyl), as shown in *Scheme 1B*.

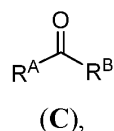
Scheme 1A



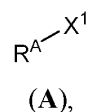
Scheme 1B



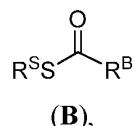
[00106] As represented in *Scheme 1A*, provided herein are methods for preparing a compound of Formula (C):



or a salt thereof, the methods comprising reacting a compound of Formula (A):



or a salt thereof, with a compound of Formula (B):



or a salt thereof, in the presence of nickel and zirconium; wherein:

R^{A} is optionally substituted alkyl;

R^{B} is optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted carbocyclyl, optionally substituted heteroaryl, or optionally substituted heterocyclyl;

optionally wherein R^{A} and R^{B} are joined together via a linker, wherein the linker is selected from the group consisting of optionally substituted alkylene, optionally substituted heteroalkylene, optionally substituted alkenylene, optionally substituted heteroalkenylene, optionally substituted alkynylene, optionally substituted heteroalkynylene, optionally substituted arylene, optionally substituted heteroarylene, optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted acylene, and combinations thereof;

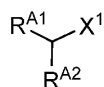
X^1 is halogen or a leaving group; and

R^{S} is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl.

[00107] In certain embodiments, R^{A} is a small molecule. In certain embodiments, R^{B} is a small molecule. Small molecules encompass complex small molecules, such as natural products, pharmaceutical agents, and fragments thereof, and intermediates thereto.

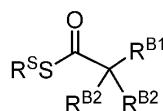
[00108] As generally defined herein, a “linker” is a group comprising optionally substituted alkylene, optionally substituted heteroalkylene, optionally substituted alkenylene, optionally substituted heteroalkenylene, optionally substituted alkynylene, optionally substituted heteroalkynylene, optionally substituted arylene, optionally substituted heteroarylene, optionally substituted carbocyclylene, optionally substituted heterocyclylene, optionally substituted acylene, or any combination thereof.

[00109] In certain embodiments, the compound of Formula (A) is of Formula (A-1):



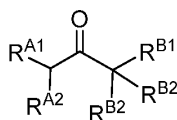
(A-1),

or a salt thereof; the compound of Formula (B) is of Formula (B-1):



(B-1),

or a salt thereof; and the compound of Formula (C) is of Formula (C-1):



(C-1),

or a salt thereof, wherein:

X^1 is halogen or a leaving group;

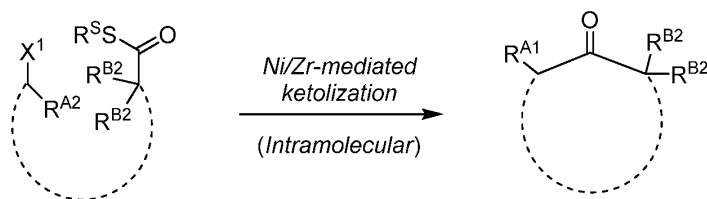
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

each instance of R^{A1} , R^{A2} , R^{B1} , and R^{B2} is independently hydrogen, optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted carbocyclyl, optionally substituted heteroaryl, or optionally substituted heterocyclyl; optionally wherein R^{A1} and R^{B1} are joined together via a linker.

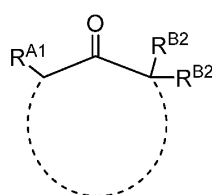
[00110] In certain embodiments, R^{A1} is a small molecule. In certain embodiments, R^{B1} and R^{B2} are independently a small molecules. Small molecules encompass complex small molecules, such as natural products, pharmaceutical agents, and fragments thereof, and intermediates thereto.

[00111] The Ni/Zr-mediated ketolization reactions provided herein may be performed in an intramolecular fashion to yield cyclic ketones as shown in *Scheme 1C*.

Scheme 1C

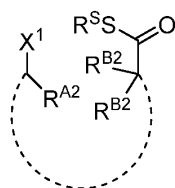


[00112] As shown in *Scheme 1C*, provided herein are methods for preparing a compound of Formula (C-2):



(C-2),

or salt thereof, comprising reacting a compound of Formula (A-B):



(A-B),

or a salt thereof, in the presence of nickel and zirconium; wherein:

R^{A2} and R^{B2} are optionally substituted alkyl, optionally substituted alkenyl, optionally substituted alkynyl, optionally substituted aryl, optionally substituted carbocyclyl, optionally substituted heteroaryl, or optionally substituted heterocyclyl;

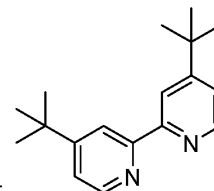
X^1 is halogen or a leaving group;

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl; and

represents a linker.

[00113] Ni/Zr-mediated ketolization reactions provided herein are carried out in the presence of nickel. In certain embodiments, the ketolization reaction is carried out in the presence of a nickel complex. Any nickel complex (*e.g.*, nickel salt, nickel complex, nickel catalyst, or nickel pre-catalyst) known or available in the art may be used in the reaction. In certain embodiments, the ketolization reaction is carried out in the presence of nickel (II). In certain

embodiment, the ketolization reaction is carried out in the presence of a nickel (0). In certain embodiments, the nickel complex is of the formula: $\text{NiX}_2\bullet(\text{ligand})$, wherein X is halogen (*e.g.*, Cl, Br, I, or F). In certain embodiments, “ligand” is a bidentate ligand. In certain embodiments, the ligand is an optionally substituted bipyridyl ligand. In certain embodiments, the nickel complex is $\text{NiX}_2\bullet(\text{tbbpy})$, wherein X is halogen (*e.g.*, Cl, Br, I, or



F), and “tbbpy” is 4,4'-bis(*tert*-butyl)-2,2'-bipyridine, having the structure:

. In certain embodiments, the nickel complex is $\text{NiCl}_2\bullet(\text{tbbpy})$. In certain embodiments, the nickel complex is $\text{NiBr}_2\bullet(\text{tbbpy})$.

[00114] In certain embodiments, the nickel complex is used after complexation of a nickel source and a “ligand” in solution. In certain embodiments, the nickel complex is of the formula: $\text{NiX}_2\bullet(\text{ligand})$; wherein X is halogen and “ligand” is a bidentate ligand. In certain embodiments, the nickel source is NiCl_2 ; the “ligand” is 4,4'-di-*tert*-butyl-2,2'-dipyridyl (tbbpy); and the resulting nickel complex is of the formula $\text{NiCl}_2\bullet(\text{tbbpy})$. In certain embodiments, the nickel source is NiBr_2 ; and the “ligand” is 4,4'-di-*tert*-butyl-2,2'-dipyridyl (tbbpy); and the resulting nickel complex is of the formula $\text{NiBr}_2\bullet(\text{tbbpy})$.

[00115] In certain embodiments, the nickel is present in a catalytic amount. In certain embodiments, the nickel is present at approximately 1-5 mol%, 5-10 mol%, 1-10 mol%, 5-20 mol%, 10-20 mol%, 20-30 mol%, 20-40 mol%, 30-40 mol%, 40-50 mol%, 50-60 mol%, 60-70 mol%, 70-80 mol%, or 80-90 mol% relative to a compound of Formula (A) or (B) in the reaction mixture. In certain embodiments, the nickel is present in from 1-50 mol%. In certain embodiments, the nickel is present in from 1-10 mol%. In certain embodiments, the nickel is present in approximately 5 mol%. In certain embodiments, the nickel is present in approximately 30 mol%. In certain embodiments, the nickel is present in a stoichiometric or excess amount relative to a compound of Formula (A) or (B) in the reaction mixture. In certain embodiments, approximately 1 equivalent of nickel is present (*i.e.*, stoichiometric). In other embodiments, greater than 1 equivalent of nickel is present (*i.e.*, excess).

[00116] As described above, the Ni/Zr-mediated ketolization reactions are carried out in the presence of zirconium. In certain embodiments, the reaction is carried out in the presence of a zirconium complex. Any zirconium source (*e.g.*, zirconium salt, complex, catalyst or precatalyst) known or available in the art may be used in the reaction. In certain

embodiments, the zirconium source is of the formula (ligand)_nZrX₂; wherein n is the number of ligands (*e.g.*, 0, 1, 2, 3, 4), and X is halogen (*e.g.*, Cl, Br, I, or F). In certain embodiments, n is 2, and the ligand is cyclopentadienyl. In certain embodiments, the zirconium source is Cp₂ZrX₂. In certain embodiments, the zirconium source is Cp₂ZrCl₂.

[00117] In certain embodiments, the zirconium is present in a catalytic amount. In certain embodiments, the zirconium is present in between 1-5 mol%, 5-10 mol%, 1-10 mol%, 5-20 mol%, 10-20 mol%, 20-30 mol%, 30-40 mol%, 40-50 mol%, 50-60 mol%, 60-70 mol%, 70-80 mol%, or 80-90 mol% relative to a compound of Formula (A) or (B) in the reaction mixture. In certain embodiments, the zirconium is present in a stoichiometric or excess amount relative to a compound of Formula (A) or (B) in the reaction mixture. In certain embodiments, approximately 1 equivalent of zirconium is present (*i.e.*, stoichiometric). In other embodiments, greater than 1 equivalent of zirconium is present (*i.e.*, excess). In certain embodiments, approximately 1.1, 1.2, 1.3, 1.4, 1.5, 1.6, 1.7, 1.8, 1.9, 2, 2.5, 3, 3.5, 4, 4.5, 5, 6, 7, 8, 9, or 10 equivalents of zirconium is present. In certain embodiments, approximately 3 equivalents of zirconium is present.

[00118] In certain embodiments, a Ni/Zr-mediated ketolization reaction provided herein is performed in the presence of one or more additional reagents or catalysts, such as a reducing metal. In certain embodiments, the reducing metal is zinc. In certain embodiments, the reducing metal is magnesium. In certain embodiments, zinc metal is used (*i.e.*, zinc(0)). In certain embodiments, magnesium metal is used (*i.e.*, magnesium(0)). In certain embodiments, the reaction is carried out in the presence of zinc powder, zinc foil, zinc beads, or any other form of zinc metal. In certain embodiments, a zinc salt is employed such as zinc acetate, zinc sulfate, zinc chloride, zinc bromide, zinc iodide, zinc fluoride, zinc sulfide, or zinc phosphate. The zinc may be present in a catalytic, stoichiometric, or excess amount. In certain embodiments, the zinc is present in excess (*i.e.*, greater than 1 equivalent) relative to a compound of Formula (A) or Formula (B). In certain embodiments, between 1 and 10 equivalents of zinc are used. In certain embodiments, approximately 1.5, 2, 2.5, 3, 3.5, 4, 4.5, 5, 5.5, 6, 7, 8, 9, or 10 equivalents of zinc are present. In certain embodiments, approximately 6 equivalents of zinc are used.

[00119] In certain embodiments, the ketolization reaction is carried out in the presence of one or more reagents which help activate zinc metal in the reaction (*e.g.*, by clearing the surface of zinc oxide). In certain embodiments, the reaction is carried out in the presence of a trialkylsilyl halide (*e.g.*, triethylsilyl chloride (TESCl)). This reagent may be present in a catalytic, stoichiometric, or excess amount. In certain embodiments, approximately 1.5, 2,

2.5, 3, 3.5, 4, 4.5, 5, 5.5, 6, 7, 8, 9, or 10 equivalents of this reagent is present. In certain embodiments, approximately 1.5 equivalents of this reagent is present.

[00120] In certain embodiments, the Ni/Zr-mediated ketolization is carried out in the presence of one or more additional reagents (*i.e.*, in addition to nickel, zirconium, and zinc).

[00121] In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a base or proton scavenger. In certain embodiments, the base is a pyridine base. In certain embodiments, the base is 2,6-di-*tert*-butyl pyridine. In certain embodiments, the base is 2,6-lutidine. In certain embodiments, the base is 2,6-di-*tert*-butyl-4-methylpyridine. In certain embodiments, the base is used in a stoichiometric or excess amount. In certain embodiments, approximately 1.5, 2, 2.5, 3, 3.5, 4, 4.5, 5, 5.5, 6, 7, 8, 9, or 10 equivalents of the base or proton scavenger is present. In certain embodiments, approximately 4 equivalents of the base or proton scavenger is employed.

[00122] In certain embodiments, the Ni/Zr-mediated ketolization described herein is carried out in a solvent. Any solvent may be used, and the scope of the method is not limited to any particular solvent or mixture of solvents. The solvent may be polar or non-polar, protic or aprotic, or a combination of solvents (*e.g.*, co-solvents). Examples of useful organic solvents are provided herein. In certain embodiments, the ketolization reaction is carried out in 1,3-dimethyl-2-imidazolidinone (DMI). In certain embodiments, the ketolization reaction is carried out in a 1,3-dimethyl-2-imidazolidinone (DMI)/tetrahydrofuran (THF) mixture. In certain embodiments, the ketolization reaction is carried out in a 1,3-dimethyl-2-imidazolidinone (DMI)/ethyl acetate (EtOAc) mixture.

[00123] The Ni/Zr-mediated ketolization reactions described herein may be carried out at any concentration in solvent. Concentration refers to the molar concentration (mol/L) of a coupling partners (*e.g.*, compounds of Formula (A) or (B)) in a solvent. In certain embodiments, the concentration is about 0.1 M. In certain embodiments, the concentration is approximately 0.5 M. In certain embodiments, the concentration is approximately 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, or 0.9 M. In certain embodiments, the concentration is greater than 1 M. In certain embodiments, the concentration is less than 0.1 M.

[00124] The Ni/Zr-mediated ketolization reactions described herein can be carried out at any temperature. In certain embodiments, the reaction is carried out at around room temperature (*i.e.*, between 18 and 24 °C). In certain embodiments, the reaction is carried out below room temperature (*e.g.*, between 0 °C and room temperature). In certain embodiments, the reaction is carried out at above room temperature (*e.g.*, between room temperature and 100 °C). In certain embodiments, the reaction is carried out at a temperature ranging from approximately

room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. [00125] In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a nickel complex, a zirconium complex, and a reducing metal. In certain embodiments, the nickel complex is NiBr₂(dtbbpy). In certain embodiments, the zirconium complex is Cp₂ZrCl₂. In certain embodiments, the reducing metal is zinc. In certain embodiments, the reaction is carried out in the presence of NiBr₂(dtbbpy), Cp₂ZrCl₂, and zinc metal. In certain embodiments, the reaction is carried out in a polar solvent such as DMI (1,3-dimethyl-2-imidazolidinone). In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. For example, in certain embodiments, the coupling is carried out under the following conditions: 5 mol% NiBr₂(dtbbpy), 1.0 equivalent Cp₂ZrCl₂, excess zinc metal, in DMI at room temperature.

[00126] In certain embodiments, the reaction is carried out in the presence of NiBr₂(dtbbpy), Cp₂ZrCl₂, zinc metal, and a base or proton scavenger. In certain embodiments, the reaction is carried out in the presence of NiBr₂(dtbbpy), Cp₂ZrCl₂, zinc metal, and (*t*-Bu)₂(Me)Py. In certain embodiments, the reaction is carried out in a mixture of DMI and EtOAc (ethyl acetate). In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. For example, in certain embodiments, the coupling is carried out under the following conditions: 30 mol% NiBr₂(dtbbpy), 3.0 equivalents Cp₂ZrCl₂, 6.0 equivalents zinc metal, and 4.0 equivalents (*t*-Bu)₂(Me)Py, in DMI-EtOAc at room temperature.

Synthesis of Halichondrins and Analogs

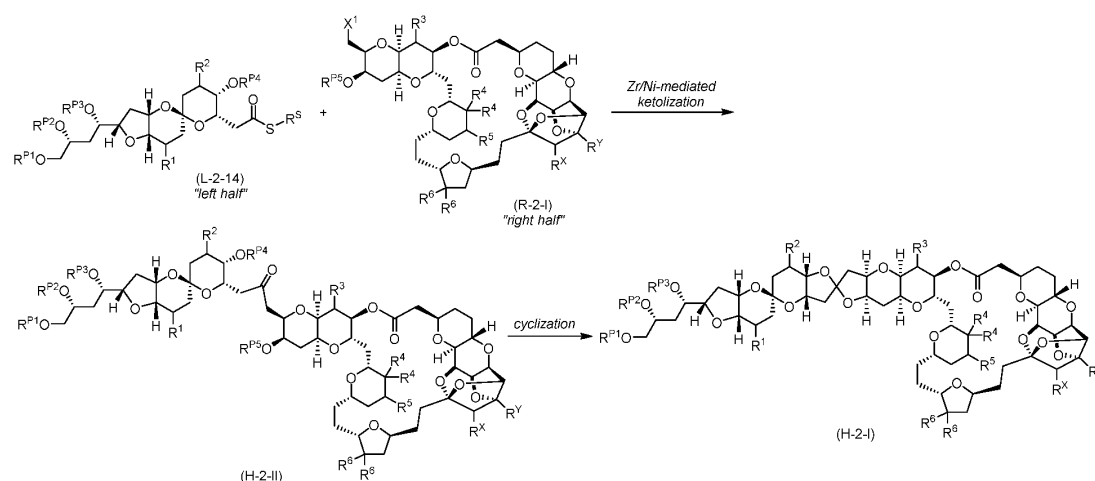
[00127] The Ni/Zr-mediated ketolization reactions provided herein can be applied to the synthesis of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C, norhalichondrin A, B, C) and analogs thereof. In certain embodiments, methods are useful in the synthesis of compounds of Formula (H3-A), such as Compound (1). In certain embodiments, the methods comprise the steps of: (1) coupling a “left half” building block

with a “right half” building block via a Ni/Zr-mediated ketolization reaction provided herein; followed by (2) cyclizing the resulting coupling product (*e.g.*, acid-mediated cyclization); optionally, followed by any necessary synthetic transformations to arrive at a desired product.

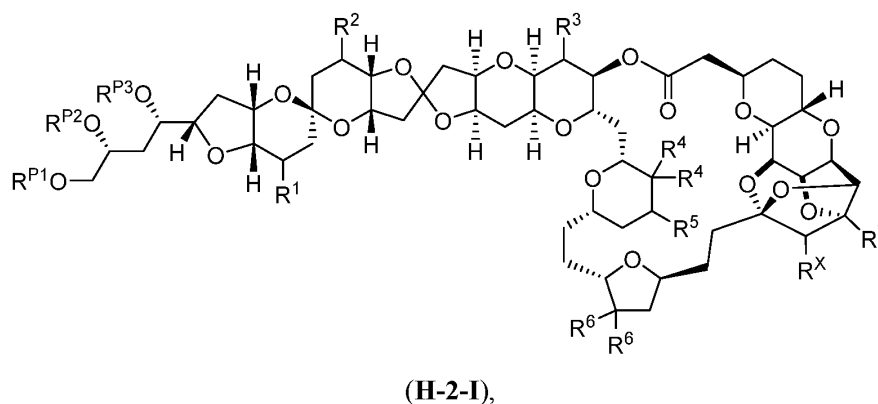
Synthesis of Halichondrins

[00128] The Ni/Zr-mediated ketolization reactions provided herein can be applied to the preparation of halichondrins (*e.g.*, halichondrin A, B, C) and analogs thereof. For example, as shown in *Scheme 2A*, coupling of a left half of Formula (L-2-14) with a right half of Formula (R-2-I) via a Ni/Zr-mediated ketolization yields a ketone of Formula (H-2-II), cyclization of which provides a compound of Formula (H-2-I), which is a halichondrin or an analog thereof, or an intermediate thereto.

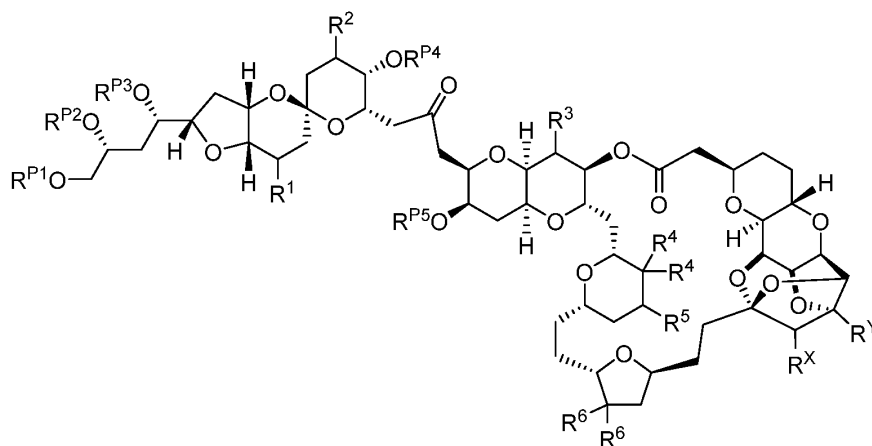
Scheme 2A



[00129] Provided herein is a method of preparing a compound of Formula (H-2-I):



or a salt thereof, the method comprising cyclizing a compound of Formula (H-2-II):

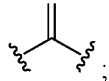


(H-2-II),

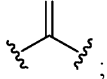
or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, optionally substituted alkyl,

or two R^6 groups are taken together to form: ;

R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

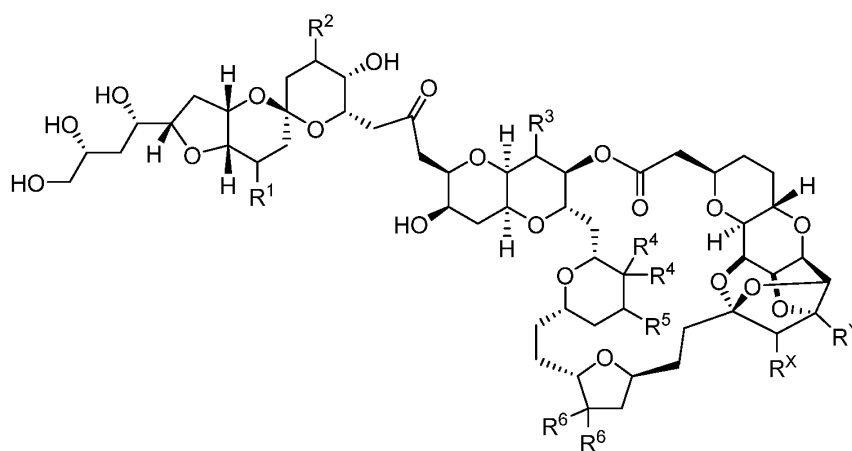
[00130] In certain embodiments, the step of cyclizing a compound of Formula (H-2-II), or a salt thereof, is carried out in the presence of an acid. The acid may be a Lewis acid or a Brønsted acid. In certain embodiments, the acid is a Brønsted acid. In certain embodiments, the acid is a sulfonic acid. In certain embodiments, the acid is a salt of a sulfonic acid. In certain embodiments, the acid is a pyridinium salt. In certain embodiments, the acid is

pyridinium *p*-toluenesulfonate (PPTS). In certain embodiments, the acid is present in a catalytic amount. In certain embodiments, the acid is present in a stoichiometric (*e.g.*, approximately 1 equivalent) or excess amount (*e.g.*, greater than 1 equivalent). In certain embodiments, the acid is present in an excess amount (*e.g.*, about 5 equivalents).

[00131] In certain embodiments, the step of cyclizing is carried out in the presence of PPTS. In certain embodiments, the step is carried out in a solvent such as CH₂Cl₂. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 20 °C. For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 5 equivalents of PPTS in CH₂Cl₂ at around 20 °C (*e.g.*, for 2 hours).

[00132] In certain embodiments, R^{P1}, R^{P2}, and R^{P3} are silyl protecting groups, and R^{P4} and R^{P5} are hydrogen. In certain embodiments, R^{P1} and R^{P2} are TBS, R^{P3} is TES, and R^{P4} and R^{P5} are hydrogen.

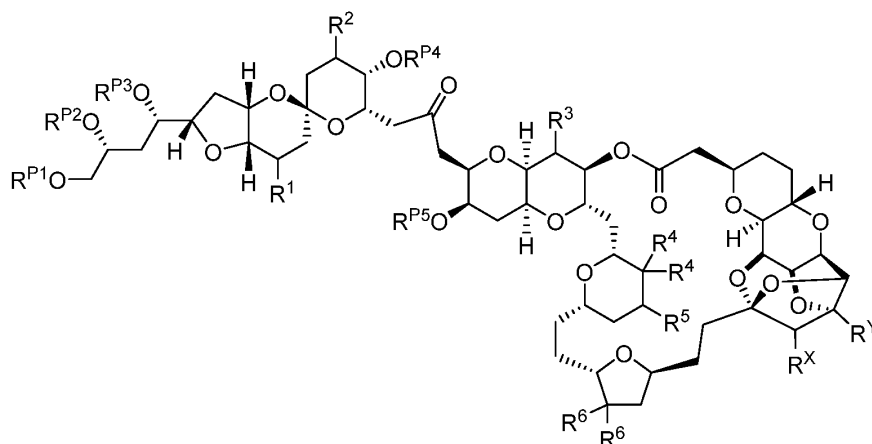
[00133] In certain embodiments, the compound of Formula (H-2-II) is of Formula (H-2-IIA):



(H-2-IIA),

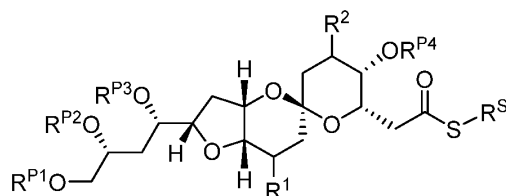
or a salt thereof.

[00134] Provided herein is a method of preparing a compound of Formula (H-2-II):



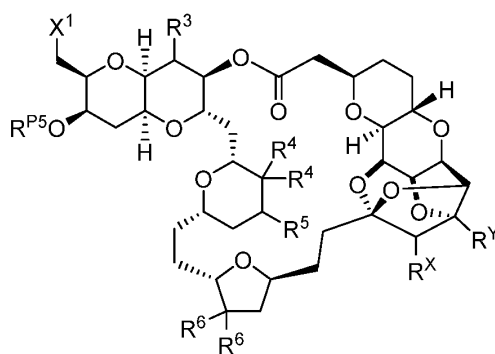
(H-2-II),

or a salt thereof, the method comprising coupling a compound of Formula (L-2-14):



(L-2-14),

or a salt thereof, with a compound of Formula (R-2-I):



(R-2-I),

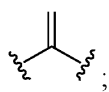
or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

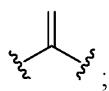
X^1 is halogen or a leaving group;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00135] In certain embodiments, the step of coupling to provide a compound of Formula (H-2-II) is a Ni/Zr-mediated ketolization provided herein. Any reagents or conditions provided herein for the Ni/Zr-mediated ketolization may be used in the coupling. In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a nickel complex, a zirconium complex, and a reducing metal. The reaction may also be carried out in the presence of one or more additional reagents, such a base or proton scavenger. In certain embodiments, the nickel complex is $NiBr_2(dtbbpy)$. In certain embodiments, the zirconium complex is Cp_2ZrCl_2 . In certain embodiments, the reducing metal is zinc. In certain embodiments, the additional base or proton scavenger is $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , zinc metal, and $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in a polar solvent such as DMI (1,3-dimethyl-2-imidazolidinone). In certain embodiments, the reaction is carried out in a mixture of DMI and EtOAc (ethyl acetate). In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to

approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature.

[00136] For example, in certain embodiments, the coupling is carried out under the following conditions: 30 mol% NiBr₂(dtbbpy), 3.0 equivalents Cp₂ZrCl₂, 6.0 equivalents zinc metal, and 4.0 equivalents (*t*-Bu)₂(Me)Py, in DMI-EtOAc at room temperature.

[00137] In certain embodiments, R^{P1}, R^{P2}, R^{P3}, R^{P4} and R^{P5} are silyl protecting groups. In certain embodiments, R^{P1} and R^{P2} are TBS; and R^{P3}, R^{P4}, and R^{P5} are TES.

[00138] In certain embodiments, the method of preparing a compound of Formula (H-2-II) further comprises one or more steps of deprotecting one or more oxygen atoms of the compound of Formula (H-2-II) (*e.g.*, to yield a compound of Formula (H-2-III), or a salt thereof). In certain embodiments, the resulting compound or salt thereof can then be used in the cyclization step to yield a compound of Formula (H-2-I), or a salt thereof. In certain embodiments, the step of deprotecting is carried out in the presence of a fluoride source (*e.g.*, when the one or more oxygen atoms are protected with silyl groups).

[00139] Examples of fluoride sources useful in the invention include, but are not limited to, metal fluorides (*e.g.*, sodium fluoride, potassium fluoride, cesium fluoride, silver fluoride) and tetraalkylammonium fluorides (*e.g.*, tetramethylammonium fluoride, tetraethylammonium fluoride, tetrabutylammonium fluoride). In certain embodiments, the fluoride source is a tetraalkylammonium fluoride. In certain embodiments, the fluoride source is tetrabutylammonium fluoride (TBAF). In certain embodiments, hydrogen fluoride (HF) is used. In certain embodiments, HF•pyridine is used as the HF source. Other examples of protecting groups useful in the present invention, and reagents useful in protection/deprotection reactions can be found in the art, *e.g.*, in *Protecting Groups in Organic Synthesis*, T. W. Greene and P. G. M. Wuts, 3rd edition, John Wiley & Sons, 1999, incorporated herein by reference.

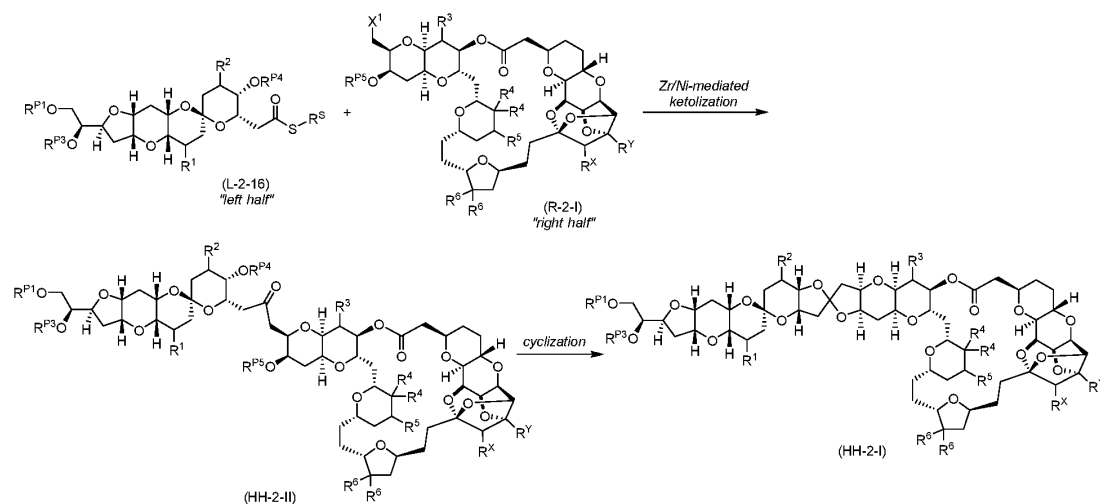
[00140] Once a compound of Formula (H-2-I), or salt thereof, is obtained, the method may comprise one or more additional steps (*e.g.*, deprotection, protection, substitution, addition, elimination) to yield a desired compound (*e.g.*, halichondrin A, B, C, or an analog thereof).

Synthesis of Homohalichondrins

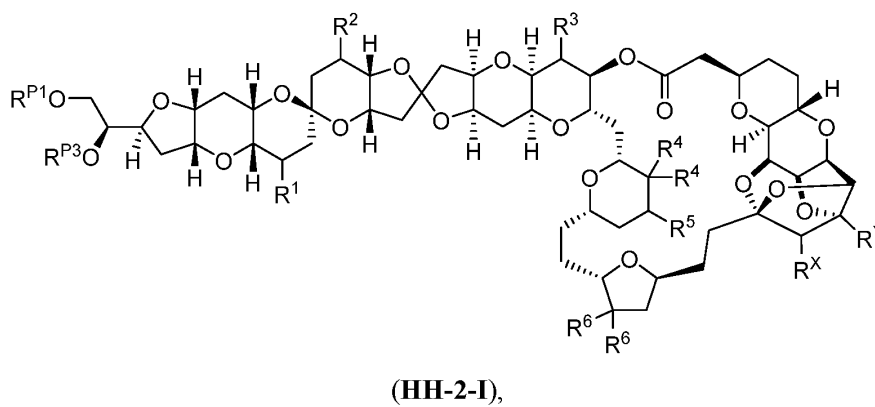
[00141] The Ni/Zr-mediated ketolization reactions provided herein can be applied to the preparation of homohalichondrins (*e.g.*, homohalichondrin A, B, C), and analogs thereof. For example, as shown in *Scheme 2B*, coupling of a left half of Formula (L-2-16) with a right half

of Formula **(R-2-I)** via a Ni/Zr-mediated ketolization yields a ketone of Formula **(HH-2-II)**, cyclization of which provides a compound of Formula **(HH-2-I)**, which is a homohalichondrin natural product or an analog thereof, or an intermediate thereto.

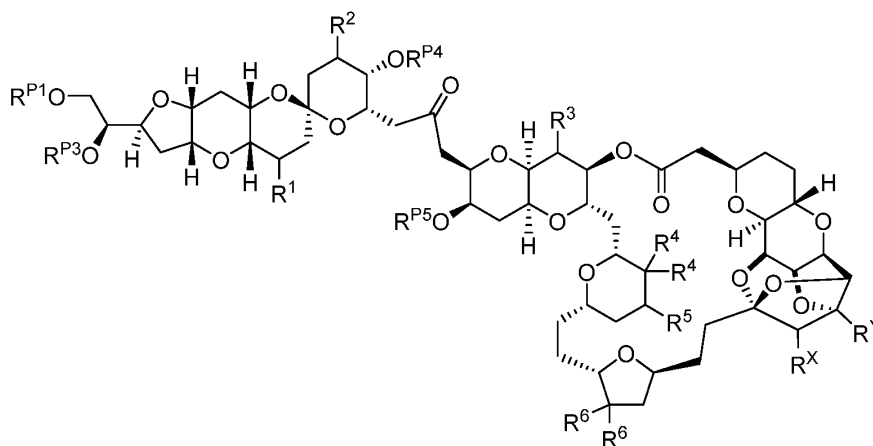
Scheme 2B



[00142] Provided herein is a method of preparing a compound of Formula **(HH-2-I)**:



or a salt thereof, the method comprising cyclizing a compound of Formula (HH-2-II):

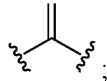


(HH-2-II),

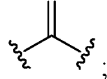
or a salt thereof, wherein:

R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R⁴ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁴ groups are taken together to form: ;

each instance of R⁶ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁶ groups are taken together to form: ;

R^{P1}, R^{P3}, R^{P4}, and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or -OR^{Xa}, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or -OR^{Ya}, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

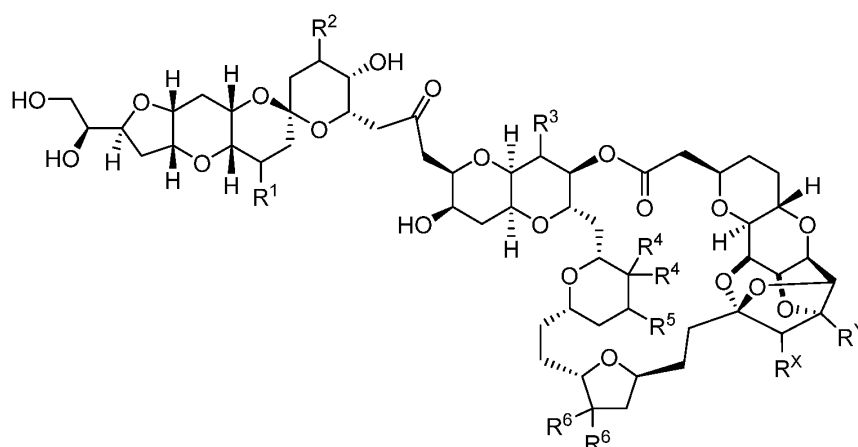
[00143] In certain embodiments, the step of cyclizing a compound of Formula (HH-2-II), or a salt thereof, is carried out in the presence of an acid. The acid may be a Lewis acid or a Brønsted acid. In certain embodiments, the acid is a Brønsted acid. In certain embodiments, the acid is a sulfonic acid. In certain embodiments, the acid is a salt of a sulfonic acid. In certain embodiments, the acid is a pyridinium salt. In certain embodiments, the acid is

pyridinium *p*-toluenesulfonate (PPTS). In certain embodiments, the acid is present in a catalytic amount. In certain embodiments, the acid is present in a stoichiometric (*e.g.*, approximately 1 equivalent) or excess amount (*e.g.*, greater than 1 equivalent). In certain embodiments, the acid is present in an excess amount (*e.g.*, about 5 equivalents).

[00144] In certain embodiments, the step of cyclizing is carried out in the presence of PPTS. In certain embodiments, the step is carried out in a solvent such as CH₂Cl₂. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 20 °C. For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 5 equivalents of PPTS in CH₂Cl₂ at around 20 °C (*e.g.*, for 2 hours).

[00145] In certain embodiments, R^{P1} and R^{P2} are silyl protecting groups; and R^{P4} and R^{P5} are hydrogen. In certain embodiments, R^{P1} is TBS; R^{P2} is TES; and R^{P4} and R^{P5} are hydrogen.

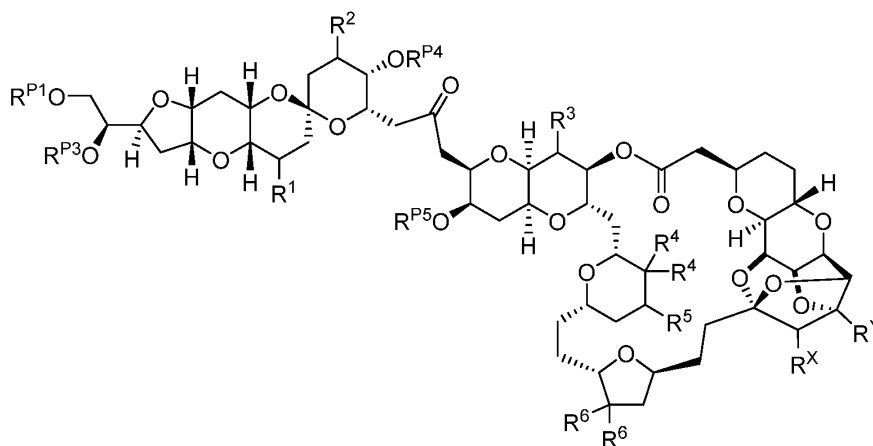
[00146] In certain embodiments, the compound of Formula (HH-2-II) is of Formula (HH-2-IIA):



(HH-2-IIA),

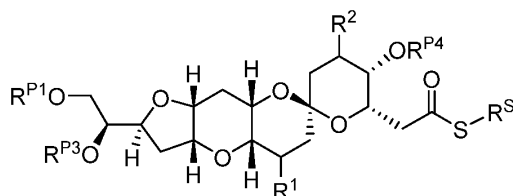
or a salt thereof.

[00147] Provided herein is a method of preparing a compound of Formula (HH-2-II):



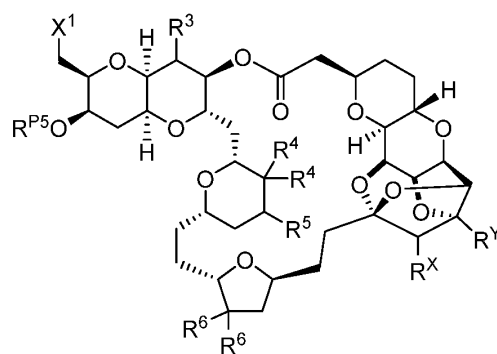
(HH-2-II),

or a salt thereof, the method comprising coupling a compound of Formula (L-2-16):



(L-2-16),

or a salt thereof, with a compound of Formula (R-2-I):



(R-2-I),

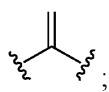
or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

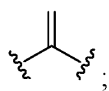
X^1 is halogen or a leaving group;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

R^{P1} , R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00148] In certain embodiments, the step of coupling to provide a compound of Formula (HH-2-II) is a Ni/Zr-mediated ketolization as provided herein. Any reagents or conditions provided herein for the Ni/Zr-mediated ketolization may be used in the coupling. In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a nickel complex, a zirconium complex, and a reducing metal. The reaction may also be carried out in the presence of one or more additional reagents, such a base or proton scavenger. In certain embodiments, the nickel complex is $NiBr_2(dtbbpy)$. In certain embodiments, the zirconium complex is Cp_2ZrCl_2 . In certain embodiments, the reducing metal is zinc. In certain embodiments, the additional base or proton scavenger is $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , zinc metal, and $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in a polar solvent such as DMI (1,3-dimethyl-2-imidazolidinone). In certain embodiments, the reaction is carried out in a mixture of DMI and EtOAc (ethyl acetate). In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to

approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature.

[00149] For example, in certain embodiments, the coupling is carried out under the following conditions: 30 mol% NiBr₂(dtbbpy), 3.0 equivalents Cp₂ZrCl₂, 6.0 equivalents zinc metal, and 4.0 equivalents (*t*-Bu)₂(Me)Py, in DMI-EtOAc at room temperature.

[00150] In certain embodiments, R^{P1}, R^{P2}, R^{P3}, R^{P4} and R^{P5} are silyl protecting groups. In certain embodiments, R^{P1} and R^{P2} are TBS; and R^{P3}, R^{P4}, and R^{P5} are TES.

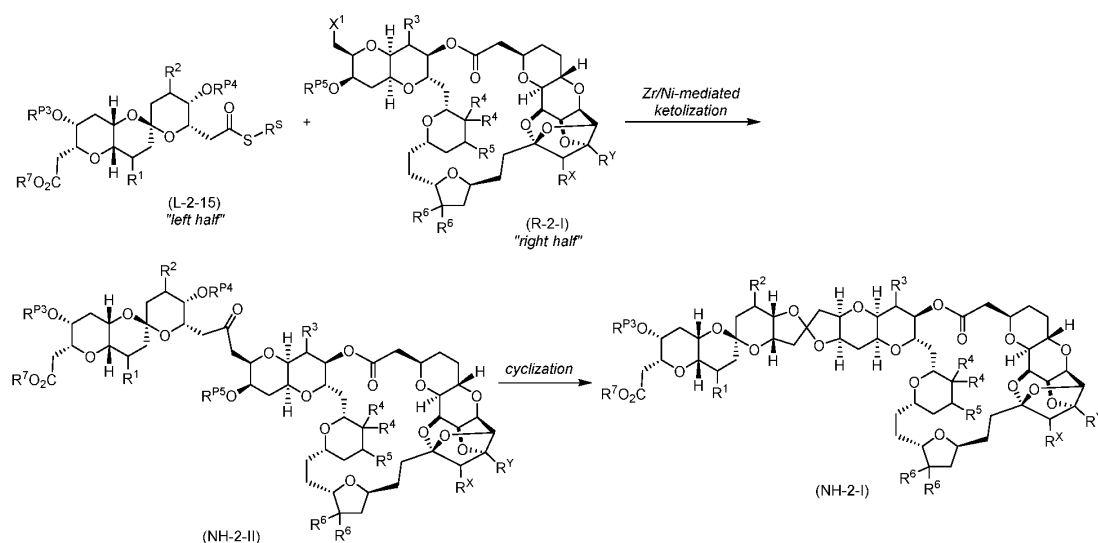
[00151] In certain embodiments, the method of preparing a compound of Formula (HH-2-II) further comprises one or more steps of deprotecting one or more oxygen atoms of the compound of Formula (HH-2-II) (*e.g.*, to yield a compound of Formula (HH-2-IIA), or a salt thereof). In certain embodiments, the resulting compound, or salt thereof, is then cyclized to yield a compound of Formula (HH-2-I), or a salt thereof. In certain embodiments, a step of deprotecting is carried out in the presence of a fluoride source (*e.g.*, when one or more oxygen atoms are protected with silyl groups). Examples of fluoride sources are provided herein.

[00152] Once a compound of Formula (HH-2-I), or salt thereof, is obtained, one or more additional steps (*e.g.*, deprotection, protection, substitution, addition, elimination) may be performed to yield a desired compound (*e.g.*, homohalichondrin A, B, C, or an analog thereof, or intermediate thereto).

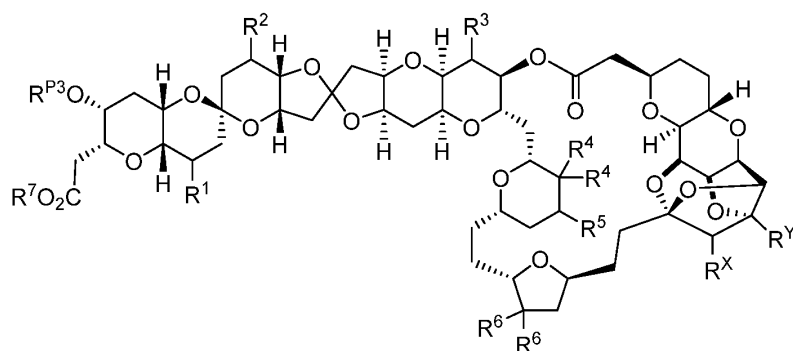
Synthesis of Norhalichondrins

[00153] The Ni/Zr-mediated ketolization reactions provided herein can be applied to the preparation of norhalichondrins (*e.g.*, norhalichondrin A, B, C) and analogs thereof. For example, as shown in *Scheme 2C*, coupling of a left half of Formula (L-2-15) with a right half of Formula (R-2-I) via a Ni/Zr-mediated ketolization yields a ketone of Formula (NH-2-II), cyclization of which provides a compound of Formula (NH-2-I), which is a norhalichondrin or an analog thereof, or intermediate thereto.

Scheme 2C

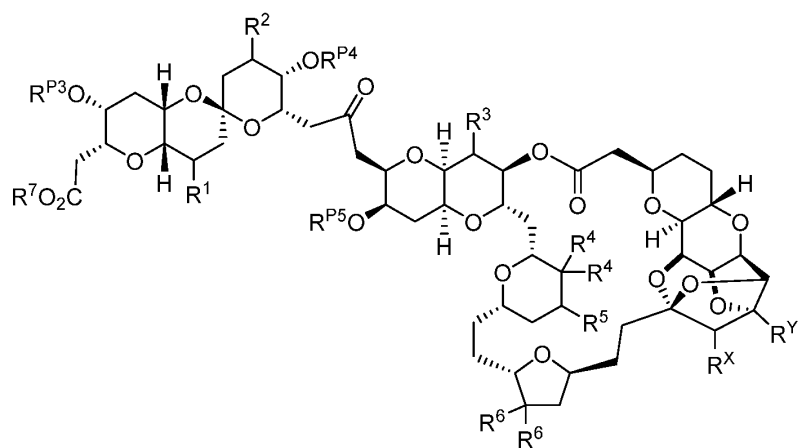


[00154] Provided herein is a method of preparing a compound of Formula (NH-2-I):



(NH-2-I),

or a salt thereof, the method comprising cyclizing a compound of Formula (NH-2-II):

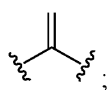


(NH-2-II),

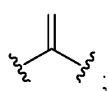
or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^7 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

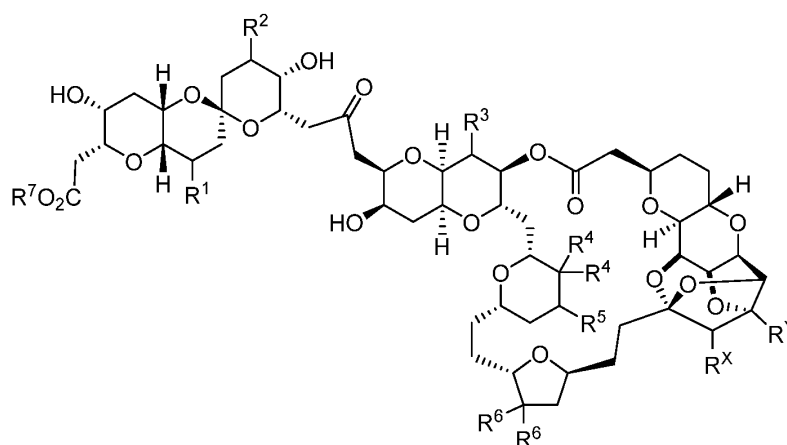
[00155] In certain embodiments, the step of cyclizing a compound of Formula (NH-2-II), or a salt thereof, is carried out in the presence of an acid. The acid may be a Lewis acid or a Brønsted acid. In certain embodiments, the acid is a Brønsted acid. In certain embodiments, the acid is a sulfonic acid. In certain embodiments, the acid is a salt of a sulfonic acid. In certain embodiments, the acid is a pyridinium salt. In certain embodiments, the acid is pyridinium *p*-toluenesulfonate (PPTS). In certain embodiments, the acid is present in a catalytic amount. In certain embodiments, the acid is present in a stoichiometric (*e.g.*, approximately 1 equivalent) or excess amount (*e.g.*, greater than 1 equivalent). In certain embodiments, the acid is present in an excess amount (*e.g.*, about 5 equivalents).

[00156] In certain embodiments, the step of cyclizing is carried out in the presence of PPTS. In certain embodiments, the step is carried out in a solvent such as CH_2Cl_2 . In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room

temperature. In certain embodiments, the reaction is carried out at around 20 °C. For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 5 equivalents of PPTS in CH₂Cl₂ at around 20 °C (*e.g.*, for 2 hours).

[00157] In certain embodiments, R^{P3} is a silyl protecting group; R⁷ is optionally substituted alkyl; and R^{P4} and R^{P5} are hydrogen. In certain embodiments, R^{P3} is TES; R⁷ is methyl; and R^{P4} and R^{P5} are hydrogen.

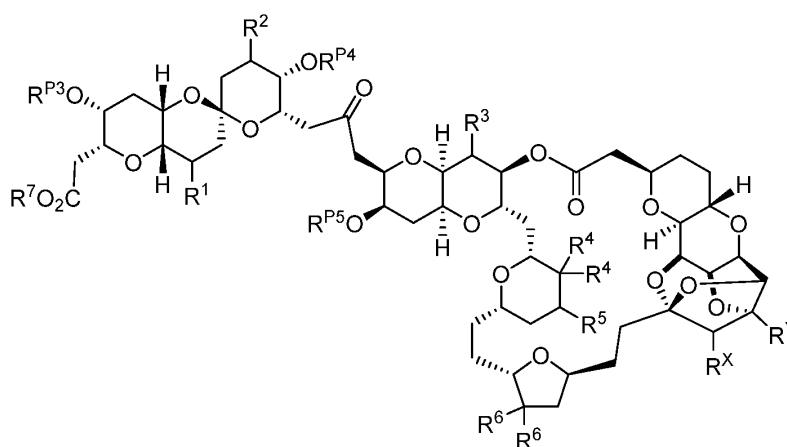
[00158] In certain embodiments, the compound of Formula (NH-2-II) is of Formula (NH-2-IIA):



(NH-2-IIA),

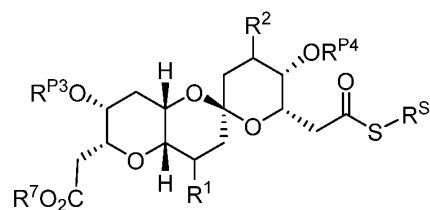
or a salt thereof.

[00159] Provided herein is a method of preparing a compound of Formula (NH-2-II):



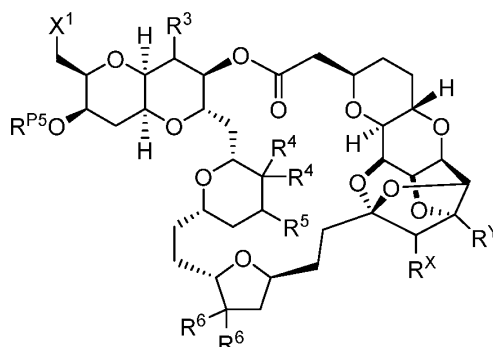
(NH-2-II),

or a salt thereof, the method comprising coupling a compound of Formula (L-2-15):



(L-2-15),

or a salt thereof, with a compound of Formula (R-2-I):



(R-2-I),

or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

X^1 is halogen or a leaving group;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^7 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00160] In certain embodiments, the step of coupling to provide a compound of Formula (NH-2-II) is a Ni/Zr-mediated ketolization provided herein. Any reagents or conditions provided herein for the Ni/Zr-mediated ketolization may be used in the coupling. In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a nickel complex, a zirconium complex, and a reducing metal. The reaction may also be carried out in the presence of one or more additional reagents, such a base or proton scavenger. In certain embodiments, the nickel complex is $NiBr_2(dtbbpy)$. In certain embodiments, the zirconium complex is Cp_2ZrCl_2 . In certain embodiments, the reducing metal is zinc. In certain embodiments, the additional base or proton scavenger is $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $NiBr_2(dtbbpy)$, Cp_2ZrCl_2 , zinc metal, and $(t-Bu)_2(Me)Py$. In certain embodiments, the reaction is carried out in a polar solvent such as DMI (1,3-dimethyl-2-imidazolidinone). In certain embodiments, the reaction is carried out in a mixture of DMI and EtOAc (ethyl acetate). In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature.

[00161] For example, in certain embodiments, the coupling is carried out under the following conditions: 30 mol% $NiBr_2(dtbbpy)$, 3.0 equivalents Cp_2ZrCl_2 , 6.0 equivalents zinc metal, and 4.0 equivalents $(t-Bu)_2(Me)Py$, in DMI-EtOAc at room temperature.

[00162] In certain embodiments, R^{P3} is a silyl protecting group; R^7 is optionally substituted alkyl; and R^{P4} and R^{P5} are silyl protecting groups. In certain embodiments, R^{P3} is TES; R^7 is methyl; and R^{P4} and R^{P5} are TES.

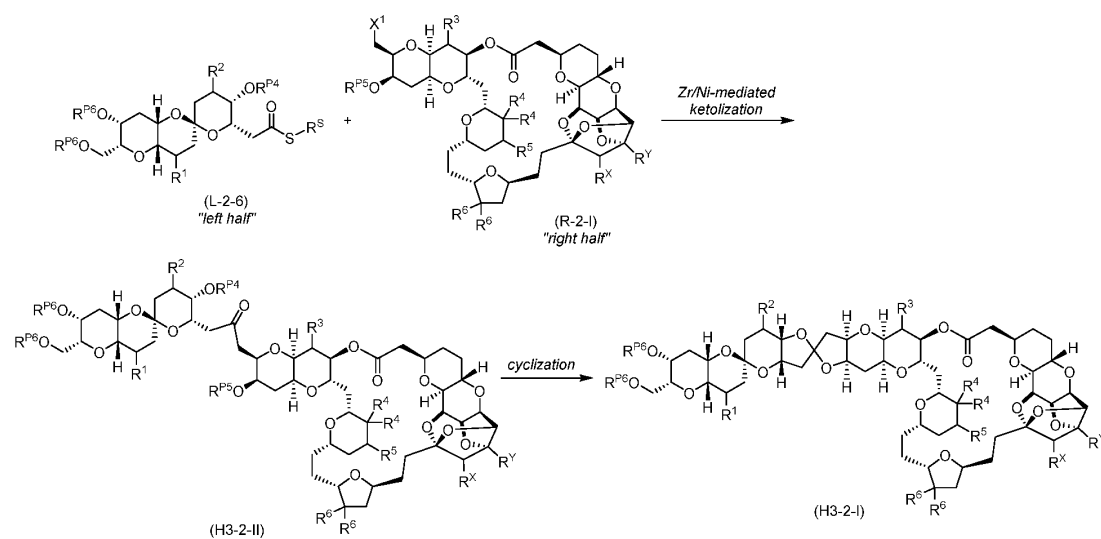
[00163] In certain embodiments, the method of preparing a compound of Formula (**NH-2-II**) further comprises one or more steps of deprotecting one or more oxygen atoms of the compound of Formula (**NH-2-II**) (*e.g.*, to yield a compound of Formula (**NH-2-IIA**), or a salt thereof). In certain embodiments, the resulting compound, or salt thereof, is then cyclized to yield a compound of Formula (**NH-2-I**), or a salt thereof. In certain embodiments, a step of deprotecting is carried out in the presence of a fluoride source (*e.g.*, when the one or more oxygen atoms are protected with silyl groups). Examples of fluoride sources are provided herein.

[00164] Once a compound of Formula (**NH-2-I**), or salt thereof, is obtained, the method may comprise one or more additional steps (*e.g.*, deprotection, protection, substitution, addition, elimination) to yield a desired compound (*e.g.*, homohalichondrin A, B, C, or an analog thereof).

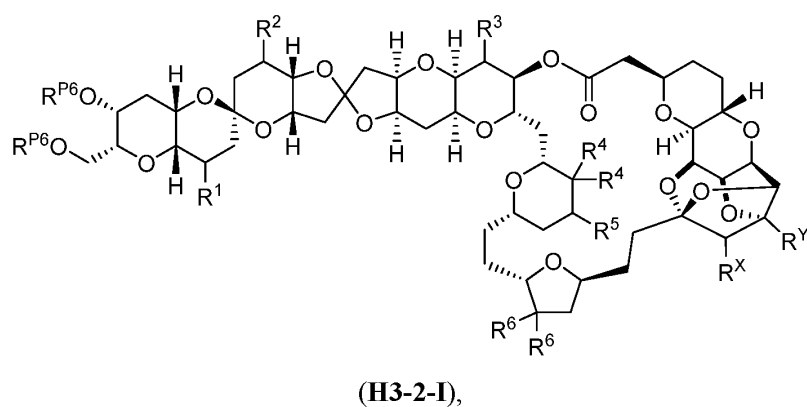
Synthesis of Additional Halichondrin Analogs

[00165] Methods for the preparation of additional halichondrin analogs are provided herein. The Ni/Zr-mediated ketolization reactions provided herein can be applied to the preparation of additional halichondrin analogs. For example, as shown in *Scheme 2D*, coupling of a left half of Formula (**L-2-6**) with a right half of Formula (**R-2-I**) via a Ni/Zr-mediated ketolization yields a ketone of Formula (**H3-2-II**), cyclization of which provides a compound of Formula (**H3-2-I**). The compound of Formula (**H3-2-I**) can be subjected to further synthetic transformation to yield a desired compound.

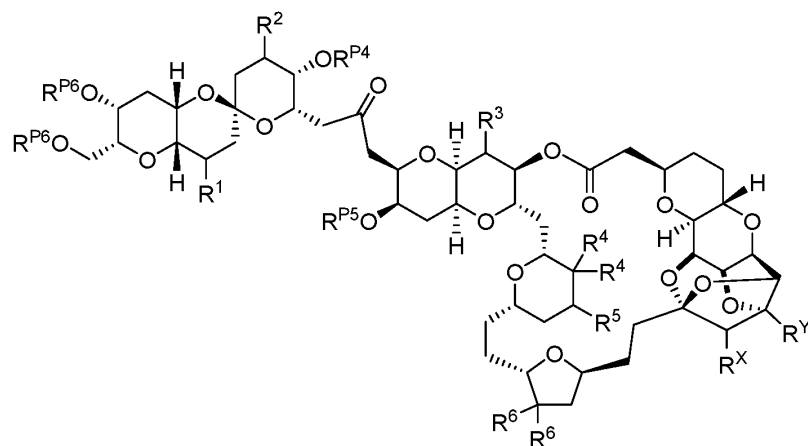
Scheme 2D



[00166] As shown in *Scheme 2D*, provided herein is a method of preparing a compound of Formula (H3-2-I):



or a salt thereof, the method comprising cyclizing a compound of Formula (H3-2-II):

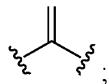


(H3-2-II),

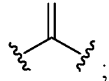
or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

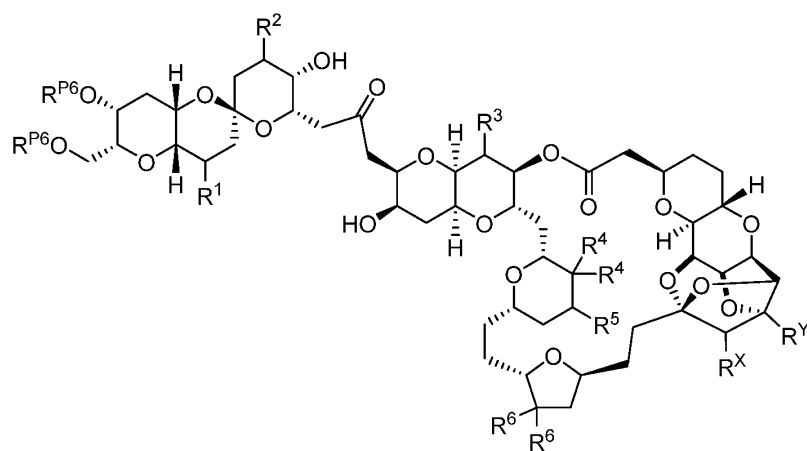
R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

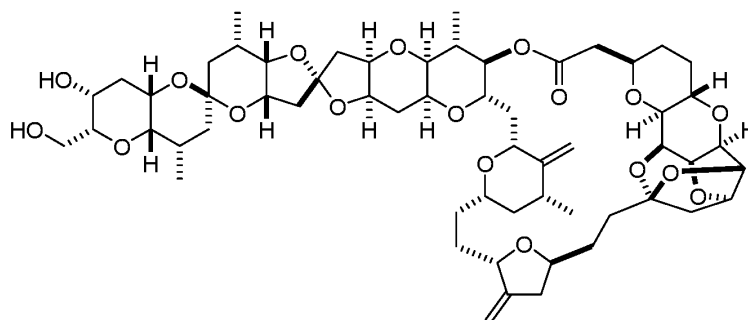
[00167] In certain embodiments, the compound of Formula (H3-2-II) is of Formula (H3-2-IIA):



(H3-2-IIA),

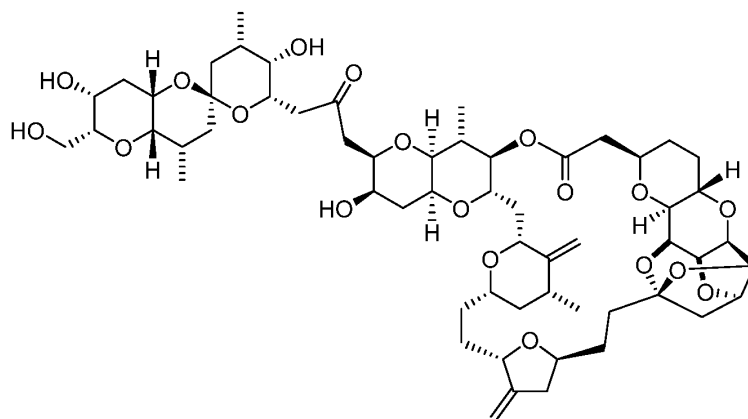
or a salt thereof.

[00168] In certain embodiments, the method is a method of preparing Compound (2):



Compound (2),

or a salt thereof, the method comprising cyclizing a compound of the formula:



Compound (C),

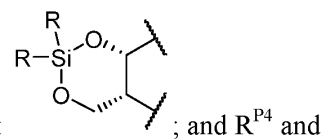
or a salt thereof.

[00169] In certain embodiments, the step of cyclizing a compound of Formula (H3-2-II), Compound (C), or a salt thereof, is carried out in the presence of an acid. The acid may be a Lewis acid or a Brønsted acid. In certain embodiments, the acid is a Brønsted acid. In certain embodiments, the acid is a sulfonic acid. In certain embodiments, the acid is a salt of a sulfonic acid. In certain embodiments, the acid is a pyridinium salt. In certain embodiments, the acid is pyridinium *p*-toluenesulfonate (PPTS). In certain embodiments, the acid is present in a catalytic amount. In certain embodiments, the acid is present in a stoichiometric (*e.g.*, approximately 1 equivalent) or excess amount (*e.g.*, greater than 1 equivalent). In certain embodiments, the acid is present in an excess amount (*e.g.*, about 5 equivalents). In certain embodiments, the step is carried out in a solvent. In certain embodiments, the reaction is carried out in dichloromethane (DCM). In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0°C to approximately 50 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0°C to approximately room temperature. In certain embodiments, the reaction is carried out at around 20 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 9-11 °C.

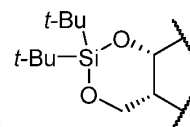
[00170] In certain embodiments, the step of cyclizing is carried out in the presence of PPTS. In certain embodiments, the step of cyclizing is carried out in the presence of PPTS in DCM. For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 5 equivalents of PPTS in DCM at around 20 °C (*e.g.*, for 2 hours). For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 5 equivalents of PPTS in DCM at around 9-11 °C (*e.g.*, for 3 hours).

[00171] In certain embodiments, two R^{P6} are oxygen protecting groups; and R^{P4} and R^{P5} are

hydrogen. In certain embodiments, two R^{P6} are joined to form:



; and R^{P4} and

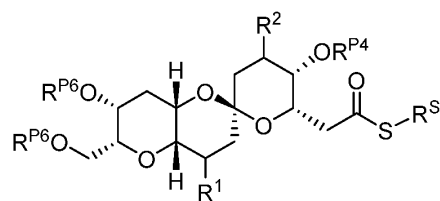


R^{P5} are hydrogen. In certain embodiments, two R^{P6} are joined to form:

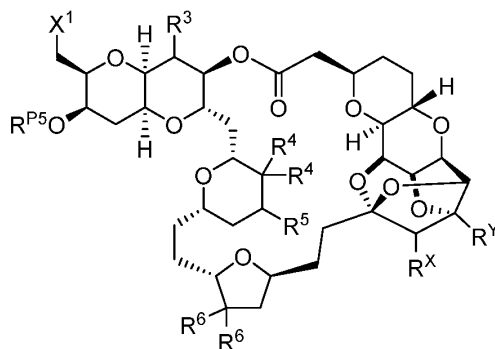
R^{P4} and R^{P5} are hydrogen. In certain embodiments, each R^{P6}, R^{P4}, and R^{P5} are each hydrogen.

In certain embodiments, one or more free hydroxyl groups of Compound (C) is substituted with an oxygen protecting group (*e.g.*, a silyl protecting group).

or a salt thereof, the method comprising coupling a compound of Formula (L-2-6):



or a salt thereof, with a compound of Formula **(R-2-I)**:



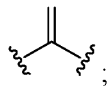
or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

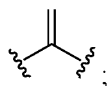
X¹ is halogen or a leaving group;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

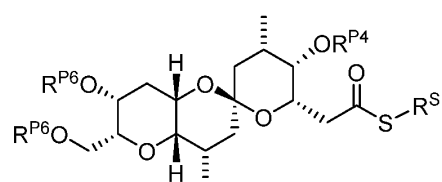
R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

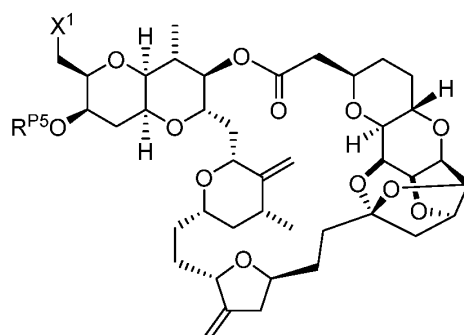
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00173] In certain embodiments, the method comprises coupling a compound of Formula (E-L):



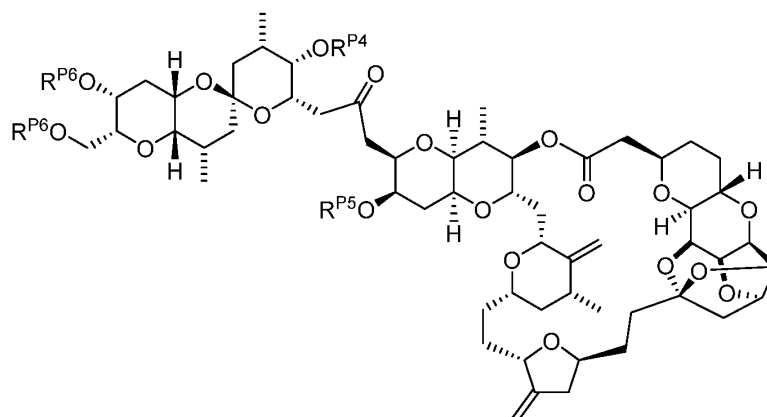
(E-L),

or a salt thereof, with a compound of the formula (E-R):



(E-R),

or a salt thereof, to yield a compound of the formula (E-1):



(E-1),

or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

X¹ is halogen or a leaving group; and

R^{P4}, R^{P5}, and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00174] In certain embodiments, the step of coupling to provide a compound of Formula (H3-2-II), (E-1), or a salt thereof, is a Ni/Zr-mediated ketolization provided herein. Any reagents or conditions provided herein for the Ni/Zr-mediated ketolization may be used in the coupling. See, e.g., the section entitled *Ni/Zr-Mediated Ketolization Reactions* above.

[00175] In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of nickel and zirconium complexes. In certain embodiments, the Ni/Zr-mediated ketolization reaction is carried out in the presence of a nickel complex, a zirconium complex, and a reducing metal.

[00176] In certain embodiments, the nickel is a nickel complex. In certain embodiments, the nickel is a nickel(II) or nickel(0) complex. In certain embodiments, the nickel complex is of the formula: NiX₂•(ligand); wherein X is halogen and “ligand” is a bidentate ligand. In certain embodiments, the nickel complex is used after complexation of a nickel source and a “ligand” in solution. In certain embodiments, the nickel source is NiCl₂; the “ligand” is 4,4'-di-tert-butyl-2,2'-dipyridyl (tbbpy); and the nickel complex is of the formula NiCl₂•(tbbpy).

In certain embodiments, the nickel source is NiBr_2 ; and the “ligand” is 4,4'-di-tert-butyl-2,2'-dipyridyl (tbbpy); and the nickel complex is of the formula $\text{NiBr}_2 \bullet (\text{tbbpy})$.

[00177] In certain embodiments, the zirconium complex is Cp_2ZrCl_2 . In certain embodiments, Cp_2ZrCl_2 is present in a stoichiometric or excess amount (e.g., from 1-4 equivalents). In certain embodiments, the reducing metal is zinc metal. In certain embodiments, the reducing metal is manganese metal. In certain embodiments, the zinc or manganese metal is present in an excess amount. The reaction may also be carried out in the presence of one or more additional reagents, such a base and/or proton scavenger. In certain embodiments, the reaction is carried out in the presence of $(t\text{-Bu})_2(\text{Me})\text{Py}$. In certain embodiments, the reaction is carried out in the presence of proton sponge (e.g., 1,8-bis(dimethylamino)naphthalene).

[00178] In certain embodiments, the reaction is carried out in the presence of $\text{NiBr}_2(\text{dtbbpy})$, Cp_2ZrCl_2 , and zinc metal. In certain embodiments, the reaction is carried out in the presence of $\text{NiBr}_2(\text{dtbbpy})$, Cp_2ZrCl_2 , and manganese metal. In certain embodiments, the reaction is carried out in the presence of $\text{NiBr}_2(\text{dtbbpy})$, Cp_2ZrCl_2 , zinc metal, and $(t\text{-Bu})_2(\text{Me})\text{Py}$. In certain embodiments, the reaction is carried out in the presence of $\text{NiBr}_2(\text{dtbbpy})$, Cp_2ZrCl_2 , manganese metal, and $(t\text{-Bu})_2(\text{Me})\text{Py}$.

[00179] In certain embodiments, the reaction is carried out in a polar solvent, such as DMI (1,3-dimethyl-2-imidazolidinone). In certain embodiments, the reaction is carried out in a mixture of DMI and EtOAc (ethyl acetate). In certain embodiments, the reaction is carried out in a mixture of DMI and ethanol. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 30 °C.

[00180] For example, in certain embodiments, the coupling is carried out under the following conditions: 30 mol% $\text{NiBr}_2(\text{dtbbpy})$, 3.0 equivalents Cp_2ZrCl_2 , 6.0 equivalents zinc metal, and 4.0 equivalents $(t\text{-Bu})_2(\text{Me})\text{Py}$, in DMI-EtOAc at room temperature.

[00181] In certain embodiments, the coupling is carried out in the presence of $\text{NiBr}_2(\text{dtbbpy})$, Cp_2ZrCl_2 , and manganese metal in DMI. For example, in certain embodiments, the coupling is carried out under the following conditions: approximately 75 mol% $\text{NiBr}_2(\text{dtbbpy})$, 3.5

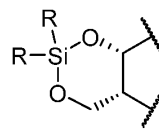
equivalents Cp_2ZrCl_2 , and 7 equivalents manganese metal in DMI at around 30 °C (*e.g.*, for 4 hours).

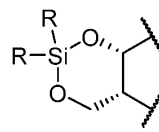
[00182] In certain embodiments, the coupling is carried out by reacting a compound of Formula (**L-2-6**), or a salt thereof, in the presence of a compound of Formula (**R-2-I**), or a salt thereof, Cp_2ZrCl_2 , and manganese metal; followed by the addition of $\text{NiBr}_2(\text{dtbbpy})$ to the reaction mixture. In certain embodiments, the coupling is carried out by reacting a compound of Formula (**L-2-6**), or a salt thereof, in the presence of a compound of Formula (**R-2-I**), or a salt thereof, Cp_2ZrCl_2 , and manganese metal in DMI; followed by the addition of $\text{NiBr}_2(\text{dtbbpy})$ in a solution of DMI to the reaction mixture.

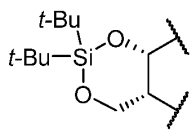
[00183] In certain embodiments, the coupling is carried out by reacting a compound of Formula (**E-L**), or a salt thereof, in the presence of a compound of Formula (**R-L**), or a salt thereof, Cp_2ZrCl_2 , and manganese metal; followed by the addition of $\text{NiBr}_2(\text{dtbbpy})$ to the reaction mixture. In certain embodiments, the coupling is carried out by reacting a compound of Formula (**E-L**), or a salt thereof, in the presence of a compound of Formula (**R-L**), or a salt thereof, Cp_2ZrCl_2 , and manganese metal in DMI; followed by the addition of $\text{NiBr}_2(\text{dtbbpy})$ in a solution of DMI to the reaction mixture.

[00184] The coupling reaction to yield a compound of Formula (**H3-2-II**), (**E-1**), or a salt thereof, can be carried out to yield any amount of product. In certain embodiments, the reaction is carried out to yield more than 1 g, 2 g, 5 g, 10 g, 20 g, 30 g, 50 g, 100 g, 200 g, 500 g, or 1 kg of product. In certain embodiments, the reaction is carried out to yield less than 1 g of product. In certain embodiments, the reaction is carried out to yield from 1 g to 100 g of product, inclusive. In certain embodiments, the reaction is carried out to yield approximately 1 g, 2 g, 5 g, 10 g, 20 g, 30 g, 40 g, 50 g, 60 g, 70 g, 80 g, 90 g, or 100 g of product.

[00185] In certain embodiments, X^1 is a halogen and R^{S} is optionally substituted pyridyl. In certain embodiments, X^1 is $-\text{I}$. In certain embodiments, R^{S} is 2-pyridyl. In certain embodiments, X^1 is $-\text{I}$; and R^{S} is 2-pyridyl.



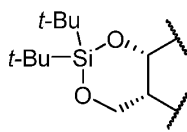
[00186] In certain embodiments, two R^{P6} are joined to form: ; and R^{P4} and R^{P5} are silyl protecting groups. In certain embodiments, two R^{P6} are joined to form:

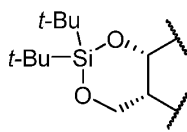


; and R^{P4} and R^{P5} are TES.

[00187] In certain embodiments, the method of preparing a compound of Formula (H3-2-II) further comprises one or more steps of deprotecting one or more oxygen atoms (*e.g.*, removing groups R^{P4} , R^{P5} , and/or R^{P6}) of the compound of Formula (H3-2-II) (*e.g.*, to yield a compound of Formula (H3-2-IIA), or a salt thereof). In certain embodiments, the resulting compound, or salt thereof, can then be used in the cyclization step to yield a compound of Formula (H3-2-I), or a salt thereof. Likewise, the method of preparing a compound of Formula (E-1) can further comprise one or more steps of deprotecting one or more oxygen atoms (*e.g.*, removing groups R^{P4} , R^{P5} , and/or R^{P6}) of the compound of Formula (E-1) (*e.g.*, to yield Compound (C), or a salt thereof). In certain embodiments, the resulting compound, or salt thereof, can then be used in the cyclization step to yield Compound (2).

[00188] In certain embodiments, a step of deprotecting is carried out in the presence of a fluoride source (*e.g.*, when R^{P4} , R^{P5} , and/or R^{P6} are silyl protecting groups). Examples of fluoride sources are provided herein. In certain embodiments, the fluoride source is TBAF. In certain embodiments, the step of deprotection is carried out in the presence of an imidazole hydrochloride. In certain embodiments, R^{P4} and R^{P5} are TES; and the step of deprotecting (to remove R^{P4} and R^{P5}) is carried out in the presence of TBAF and imidazole hydrochloride. In

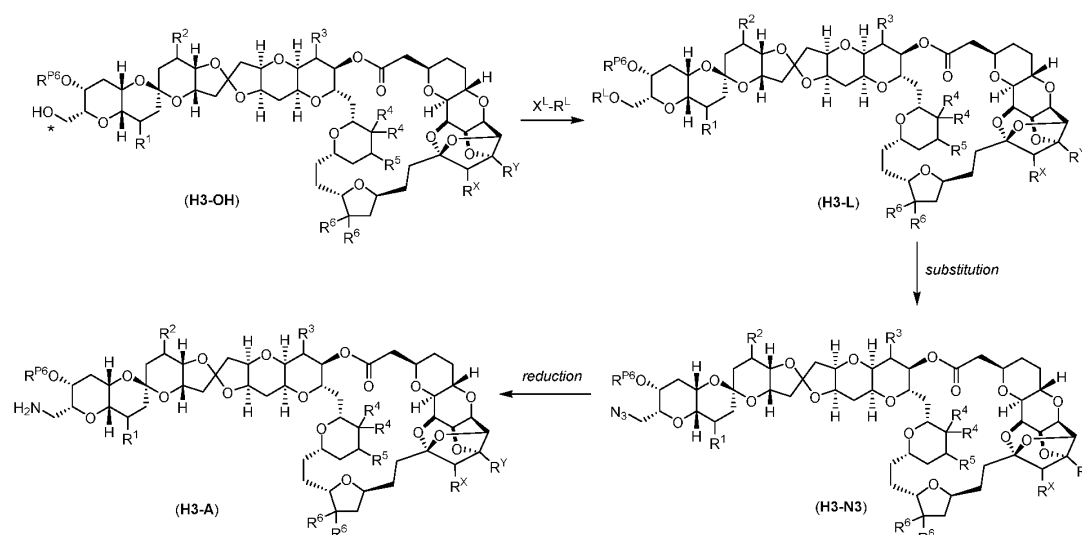


certain embodiments, two R^{P6} are joined to form: ; R^{P4} and R^{P5} are TES; and the step of deprotecting (to remove R^{P6} , R^{P4} , and R^{P5}) is carried out in the presence of TBAF and imidazole hydrochloride. In certain embodiments, the reaction is carried out in a solvent such as THF.

[00189] Once a compound of Formula (H3-2-I), (E-1), or salt thereof, is obtained, the method may comprise one or more additional steps (*e.g.*, deprotection, protection, substitution, addition, elimination) to yield a desired compound.

Synthesis of Amino Analogs of Halichondrins

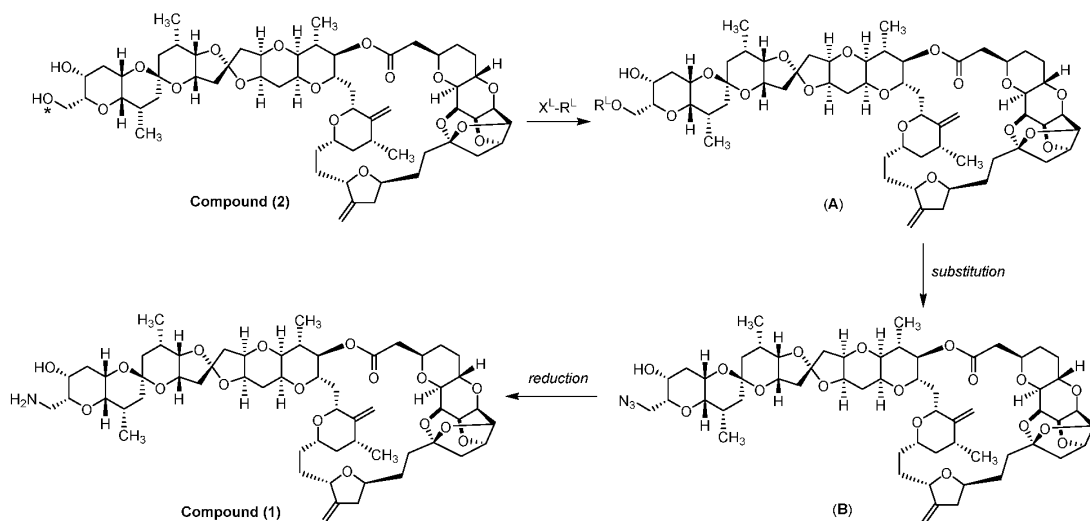
[00190] Provided herein are methods for preparing amino analogs of halichondrins, such as compound of Formula (H3-A). For example, as shown below in *Scheme 4*, compounds of Formula (H3-A) can be prepared by converting compounds of Formula (H3-OH). The primary hydroxyl group (denoted by * in *Scheme 4*) is converted to a leaving group $-OR^L$ by treatment of a compound of Formula (H3-OH) with a reagent of formula X^L-R^L . The group $-OR^L$ can then be substituted for an amine or amine precursor. In certain embodiments, the method comprises substituting the primary $-OR^L$ group with an azide ($-N_3$) (*i.e.*, to yield a compound of Formula (H3-N3)). The azide moiety can then be reduced to an amine to yield a compound of Formula (H3-A).

Scheme 4

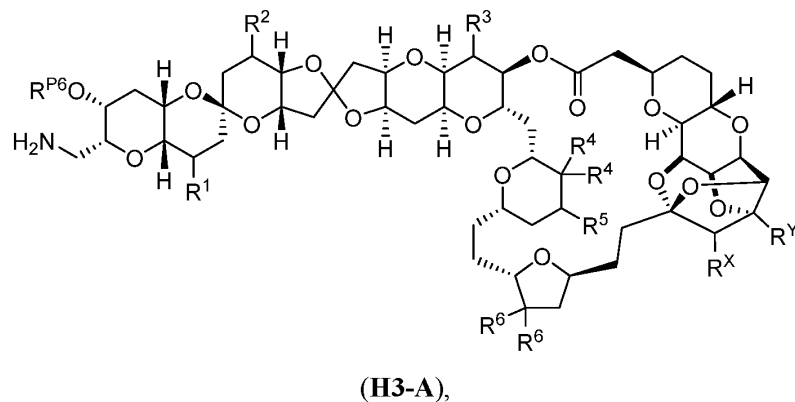
[00191] In certain embodiments, the compound of Formula (H3-A) is a Compound (1), or a salt thereof. Therefore, provided herein are methods for preparing Compound (1) and salts thereof. For example, as shown below in *Scheme 2*, Compound (1) can be prepared by converting Compound (2) to a compound of Formula (A). In this step, the primary hydroxyl group of Compound (2) (denoted by * in *Scheme 2*) is converted to a leaving group $-OR^L$ by treatment of Compound (2) with a reagent of formula X^L-R^L . In certain embodiments, the leaving group is a sulfonate (*i.e.*, R^L is optionally substituted sulfonyl). The group $-OR^L$ can then be substituted for an amine or amine precursor. In certain embodiments, the method comprises substituting the primary $-OR^L$ group with an azide ($-N_3$) (*i.e.*, to yield a compound

of Formula (B)). The azide moiety of a compound of Formula (B) can then be reduced to an amine to yield Compound (1).

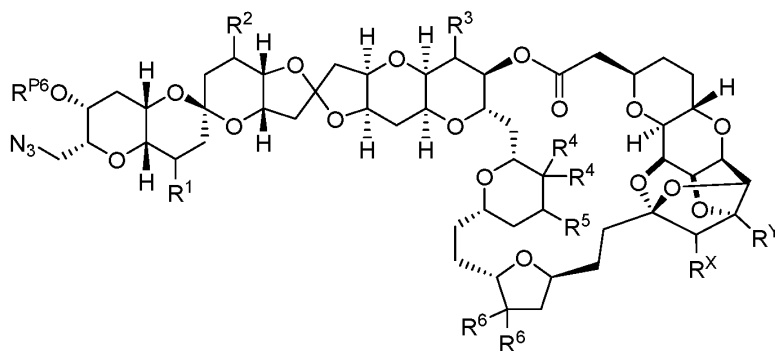
Scheme 2



[00192] As shown in *Scheme 4* above, provided herein is a method of preparing a compound of Formula (H3-A):



or a salt thereof, the method comprising a step of reducing a compound of Formula (H3-N3):



(H3-N3),

or a salt thereof, wherein:

R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R⁴ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁴ groups are taken together to form: ;

each instance of R⁶ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁶ groups are taken together to form: ;

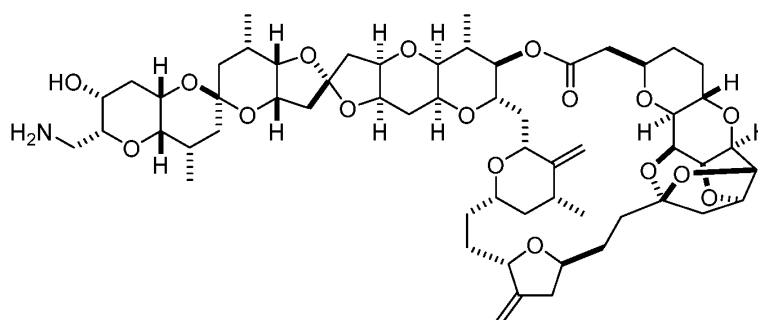
R^{P4}, R^{P5}, and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or -OR^{Xa}, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or -OR^{Y_a}, wherein R^{Y_a} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

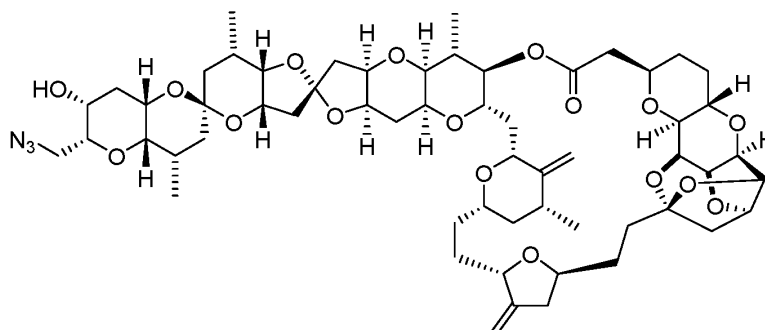
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00193] In certain embodiments, as shown in *Scheme 2*, the method provided herein is a method for preparing Compound (1):



Compound (1),

or a salt thereof, the method comprising reducing a compound of Formula (B):



(B),

or a salt thereof.

[00194] The step of reducing to form a compound of Formula (**H3-A**), Compound (1), or a salt thereof, may be carried out in the presence of any reagents or conditions capable of reducing an azide to an amine (see, *e.g.*, *Chem. Rev.*, 1988, 88 (2), pp 297–368). In certain embodiments, the step of reducing is carried out in the presence of a phosphine reagent (*i.e.*, the Staudinger reaction). In certain embodiments, the phosphine is a trialkylphosphine. In certain embodiments, the phosphine is a triarylphosphine. In certain embodiments, the phosphine is triphenylphosphine (Ph₃P). In certain embodiments, the phosphine reagent is polymer-bound phosphine. In certain embodiments, the phosphine reagent is polymer-bound triphenylphosphine. In certain embodiments, treatment with the phosphine is followed by treatment with water, *e.g.*, an aqueous work-up.

[00195] In certain embodiments, approximately 1 equivalent of the phosphine reagent is used. In certain embodiments, greater than 1 equivalent of the phosphine reagent is used. In certain embodiments, approximately 1-10 equivalents of the phosphine reagent is used. In

certain embodiments, approximately 1-5 equivalents of the phosphine reagent is used. In certain embodiments, approximately 3 equivalents of the phosphine is used. In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the reaction is carried out in THF. In certain embodiments, the reaction is carried out in THF and water. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reactions is carried out at around 25 °C.

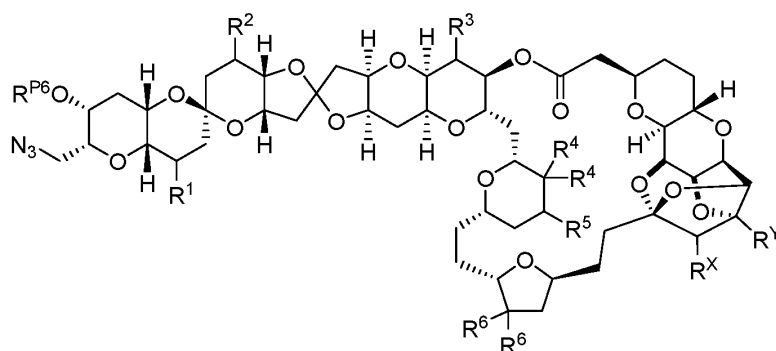
[00196] In certain embodiments, the reaction is carried out in the presence of polymer-bound PPh_3 in THF and water. In certain embodiments, the reaction is carried out under the following conditions: 3 equivalents polymer-bound PPh_3 in THF and water at around 25 °C (*e.g.*, for 70 hours).

[00197] In certain embodiments, the product is purified and isolated by precipitation. In certain embodiments, the product is purified by column chromatography. In certain embodiments, the product is isolated and purified using a combination of column chromatography and precipitation.

[00198] In certain embodiments, R^{P6} is hydrogen. In certain embodiments, R^{P6} is an oxygen protecting group. In certain embodiments, R^{P6} is a silyl protecting group. In certain embodiments, one or more free hydroxyl groups of Compound (B) and Compound (1) is substituted with an oxygen protecting group (*e.g.*, a silyl protecting group).

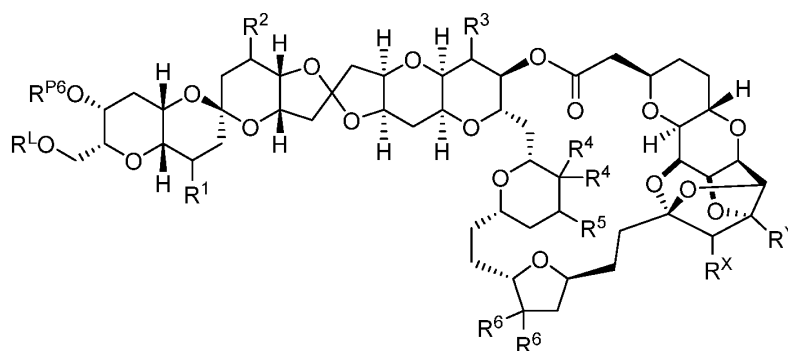
[00199] Other reagents and conditions may be used to convert the azide of Compound (B), or a compound of Formula (**H3-N3**), to an amine. For example, in certain embodiments, the step of reducing is carried out in the presence of palladium and hydrogen (*e.g.*, Pd/C and H_2). In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H^-) source.

[00200] As shown in *Scheme 4*, also provided herein is a method of preparing a compound of Formula (H3-N3):



(H3-N3),

or a salt thereof, the method comprising a step of reacting a compound of Formula (H3-L):



(H3-L),

or a salt thereof, in the presence of an azide, to yield a compound of Formula (H3-N3), or a salt thereof, wherein:

R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

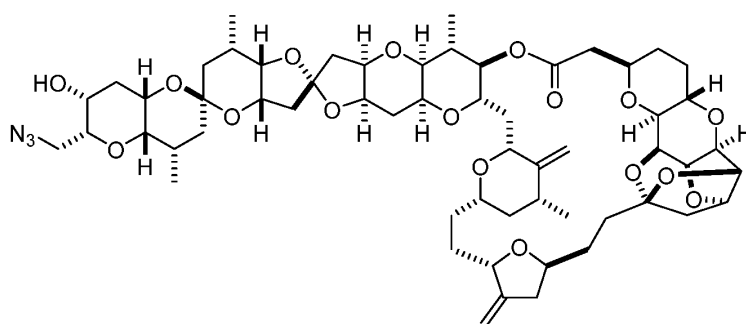
alkyl, or two R^6 groups are taken together to form: ;

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

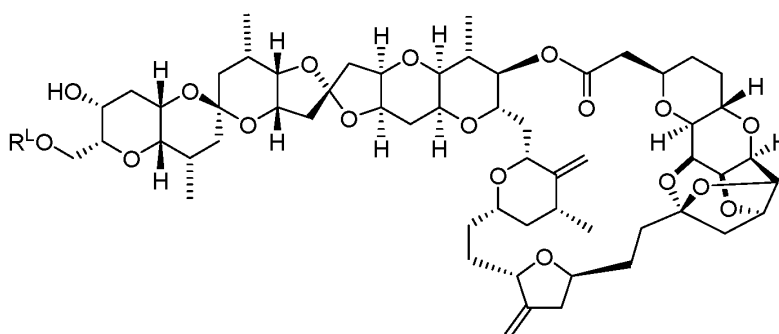
R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocycl.

[00201] In certain embodiments, as shown in *Scheme 2*, the method is a method of preparing a compound of Formula (B):



(B),

or a salt thereof, the method comprising reacting a compound of Formula (A):



(A),

or a salt thereof, in the presence of an azide, wherein:

R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

[00202] The reaction to form a compound of Formula (H3-N3), Compound (B), or a salt thereof, is carried out in the presence of an azide. In certain embodiments, the azide is an azide salt. In certain embodiments, the azide sodium azide (NaN_3) or potassium azide (KN_3).

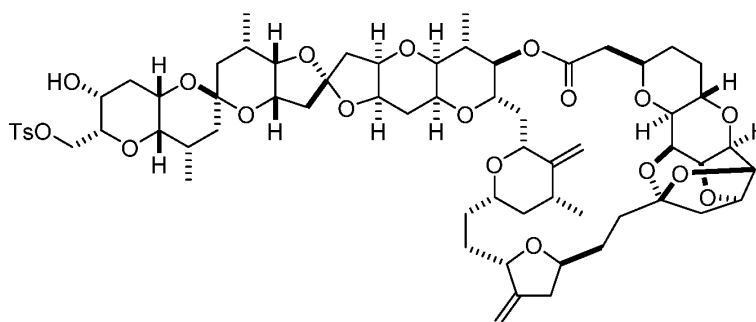
In certain embodiments, the azide is a tetraalkylammonium azide (*i.e.*, [(alkyl)₄N]N₃). In certain embodiments, the azide is tetrabutylammonium azide ([*n*-Bu₄N]N₃). In certain embodiments, approximately 1 equivalent of the azide is present. In certain embodiments, greater than 1 equivalent of the azide is present. In certain embodiments, approximately 1-10 equivalents of azide are present. In certain embodiments, approximately 5-10 equivalents are present. In certain embodiments, approximately 8 equivalents of azide is present.

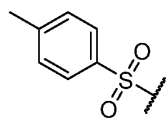
[00203] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is a polar solvent. In certain embodiments, the solvent is an apolar solvent. In certain embodiments, the solvent is toluene. In certain embodiments, the reaction is carried out at above room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from room temperature to approximately 150 °C. In certain embodiments, the reaction is carried out at approximately 100 °C.

[00204] In certain embodiments, the reaction is carried out in the presence of tetrabutylammonium azide ([*n*-Bu₄N]N₃) in toluene. In certain embodiments, the reaction is carried out in the presence of tetrabutylammonium azide ([*n*-Bu₄N]N₃) in toluene at approximately 100 °C. In certain embodiments, the reaction is carried out under the following conditions: 8 equivalents of tetrabutylammonium azide ([*n*-Bu₄N]N₃) in toluene at approximately 100 °C (*e.g.*, for 5 hours).

[00205] In certain embodiments, R^{P6} is hydrogen and R^L is Ts. In certain embodiments, R^{P6} is an oxygen protecting group and R^L is Ts. In certain embodiments, R^{P6} is a silyl protecting group and R^L is Ts. In certain embodiments, one or more free hydroxyl groups of Compound (A) and Compound (B) is substituted with an oxygen protecting group (*e.g.*, a silyl protecting group).

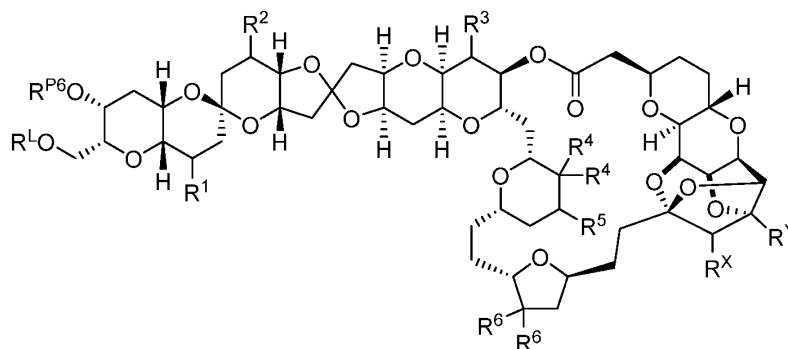
[00206] In certain embodiments, the compound of Formula (A) is the following:





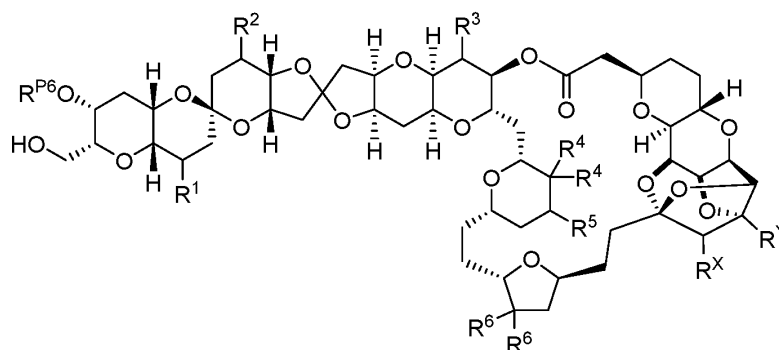
or a salt thereof. “Ts” is a tosyl group of the formula:

[00207] Also provided herein is a method of preparing a compound of Formula (H3-L):



(H3-L),

or a salt thereof, the method comprising a step of reacting a compound of Formula (H3-OH):



(H3-OH),

or a salt thereof, in the presence of a reagent of the formula X^L-R^L , to yield a compound of Formula (H3-L), or a salt thereof, wherein:

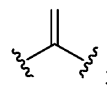
R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl;

X^L is halogen or a leaving group;

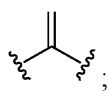
R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R⁴ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁴ groups are taken together to form: ;



each instance of R^6 is independently hydrogen, halogen, or optionally substituted

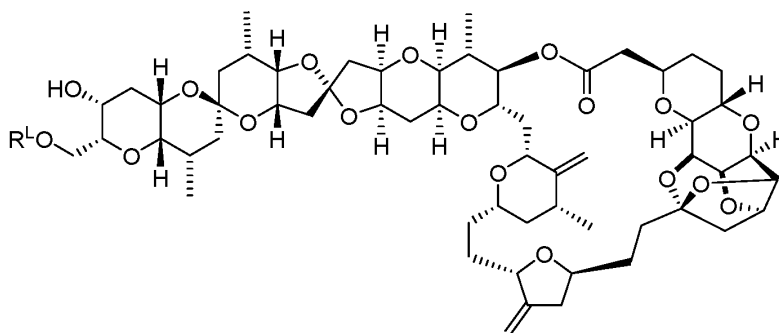
alkyl, or two R^6 groups are taken together to form: ;

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

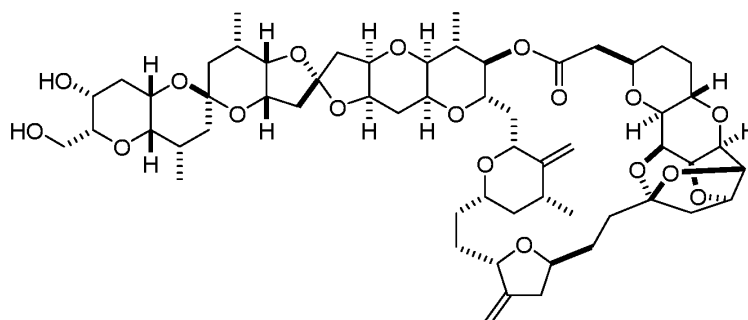
R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocycl.

[00208] In certain embodiments, the method is a method of preparing a compound of Formula (A):



(A),

or a salt thereof, the method comprising reacting Compound (2):



Compound (2),

or a salt thereof, in the presence of a reagent of the formula X^L-R^L , wherein:

X^L is halogen or a leaving group; and

R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

[00209] The reaction to form a compound of Formula (H3-L), Compound (A), or a salt thereof, is carried out in the presence of a reagent of the formula X^L-R^L . The overall transformation converts the primary hydroxyl group of the starting material to a leaving group (*e.g.*, a sulfonyl leaving group) of formula $-OR^L$.

[00210] In certain embodiments, the reagent of the formula X^L-R^L is a sulfonating agent. Sulfonating reagents capable of converting a free hydroxyl group to a sulfonate leaving group are known in the art. In certain embodiments, the reagent of the formula X^L-R^L is a sulfonyl halide (*i.e.*, wherein R^L is optionally substituted sulfonyl). In certain embodiments, the reagent is a tosyl halide (*i.e.*, X^L-Ts). In certain embodiments, the reagent is a sulfonyl chloride (X^L is chlorine and R^L is optionally substituted sulfonyl). In certain embodiments, the reagent is tosyl chloride (TsCl). In certain embodiments, approximately 1 equivalent of the reagent is used. In certain embodiments, greater than 1 equivalent of the reagent is used. In certain embodiments, approximately 3 equivalents of the reagent is used.

[00211] In certain embodiments, the reaction is carried out in the presence of one or more additional reagents. In certain embodiments, the reaction is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amine base. In certain embodiments, the base is a trialkylamine base. Examples of amine bases include, but are not limited to, triethylamine (TEA) and diisopropylethylamine (DIPEA). In certain embodiments, the base is triethylamine (TEA). In certain embodiments, the base is a heterocyclic base. Examples of heterocyclic bases include, but are not limited to, pyridine and imidazole bases. In certain embodiments, approximately 1 equivalent of the base is used. In certain embodiments, greater than 1 equivalent of the base is used. In certain embodiments, an excess (*e.g.*, approximately 6 equivalents) of the base is used.

[00212] In certain embodiments, the reaction is carried out in the presence of a Lewis acid. In certain embodiments, the Lewis acid is dibutyltin oxide. In certain embodiments, the Lewis acid is present in 1 equivalent or less (*e.g.*, 0.5 equivalents).

[00213] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is dichloromethane (DCM). In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out

at around room temperature. In certain embodiments, the reaction is carried out at around 25 °C.

[00214] In certain embodiments, the reaction is carried out in the presence of TsCl, TEA, and a Lewis acid. In certain embodiments, the reaction is carried out in the presence of TsCl, TEA, and dibutyltin oxide. In certain embodiments, the reaction is carried out in the presence of TsCl, TEA, and dibutyltin oxide in DCM. In certain embodiments, the reaction is carried out in the presence of TsCl, TEA, and dibutyltin oxide in DCM at around 25 °C. In certain embodiments, the reaction is carried out under the following conditions: 3 equivalents TsCl, excess TEA (*e.g.*, approximately 6 equivalents), and less than 1 equivalent of dibutyltin oxide (*e.g.*, 0.6 equivalents) in DCM at approximately 25 °C (*e.g.*, for 3 hours).

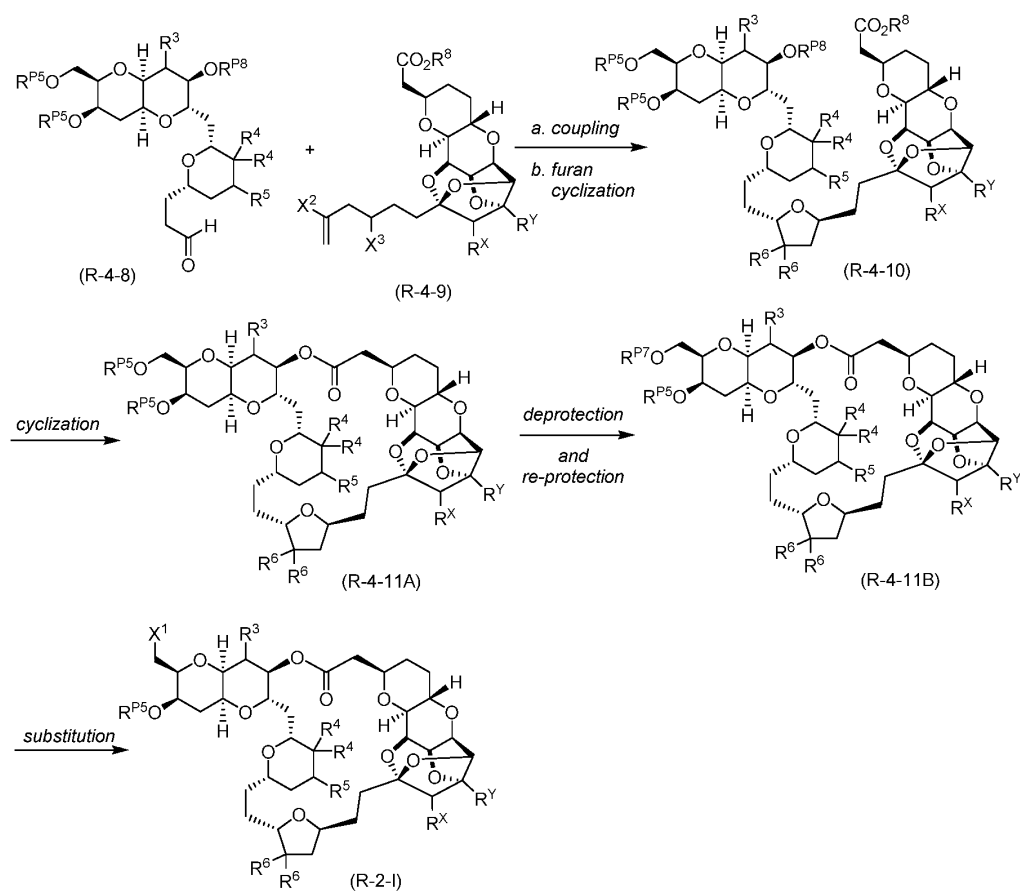
[00215] In certain embodiments, R^{P6} is hydrogen and R^L is Ts. In certain embodiments, R^{P6} is an oxygen protecting group, and R^L is Ts. In certain embodiments, R^{P6} is a silyl protecting group, and R^L is Ts. In certain embodiments, one or more free hydroxyl groups of Compound (A) and Compound (2) is substituted with an oxygen protecting group (*e.g.*, a silyl protecting group).

[00216] Methods for preparing the starting materials (*i.e.*, compounds of Formula (H3-OH), Compound (2), and salts thereof) are provided herein, *e.g.*, under the subsection entitled *Synthesis of Additional Halichondrin Analogs*.

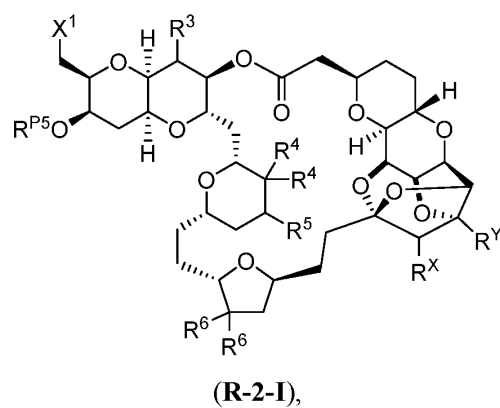
Preparation of “Right Half” Building Blocks

[00217] Also provided herein are methods useful in the preparation of “right half” building blocks of halichondrins (*e.g.*, halichondrin A, B, C; homohalichondrin A, B, C, norhalichondrin A, B, C, and analogs thereof). For example, as described above, compounds of Formula (R-2-I) are useful as right half building blocks. As shown below in *Scheme 3A*, a compound of Formula (R-2-I) can be prepared by substitution of a compound of Formula (R-4-11B) (*i.e.*, substitution of the group –OR^{P7} with the group –X¹). A compound of Formula (R-4-11B) can be prepared by deprotecting and re-protecting one or more oxygen atoms of a compound of Formula (R-4-11A), thereby converting one occurrence of the group –OR^{P5} to the group –OR^{P7}). As also shown in *Scheme 3A*, a compound of Formula (R-4-11) can be prepared by cyclizing a compound of Formula (R-4-10). Furthermore, a compound of Formula (R-4-10) can be obtained by coupling a compound of Formula (R-4-8) with a compound of Formula (R-4-9).

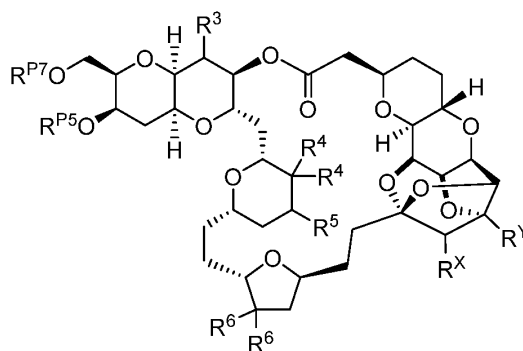
Scheme 3A



[00218] As shown in *Scheme 3A*, provided herein is a method of preparing a compound of Formula (R-2-I):



or a salt thereof, the method comprising reacting a compound of Formula (R-4-11B):



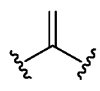
(R-4-11B),

or a salt thereof, in the presence of a nucleophile, thereby substituting the group $-\text{OR}^{\text{P}7}$ with the group $-\text{X}^1$; wherein:

X^1 is halogen or a leaving group;

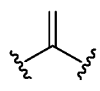
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form:



each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form:



$\text{R}^{\text{P}5}$ is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

$\text{R}^{\text{P}7}$ is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group;

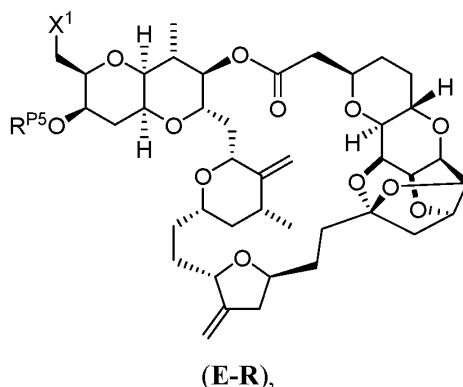
optionally wherein $\text{R}^{\text{P}5}$ and $\text{R}^{\text{P}7}$ are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^{X} is hydrogen or $-\text{OR}^{\text{Xa}}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

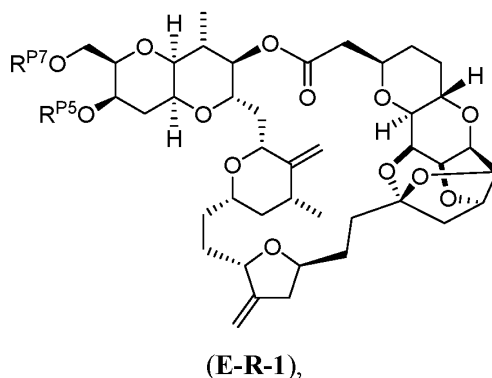
R^{Y} is hydrogen or $-\text{OR}^{\text{Ya}}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00219] In certain embodiments, method is a method of preparing a compound of Formula (E-R):



or a salt thereof, the method comprising reacting a compound of Formula (E-R-1):



or a salt thereof, in the presence of a nucleophile, thereby substituting the group $-OR^{P7}$ with the group $-X^1$; wherein:

X^1 is halogen or a leaving group;

R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{P5} and R^{P7} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00220] As described above, the method of preparing a compound of Formula (R-2-I), (E-R), or a salt thereof, comprises a step of reacting a compound of Formula (R-4-11B), or a salt thereof, in the presence of a nucleophile, thereby substituting the leaving group $-OR^{P7}$ with the group $-X^1$. In certain embodiments, the nucleophile is a halide anion (*e.g.*, Cl^- , Br^- , I^- , F^-). In certain embodiments, the reaction is carried out in the presence of a halide salt. In certain

embodiments, the reaction is carried out in the presence of an iodide salt (*e.g.*, NaI, KI), thereby substituting the leaving group $-\text{OR}^{\text{P7}}$ with the group $-\text{I}$. In certain embodiments, the iodide salt is sodium iodide (NaI). In certain embodiments, the reaction is carried out in the presence of NaI. In certain embodiments, the reaction is carried out in a polar solvent (*e.g.*, DMF or DMI). In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature.

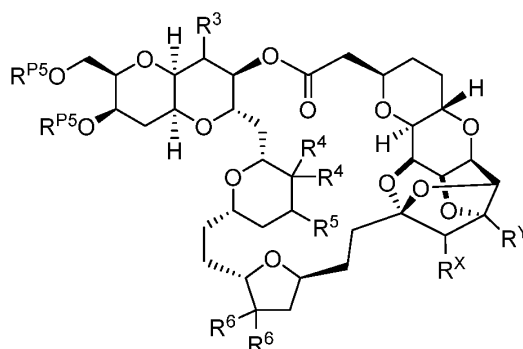
[00221] In certain embodiments, the reaction is carried out in the presence of NaI in DMI at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 5 equivalents of NaI in DMF at room temperature (*e.g.*, for 2-5 hours). For example, in certain embodiments, the reaction is carried out under the following conditions: 5 equivalents of NaI in DMI at room temperature (*e.g.*, for 2-5 hours).

[00222] In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is a leaving group. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{O-sulfonyl}$. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{OMs}$. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{OTs}$. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{OTf}$. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{O-acyl}$. In certain embodiments, the group $-\text{OR}^{\text{P7}}$ is $-\text{O-phosphoryl}$. In certain embodiments, R^{P5} is a silyl protecting group. In certain embodiments, R^{P5} is TES. In certain embodiments, $-\text{OR}^{\text{P7}}$ is $-\text{OTf}$ and R^{P5} is TES.

[00223] As shown in *Scheme 3A*, a compound of Formula (**R-4-11B**) can be prepared by deprotecting and re-protecting one or more oxygen atoms of a compound of Formula (**R-4-11A**), thereby converting one occurrence of the group $-\text{OR}^{\text{P5}}$ to the group $-\text{OR}^{\text{P7}}$.

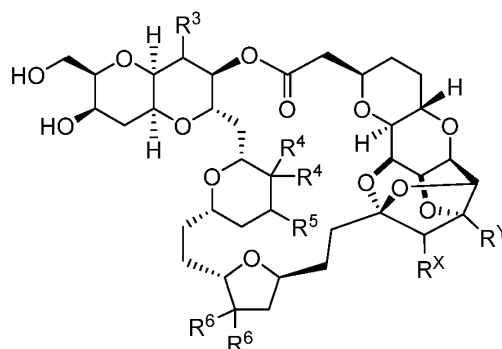
[00224] For example, in certain embodiments, provided herein is a method of preparing a compound of Formula (**R-4-11B**), or a salt thereof, the method comprising:

(a) a step of deprotecting a compound of Formula (**R-4-11A**):



(**R-4-11A**),

or a salt thereof, to yield a compound of Formula (R-4-11C):

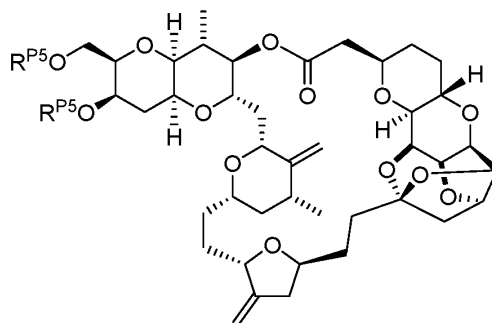


(R-4-11C),

or a salt thereof, following by (b) one or more steps of re-protecting the compound of Formula (R-4-11C), or a salt thereof, to yield a compound of Formula (R-4-11B), or a salt thereof.

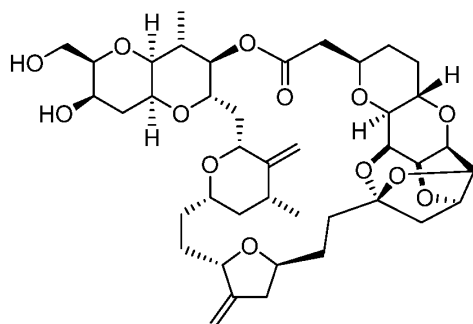
[00225] In certain embodiments, the method comprises:

(a) a step of deprotecting a compound of Formula (E-R-2):



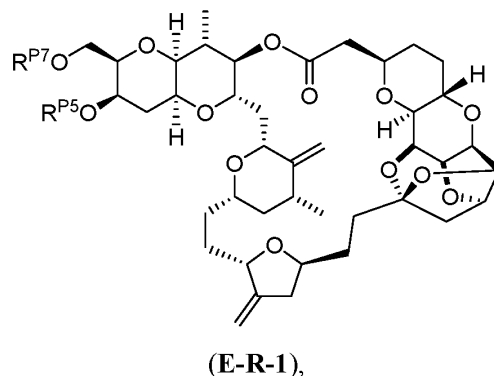
(E-R-2),

or a salt thereof, to yield a compound of the formula:



or a salt thereof; and

(b) one or more steps of re-protecting the product of step (a) to yield a compound of Formula (E-R-1):



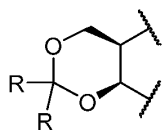
or a salt thereof, wherein:

R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

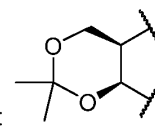
R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{P5} and R^{P7} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00226] As shown above, the compounds of Formula (R-4-11A) and (E-R-2) can be deprotected to remove the groups R^{P5} (*i.e.*, step (a)). In certain embodiments, the R^{P5} groups are silyl protecting groups; and step (a) is carried out in the presence of a fluoride source. In certain embodiments, the fluoride source is tetrabutylammonium fluoride (TBAF). In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the formula:



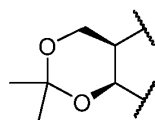
; and step (a) is carried out in the presence of an acid. In certain embodiments,



two R^{P5} are joined with the intervening atoms to form a ring of the formula: ; and step (a) is carried out in the presence of an acid. In certain embodiments, the acid is *p*-toluenesulfonic acid (TsOH). In certain embodiments, the acid is *p*-toluenesulfonic acid monohydrate (TsOH•H₂O). In certain embodiments, the acid is present in a catalytic amount.

[00227] In certain embodiments, the step of deprotecting is carried out in DCM and an alcohol (*e.g.*, ROH). In certain embodiments, the deprotection is carried out in DCM and MeOH. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the deprotection is carried out at around room temperature. In certain embodiments, the deprotection is carried out at around 25 °C.

[00228] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring



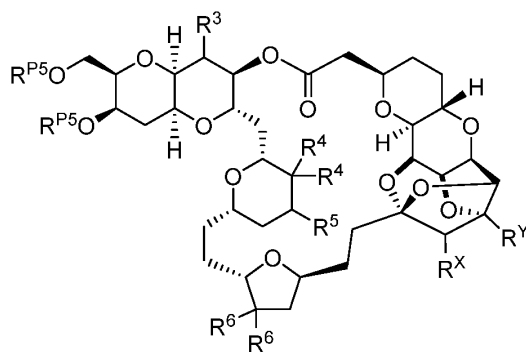
of the formula: ; and the deprotection is carried out in the presence of TsOH•H₂O in DCM and an alcohol. In certain embodiments, the deprotection is carried out under the following conditions: catalytic TsOH•H₂O (*e.g.*, 0.02 equiv) in DCM and MeOH at around 25 °C (*e.g.*, for 4 hours).

[00229] In certain embodiments, on compound of Formula (R-4-11B) or (E-R-1), -OR^{P7} is a sulfonate leaving group and R^{P5} is a silyl protecting group; and step (b) is carried out in the presence of a sulfonating reagent and a base (thereby installing R^{P7} as a sulfonyl group), followed by a silylating reagent and a base (thereby installing R^{P5} as a silyl group). In certain embodiments, the sulfonating reagent is a triflating agent. In certain embodiments, the sulfonating reagent is Tf₂O. In certain embodiments, the silylating reagent is TESOTf. In certain embodiments, the base is an amine or pyridine base. In certain embodiments, the base is 2,4,6-collidine.

[00230] In certain embodiments, the steps of protecting are carried out in a solvent. In certain embodiments, the solvent is DCM. In certain embodiments, the steps of protecting are carried out at below room temperature (*e.g.*, from about -78 °C to -40 °C; from about -78 °C to 0 °C; from about -78 °C to room temperature).

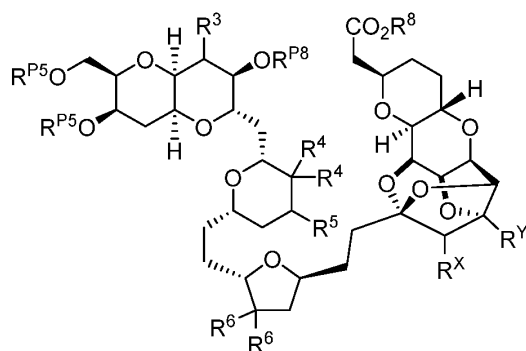
[00231] In certain embodiments, -OR^{P7} is -OTf and R^{P5} is TES; and step (b) is carried out in the presence of Tf₂O and a base, followed by TESOTf and a base. In certain embodiments, the reaction is carried out in the presence of Tf₂O and 2,4,6-collidine in DCM, followed by addition of TESOTf. In certain embodiments, the reaction is carried out under the following conditions: approximately 1.4 equivalents of Tf₂O and 5 equivalents of 2,4,6-collidine in DCM at around -78 °C, followed by addition of 1.4 equivalents of TESOTf and warming to around -40 °C.

[00232] As shown in *Scheme 3A*, also provided herein is a method of preparing a compound of Formula (R-4-11A):



(R-4-11A),

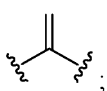
or a salt thereof, the method comprising cyclizing a compound of Formula (R-4-10):



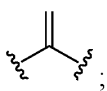
(R-4-10),

or a salt thereof, wherein:

R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, optionally substituted alkyl,

or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, optionally substituted alkyl,

or two R^6 groups are taken together to form: ;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally, wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring;

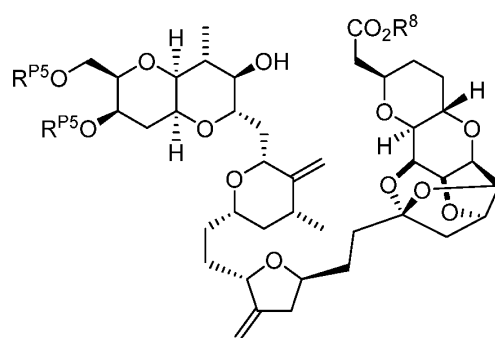
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

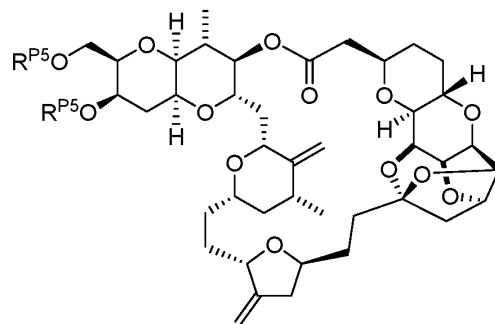
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00233] In certain embodiments, the method comprises cyclizing a compound of Formula **(E-R-3)**:



(E-R-3),

or a salt thereof, to yield a compound of Formula **(E-R-2)**:



(E-R-2),

or a salt thereof, wherein:

each instance of R^{P5} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R⁸ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00234] In certain embodiments, the step of cyclizing a compound of Formula (**R-4-10**), (**E-R-3**), or a salt thereof, is carried out in the presence of an anhydride reagent. In certain embodiments, the anhydride reagent is a benzoic anhydride. In certain embodiments, the reagent is a nitrobenzoic anhydride. In certain embodiments, the anhydride is 2-methyl-6-nitrobenzoic anhydride (MNBA). The anhydride reagent may be present in a catalytic, stoichiometric, or excess amount. In certain embodiments, the anhydride reagent is present in excess (*i.e.*, greater than 1 equivalent) relative to a compound of Formula (**R-4-10**) or (**E-R-3**). In certain embodiments, the anhydride is present in approximately 3 equivalents.

[00235] In certain embodiments, the reaction is carried out in the presence of a nucleophilic reagent capable of activating the carboxyl group $-\text{CO}_2\text{R}^8$ or $-\text{CO}_2\text{H}$. In certain embodiments, the nucleophilic reagent is a pyridine. In certain embodiments, the nucleophilic reagent is 4-dimethylaminopyridine (DMAP). In certain embodiments, the nucleophilic reagent is present in excess (*i.e.*, greater than 1 equivalent) relative to a compound of Formula (**R-4-10**) or (**E-R-3**). In certain embodiments, the reagent is present in approximately 6 equivalents.

[00236] In certain embodiments, the step of cyclizing is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amine base. In certain embodiments, the base is a trialkylamine base (*e.g.*, trimethylamine, triethylamine, tributylamine, diisopropyl ethylamine). In certain embodiments, the base is a heteroaryl base (*e.g.*, a pyridine base, an imidazole base). In certain embodiments, the base is diisopropyl ethylamine (DIPEA). In certain embodiments, the base is present in excess (*i.e.*, greater than 1 equivalent) relative to a compound of Formula (**R-4-10**). In certain embodiments, the base is present in approximately 6 equivalents.

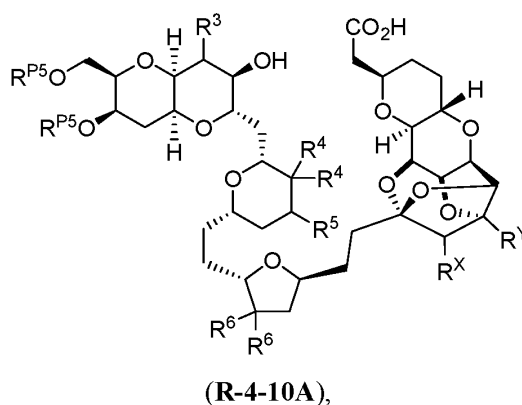
[00237] In certain embodiments, the step of cyclizing is carried out in a solvent (*e.g.*, toluene). In certain embodiments, the reaction is carried out at above room temperature. In certain embodiments, the deprotection is carried out in DCM and MeOH. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at approximately 70 °C or 80 °C.

[00238] In certain embodiments, the step of cyclizing is carried out in the presence of an anhydride reagent, a nucleophilic reagent, and a base. In certain embodiments, the anhydride

reagent is 2-methyl-6-nitrobenzoic anhydride. In certain embodiments, the nucleophilic reagent is DMAP. In certain embodiments, the base is a trialkylamine base such as DIPEA. In certain embodiments, the step is carried out in the presence of 2-methyl-6-nitrobenzoic anhydride (MNBA), 4-dimethylaminopyridine (DMAP), and diisopropyl ethylamine (DIPEA).

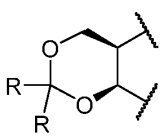
[00239] For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 6 equivalents MNBA, 12 equivalents DMAP, and 6 equivalents DIPEA, in toluene at around 70 °C. For example, in certain embodiments, the step of cyclizing is carried out under the following conditions: 3 equivalents MNBA, 6 equivalents DMAP, and 6 equivalents DIPEA, in toluene at around 80 °C (*e.g.*, for 6 hours). In certain embodiments, the reaction entails slow addition (*i.e.*, dropwise addition) of the compound of Formula (R-4-10) or (E-R-3), or salt thereof, to the reaction mixture.

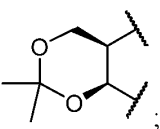
[00240] In certain embodiments, the compound of Formula (R-4-10) is of the Formula (R-4-10A):



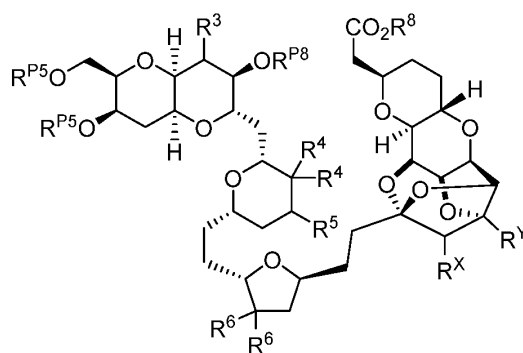
or a salt thereof.

[00241] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

of the formula: ; and R⁸ is hydrogen. In certain embodiments, two R^{P5} are

joined with the intervening atoms to form a ring of the formula: ; and R⁸ is hydrogen.

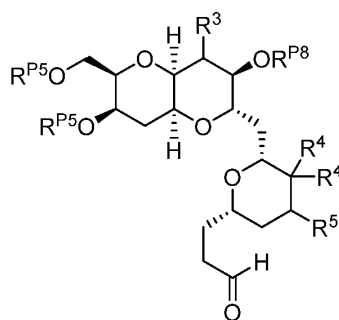
[00242] Also provided herein is a method of preparing a compound of Formula (R-4-10):



(R-4-10),

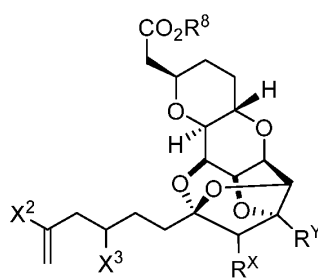
or a salt thereof, the method comprising the steps of:

(a) coupling a compound of Formula (R-4-8):



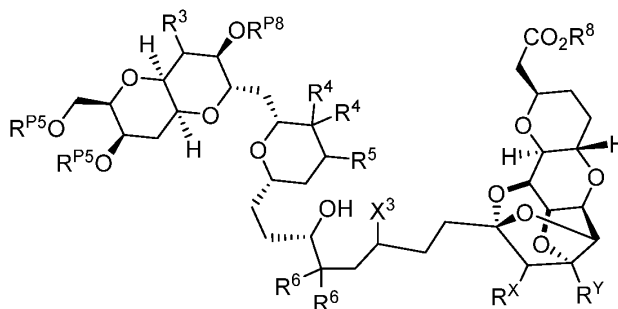
(R-4-8),

or a salt thereof, with a compound of Formula (R-4-9):



(R-4-9),

or a salt thereof, to yield a compound of Formula (R-4-10B):



(R-4-10B),

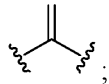
or a salt thereof, followed by

(b) cyclizing a compound of Formula (R-4-10B), or a salt thereof, to yield a compound of Formula (R-4-10), or a salt thereof, wherein:

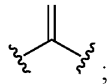
X^3 and X^2 are each independently halogen or a leaving group;

R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring;

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

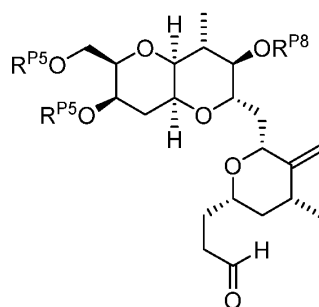
R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

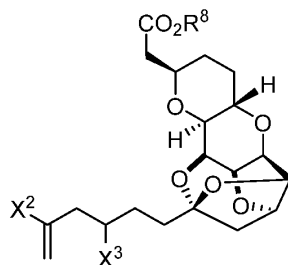
[00243] In certain embodiments, the method comprises:

(a) a step of coupling a compound of Formula (E-R-4):



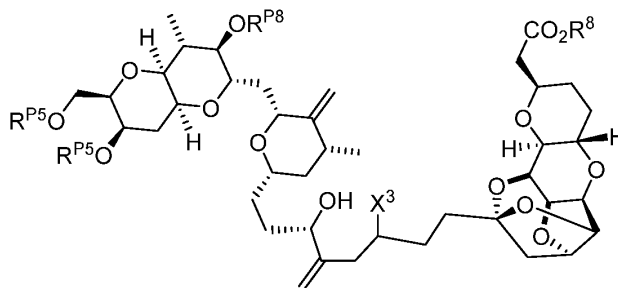
(E-R-4),

or a salt thereof, with a compound of Formula (E-R-5):



(E-R-5),

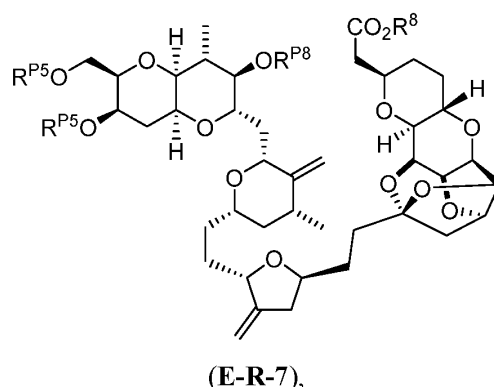
or a salt thereof, to yield a compound of Formula (E-R-6):



(E-R-6),

or a salt thereof, followed by

(b) a step of cyclizing a compound of Formula (E-R-6), or a salt thereof, to yield a compound of Formula (E-R-7):



or a salt thereof, or a salt thereof, wherein:

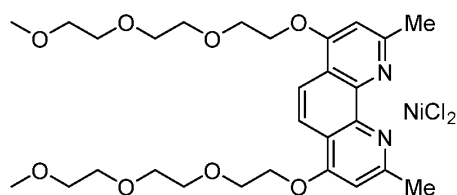
X^3 and X^2 are each independently halogen or a leaving group;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

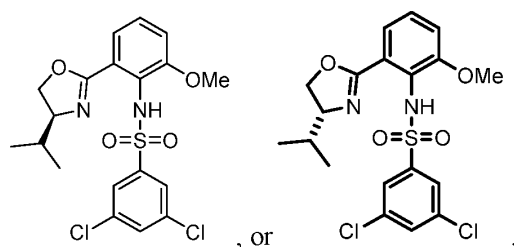
[00244] In certain embodiments, step (a) above (to prepare a compound of Formula (R-4-10B), (E-R-6), or a salt thereof) is a Ni/Cr-mediated reductive coupling reaction; and step (b) above (to prepare a compound of Formula (R-4-10), (E-R-7), or a salt thereof) is an acid-promoted or base-promoted intramolecular furan cyclization. Reagents and conditions for steps (a) and (b) can be found in, *e.g.*, international PCT application publications, WO 2016/176560, published November 3, 2016, and WO 2016/003975, published January 7, 2016, the entire contents of which is incorporated herein by reference.

[00245] The Ni/Cr-mediated reductive coupling (*i.e.*, step (a)) is carried out in the presence of nickel and chromium. In certain embodiments, the nickel is a nickel complex. Examples of nickel complexes include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the nickel complex is $(Et)_2Phen\bullet NiCl_2$. In certain embodiments, the nickel complex is the following:



In certain embodiments, the nickel complex is present in a catalytic amount.

[00246] In certain embodiments, the chromium is a chromium complex. In certain embodiments, the chromium complex is prepared from a chromium salt and a chiral ligand. In certain embodiments, the chromium salt is CrCl_2 or CrCl_3 . In certain embodiments, the chiral ligand is a chiral sulfonamide. Examples of chiral ligands include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the chiral ligand is (*S*)-**4-G**. In certain embodiments, the chiral sulfonamide ligand is one of the following:



or a salt thereof. In certain embodiments, the chromium complex is present in a catalytic amount.

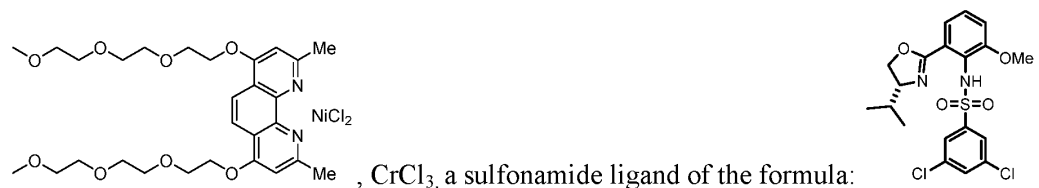
[00247] The Ni/Cr-mediated reductive coupling may be carried out in the presence of one or more additional reagents. In certain embodiments, the coupling is carried out in the presence of a lithium salt (*e.g.*, LiCl or LiBr). In certain embodiments, the coupling is carried out in the presence of a reducing metal such as zinc or manganese (*e.g.*, zinc or manganese metal). In certain embodiments, the coupling is carried out in the presence of zirconium (*e.g.*, ZrCp_2Cl_2). In certain embodiments, the reducing metal is zinc metal. In certain embodiments, the metal is manganese metal. In certain embodiments, the coupling is carried out in the presence of a base or proton scavenger (*e.g.*, 2,6-di-*tert*-butyl-4-methylpyridine). In certain embodiments, the coupling is carried out in the presence of proton sponge (*e.g.*, 1,8-bis(dimethylamino)naphthalene).

[00248] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is acetonitrile (MeCN). In certain embodiments, the deprotection is carried out in DCM and MeOH. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100°C . In certain embodiments, the reaction is carried out at a temperature ranging from approximately

room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 30 °C.

[00249] In certain embodiments, the Ni/Cr-mediated reductive coupling is carried out in the presence of a nickel complex, a chromium salt, a sulfonamide ligand, a lithium salt, a zirconium complex, a reducing metal, and a base or proton scavenger. In certain embodiments, the step of coupling is carried out in the presence of (Et)₂Phen•NiCl₂, CrCl₂, (S)-**4-G**, LiCl, ZrCp₂Cl₂, manganese metal, and a base or proton scavenger (*e.g.*, 2,6-di-*tert*-butyl-4-methylpyridine). For example, in certain embodiments, the reaction is carried out under the following conditions: 2 mol% (Et)₂Phen•NiCl₂, 10 mol% CrCl₂, 10 mol% ligand (S)-**4-G**, 2 equivalents LiCl, 2.5 equivalents ZrCp₂Cl₂, excess manganese metal, and 2.5 equivalents 2,6-di-*tert*-butyl-4-methylpyridine, in MeCN at room temperature (*e.g.*, for 2 hours).

[00250] In certain embodiments, the Ni/Cr-mediated reductive coupling is carried out in the presence of a nickel complex, a chromium salt, a sulfonamide ligand, a zirconium complex, a reducing metal, and a base or proton scavenger. In certain embodiments, the coupling is carried out in the presence of: a nickel complex of the formula:



Cp₂ZrCl₂, manganese metal, and a base or proton scavenger (*e.g.*, 2,6-di-*tert*-butyl-4-methylpyridine and/or proton sponge (*e.g.*, 1,8-bis(dimethylamino)naphthalene)). In certain embodiments, the reaction is carried out in MeCN. In certain embodiments, the reaction is carried out at around 30 °C. For example, the coupling can be carried out under the following

conditions: 3 mol% of a nickel complex of the formula:

The nickel complex is the same as shown above, consisting of a 1,10-bis(4-methoxyphenyl)-2,9-bis(2-methoxyethoxy)anthracene-2,9-dimethanol ligand coordinated to NiCl₂.

20 mol% CrCl₃, 20 mol% of a sulfonamide ligand of the formula:

The sulfonamide ligand is (S)-1-(4-chlorophenyl)-2-methoxy-2-oxo-1,2,3,4-tetrahydro-1H-benzothiazole-5-carboxamide.

equivalents Cp_2ZrCl_2 , 2 equivalents manganese metal, and 2 equivalents of 2,6-di-*tert*-butyl-4-methylpyridine, and proton sponge in MeCN at around 30 °C.

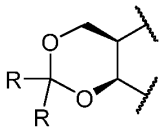
[00251] In certain embodiments, step (b) (to prepare a compound of Formula (R-4-10), (E-R-7), or a salt thereof) is carried out in the presence of a Lewis acid. In certain embodiments, the Lewis acid is AgOTf. In certain embodiments, the Lewis acid is Ag_2O . In certain embodiments, the Lewis acid is SrCO_3 . The Lewis acid may be present in a catalytic, stoichiometric, or excess amount. In other embodiments, step (b) is carried out in the presence of a base. In certain embodiments, the base is a carbonate salt. In certain embodiments, the base is potassium carbonate (K_2CO_3).

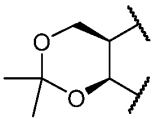
[00252] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is MeOH. In certain embodiments, the solvent is MeCN. In certain embodiments, the reaction is carried out in MeOH and water. In certain embodiments, the reaction is carried out at above room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at from 50-60 °C. In certain embodiments, the reaction is carried out at around 60 °C. In certain embodiments, the reaction is carried out at around 55 °C.

[00253] In certain embodiments, in addition to affecting the furan cyclization, the reaction conditions are sufficient to hydrolyze the ester $-\text{CO}_2\text{R}^8$ (wherein R^8 is hydrogen in the product (E-R-7) or (R-4-10)).

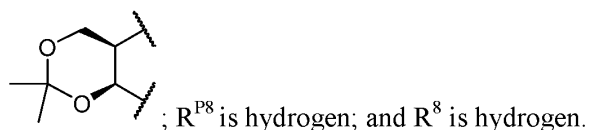
[00254] For example, in certain embodiments, the reaction is carried out under the following conditions: 10 equivalents K_2CO_3 in MeCN at 60 °C (*e.g.*, for 3 hours). In certain embodiments, the reaction is carried out in the presence of K_2CO_3 , in MeOH and water, at around 55 °C. As another example, the reaction can be carried out under the following conditions: 10 equivalents K_2CO_3 in MeOH and water at around 55 °C (*e.g.*, for 23 hours).

[00255] In certain embodiments, two $\text{R}^{\text{P}5}$ are joined with the intervening atoms to form a ring

of the formula: ; $\text{R}^{\text{P}8}$ is hydrogen; and R^8 is optionally substituted alkyl or hydrogen. In certain embodiments, two $\text{R}^{\text{P}5}$ are joined with the intervening atoms to form a

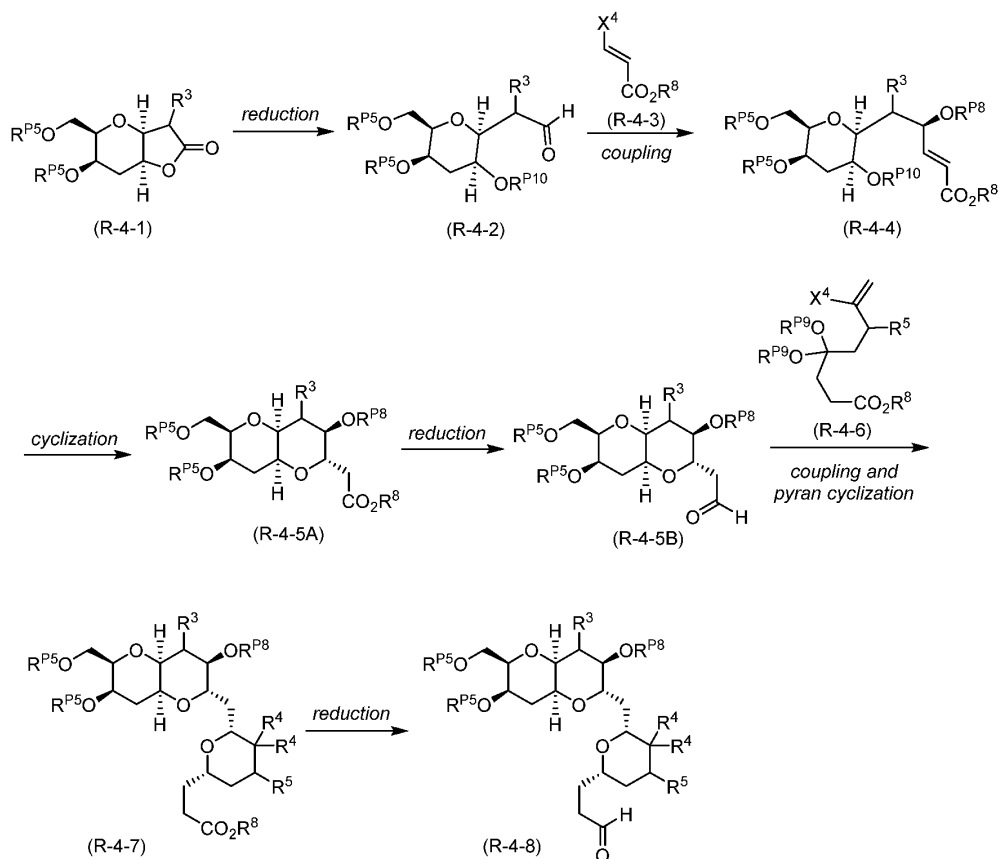
ring of the formula: ; $\text{R}^{\text{P}8}$ is hydrogen; and R^8 is methyl. In certain

embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the formula:

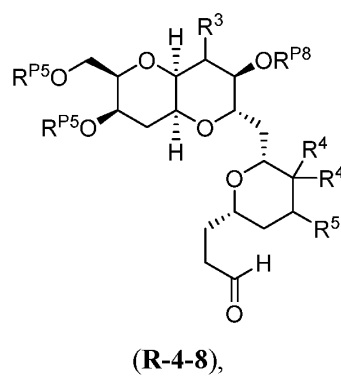


[00256] As shown in *Scheme 3B*, a compound of Formula (**R-4-8**) can be prepared by reducing the ester moiety ($-\text{CO}_2R^8$) of a compound of Formula (**R-4-7**) to an aldehyde moiety. A compound of Formula (**R-4-7**) can be prepared by coupling a compound of Formula (**R-4-5B**) with a compound of Formula (**R-4-6**), followed by formation of the pyran ring via cyclization of the adduct, or a deprotected form of the adduct. In turn, a compound of Formula (**R-4-5B**) can be prepared by reducing the ester moiety ($-\text{CO}_2R^8$) of a compound of Formula (**R-4-5A**) to an aldehyde moiety. A compound of Formula (**R-4-5A**) can be prepared by cyclization of a compound of Formula (**R-4-4**), which can be prepared by coupling a compound of Formula (**R-4-2**) with an olefin of Formula (**R-4-3**). As shown in *Scheme 3B*, a compound of Formula (**R-4-2**) can be prepared by reducing the lactone of a compound of Formula (**R-4-1**).

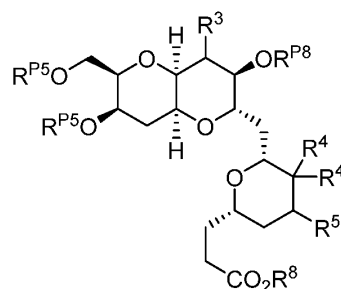
Scheme 3B



[00257] As shown in *Scheme 3B*, provided herein is a method of preparing a compound of Formula **(R-4-8)**:



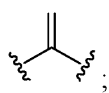
or a salt thereof, the method comprising reducing a compound of Formula (R-4-7):



(R-4-7),

or a salt thereof, wherein:

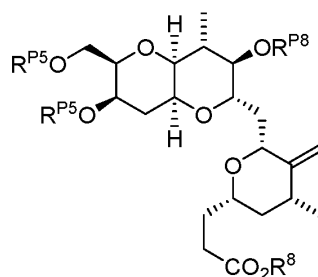
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

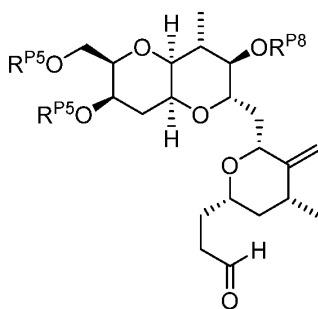
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00258] In certain embodiments, the method comprises a step of reducing a compound of Formula (E-R-8):



(E-R-8),

or a salt thereof, to yield a compound of Formula (E-R-4):



(E-R-4),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

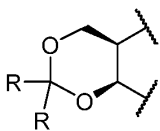
[00259] The step of reducing a compound of Formula (R-4-7), (E-R-8), or a salt thereof, converts the ester group $-CO_2R^8$ to an aldehyde group. In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H^-) source. Any hydride source known in the art may be used in this transformation. Examples of hydride sources include, but are not limited to, lithium aluminum hydride (LAH), sodium borohydride ($NaBH_4$), lithium borohydride, and diisobutylaluminum hydride (DIBAL). In certain embodiments, the hydride source is diisobutylaluminum hydride (DIBAL). In certain embodiments, the hydride source is present in a stoichiometric or excess amount.

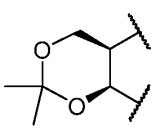
[00260] The step of reducing may optionally comprise reducing the $-CO_2R^8$ moiety to an alcohol, followed by oxidation of the resulting alcohol to an aldehyde to yield a compound of Formula (R-4-7), (E-R-8), or a salt thereof.

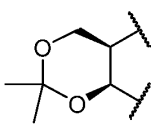
[00261] In certain embodiments, the step of reducing is carried out in the presence of DIBAL. In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is DCM. In certain embodiments, the reaction is carried out at below room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately $-78\text{ }^\circ\text{C}$ to approximately room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately $-78\text{ }^\circ\text{C}$ to approximately $0\text{ }^\circ\text{C}$. In certain embodiments, the reaction is carried out at around $-78\text{ }^\circ\text{C}$.

For example, in certain embodiments, the reaction is carried out under the following conditions: DIBAL in DCM at around -78 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: approximately 2.3 equivalents DIBAL in DCM at around -78 °C (*e.g.*, for 1-2 hours).

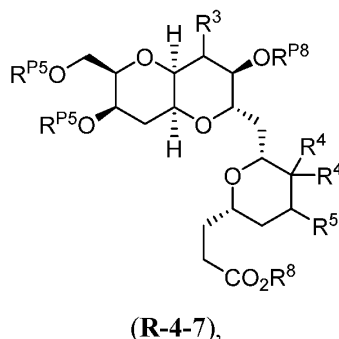
[00262] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

of the formula: ; R^{P8} is hydrogen; and R⁸ is optionally substituted alkyl. In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the

formula: ; R^{P8} is hydrogen, and R⁸ is ethyl. In certain embodiments, two R^{P5} are

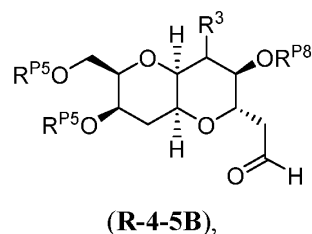
joined with the intervening atoms to form a ring of the formula: ; R^{P8} is hydrogen, and R⁸ is methyl.

[00263] As shown in *Scheme 3B*, also provided herein is a method of preparing a compound of Formula **(R-4-7)**:

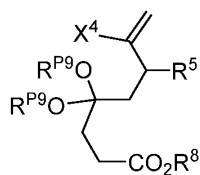


or a salt thereof, the method comprising the steps of:

(a) coupling a compound of Formula **(R-4-5B)**:

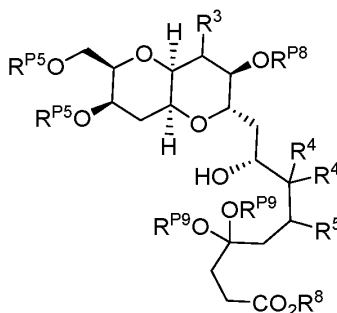


or a salt thereof, with a compound of Formula (R-4-6):



(R-4-6),

or a salt thereof, to yield a compound of Formula (R-4-7A):



(R-4-7A),

or a salt thereof; and

(a-i) deprotecting and cyclizing a compound of Formula (R-4-7A), or a salt thereof, to give a compound of Formula (R-4-7), or a salt thereof; wherein:

R³ and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R⁴ is independently hydrogen, halogen, or optionally substituted

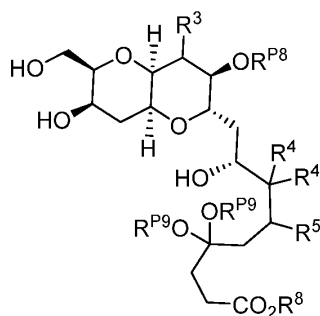
alkyl, or two R⁴ groups are taken together to form: ;

each instance of R^{P5}, R^{P8}, and R^{P9} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and optionally, wherein two R^{P9} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R⁸ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00264] In certain embodiments, after the step of coupling the compounds of Formulae (**R-4-5B**) and (**R-4-6**) (*i.e.*, step (a)), the method comprises:

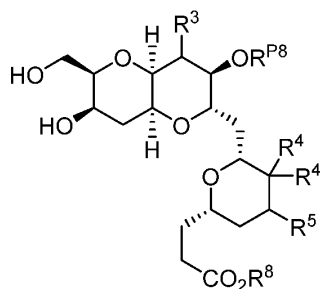
(b) a step of deprotecting a compound of Formula (**R-4-7A**), or a salt thereof, to yield a compound of Formula (**R-4-7B**):



(**R-4-7B**),

or a salt thereof;

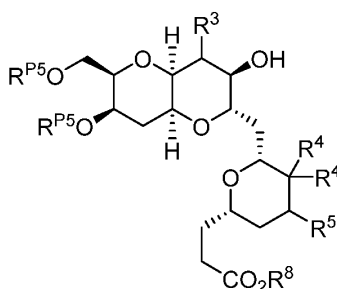
(c) a step of cyclizing to yield a compound of Formula (**R-4-7C**):



(**R-4-7C**),

or a salt thereof; and optionally

(d) a step of re-protecting the compound of Formula (**R-4-7C**), or a salt thereof, at one or more oxygen atoms to yield a compound of Formula to yield a compound of Formula (**R-4-7B**):

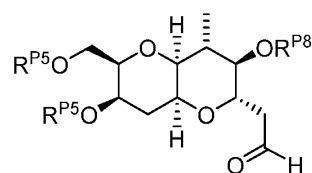


(**R-4-7B**),

or a salt thereof.

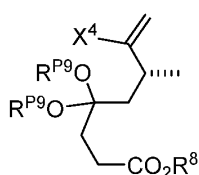
[00265] In certain embodiments, the method comprises:

(a) a step of coupling a compound of Formula (E-R-9):



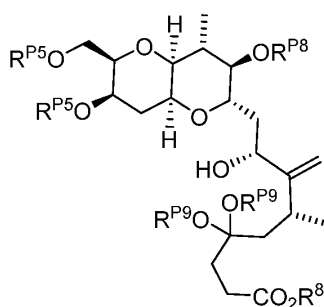
(E-R-9),

or a salt thereof, with a compound of Formula (E-R-10):



(E-R-10),

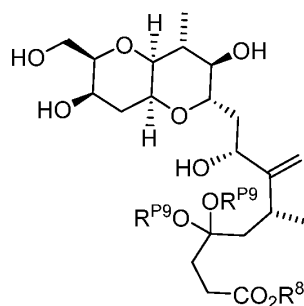
or a salt thereof, to yield a compound of Formula (E-R-11):



(E-R-11),

or a salt thereof;

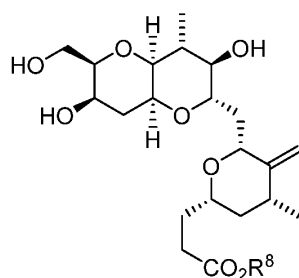
(b) a step of deprotecting a compound of Formula (E-R-11), or a salt thereof, under conditions sufficient to remove the groups R^{P5} and R^{P8} , to yield a compound of Formula (E-R-12):



(E-R-12),

or a salt thereof; and

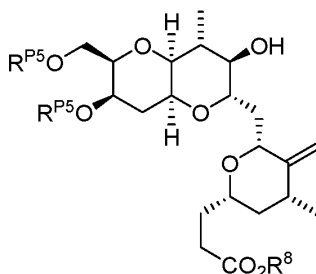
(c) a step of deprotecting and cyclizing the compound of Formula (E-R-12), or salt thereof, to yield a compound of Formula (E-R-13):



(E-R-13),

or a salt thereof;

(d) a step of protecting the compound of Formula (E-R-13), or a salt thereof, to yield a compound of Formula (E-R-14):



(E-R-14),

or a salt thereof, or a salt thereof; wherein:

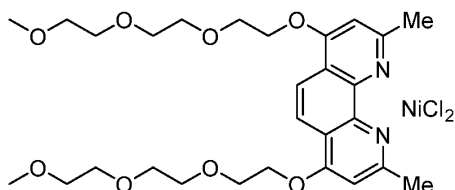
each instance of R^{P5} , R^{P8} , and R^{P9} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and optionally wherein two R^{P9} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00266] In certain embodiments, step (a) above (to form a compound of Formula (R-4-7A), (E-R-11), or a salt thereof) is a Ni/Cr-mediated reductive coupling reaction; and step (a-i) or (c) (to form a compound of Formula (R-4-7), (E-R-13), or a salt thereof) is a ketal deprotection and an acid-promoted intramolecular pyran cyclization. Reagents and conditions for steps (a), (a-i), and/or (c) above can be found in, *e.g.*, international PCT publications, WO

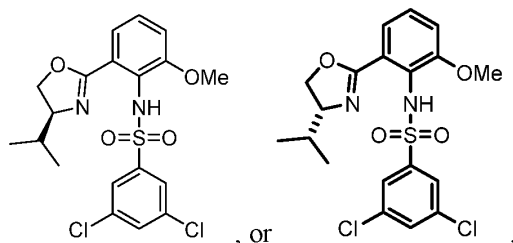
2016/176560, published November 3, 2016, and WO 2016/003975, published January 7, 2016; the entire contents of each of which is incorporated herein by reference.

[00267] The Ni/Cr-mediated reductive coupling (*i.e.*, steps (a)) is carried out in the presence of nickel and chromium. In certain embodiments, the nickel is a nickel complex. Examples of nickel complexes include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the nickel complex is $(\text{Et})_2\text{Phen}\bullet\text{NiCl}_2$. In certain embodiments, the nickel complex is the following:



In certain embodiments, the nickel complex is present in a catalytic amount.

[00268] In certain embodiments, the chromium is a chromium complex. In certain embodiments, the chromium complex is prepared from a chromium salt and a chiral ligand. In certain embodiments, the chromium salt is CrCl_2 or CrCl_3 . In certain embodiments, the chiral ligand is a chiral sulfonamide. Examples of chiral ligands include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the chiral ligand is (*S*)-**4-G**. In certain embodiments, the sulfonamide ligand is one of the following:



or a salt thereof. In certain embodiments, the chromium complex is present in a catalytic amount.

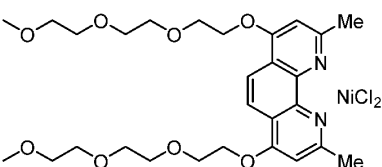
[00269] The Ni/Cr-mediated reductive coupling may be carried out in the presence of one or more additional reagents. In certain embodiments, the coupling is carried out in the presence of a lithium salt (*e.g.*, LiCl). In certain embodiments, the coupling is carried out in the presence of a reducing metal such as zinc or manganese (*e.g.*, zinc or manganese metal). In certain embodiments, the reducing metal is zinc metal. In certain embodiments, the reducing metal is manganese metal. In certain embodiments, the coupling is carried out in the presence of zirconium (*e.g.*, ZrCp_2Cl_2). In certain embodiments, the coupling is carried out in the presence of a base or proton scavenger (*e.g.*, 2,6-di-*tert*-butyl-4-methylpyridine). In certain

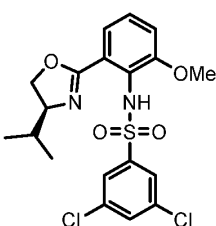
embodiments, the coupling is carried out in the presence of a proton sponge (e.g., 1,8-bis(dimethylamino)naphthalene).

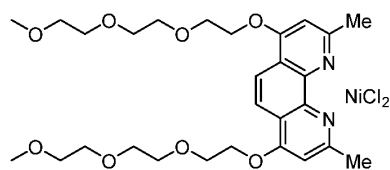
[00270] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is MeCN. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 40 °C.

[00271] In certain embodiments, the Ni/Cr-mediated reductive coupling is carried out in the presence of a nickel complex, a chromium salt, a sulfonamide ligand, a lithium salt, a zirconium complex, a reducing metal, and a base or proton scavenger. In certain embodiments, the step of coupling is carried out in the presence of (Et)₂Phen•NiCl₂, CrCl₂, (*S*)-**4-F**, LiCl, manganese metal, and ZrCp₂Cl₂. For example, in certain embodiments, the reaction is carried out under the following conditions: 2 mol% (Et)₂Phen•NiCl₂, 10 mol% CrCl₂, 10 mol% ligand (*S*)-**4-F**, 2 equivalents LiCl, excess manganese metal, 2.5 equivalents ZrCp₂Cl₂, in MeCN at room temperature (e.g., for 3 hours).

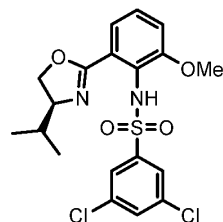
[00272] In certain embodiments, the coupling is carried out in the presence of: a nickel

complex of the formula: , CrCl₂, a sulfonamide ligand of

the formula: , Cp₂ZrCl₂, manganese metal, and a base or proton scavenger (e.g., 2,6-di-*tert*-butyl-4-methylpyridine and/or proton sponge (e.g., 1,8-Bis(dimethylamino)naphthalene)). In certain embodiments, the reaction is carried out in MeCN at around 40 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: 0.5 mol% or more of a nickel complex of the formula:



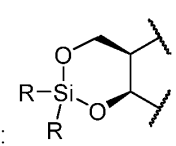
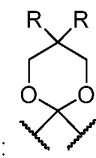
, 20 mol% CrCl₂, 20 mol% a sulfonamide ligand of the

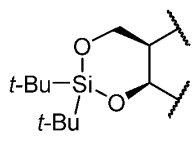
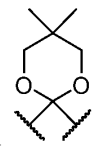


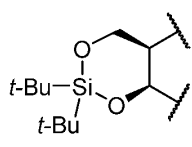
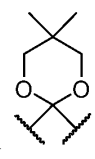
formula: , 1.1 equivalent Cp₂ZrCl₂, 4 equivalents manganese metal, and proton sponge in MeCN at around 40 °C (*e.g.*, for 19 hours).

[00273] In certain embodiments, R^{P5} and R^{P8} are silyl protecting groups; and the deprotection in step (b) is carried out in the presence of a fluoride source. In certain embodiments, the fluoride source is tetrabutylammonium fluoride (TBAF).

[00274] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

of the formula: ; two R^{P9} are joined together to form: ; R^{P8} is optionally substituted benzyl or optionally substituted silyl protecting group; and R⁸ is optionally substituted alkyl. In certain embodiments, two R^{P5} are joined with the intervening atoms to

form a ring of the formula: ; two R^{P9} are joined together to form ; R^{P8} is MPM; R⁸ is ethyl. In certain embodiments, two R^{P5} are joined with the intervening atoms to

form a ring of the formula: ; two R^{P9} are joined together to form ; R^{P8} is TBS; and R⁸ is methyl.

[00275] The ketal deprotection and acid-promoted intramolecular pyran cyclization in steps (a-i) and (c) (to form a compound of Formula (**R-4-7**), (**E-R-13**), or a salt thereof) involves deprotecting the ketal of the starting material, followed by a cyclization reaction to provide the new six-membered ring of the compound of Formula (**R-4-7**) or (**E-R-13**). The deprotecting and cyclizing may be done in the same step, or in separate steps, and in either order. In certain

embodiments, the step of deprotecting and cyclizing is carried out in the presence of an acid (*e.g.*, Lewis acid or Brønsted acid). In certain embodiments, the acid is a Lewis acid. In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of a hydride source.

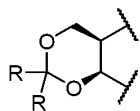
[00276] In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of a trialkylsilyl sulfonate or trialkylsilyl halide. In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of triethylsilyl trifluoromethylsulfonate (TESOTf). In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of trimethylsilyl trifluoromethylsulfonate (TMSOTf). In certain embodiments, the TESOTf or TMSOTf is present in a stoichiometric or excess amount.

[00277] In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of a trialkylsilane. In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of triethylsilane (Et_3SiH). In certain embodiments, the Et_3SiH is present in a stoichiometric or excess amount.

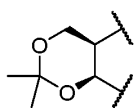
[00278] In certain embodiments, the reaction is carried out in a solvent (*e.g.*, CH_2Cl_2). In certain embodiments, the reaction is carried out at below room temperature. In certain embodiments, the reaction is carried out at approximately 0 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately 0 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature.

[00279] In certain embodiments, the step of deprotecting and cyclizing is carried out in the presence of a Lewis acid and a hydride source. In certain embodiments, the reaction is carried out in the presence of TESOTf and triethylsilane. In certain embodiments, the reaction is carried out in the presence of TESOTf and triethylsilane in DCM at around 0 °C. In certain embodiments, the reaction is carried out in the presence of TMSOTf and triethylsilane. In certain embodiments, the reaction is carried out in the presence of TMSOTf and triethylsilane in DCM at a temperature ranging from approximately -78 °C to approximately 0 °C. In certain embodiments, the reaction is carried out under the following conditions: 10 equivalents triethylsilane, 5 equivalents TESOTf, in DCM at around 0 °C (*e.g.*, for 3 hours). As another example, in certain embodiments, the reaction is carried out under the following conditions: 5 equivalents triethylsilane, 5 equivalents TMSOTf, in DCM at temperature ranging from approximately -78 °C to approximately 0 °C (*e.g.*, for 1 hour).

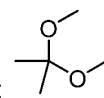
[00280] In certain embodiments, the step or re-protecting a compound of Formula (R-4-7C), (E-R-13), or a salt thereof (*i.e.*, step (d)), is carried out to install the R^{P5} groups. In certain embodiments, the resulting R^{P5} groups are joined together to form the following formula:



. In certain embodiments, the R^{P5} groups are of the following formula:



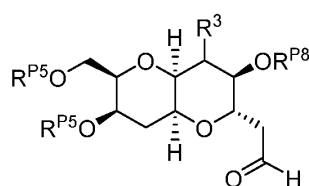
. In certain embodiments, the reaction is carried out in the presence of a ketal or



ketone; and an acid. In certain embodiments, the ketal is of the formula: (2,2-dimethoxypropane). In certain embodiments, the acid is pyridinium *p*-toluenesulfonate (PPTS). In certain embodiments, the reaction is carried out in the presence of 2,2-dimethoxypropane and PPTS. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, THF). In certain embodiments, the reaction is carried out in the presence of 2,2-dimethoxypropane and PPTS in THF at around 40 °C. In certain embodiments, the protection is carried out under the following conditions: 4 equivalents 2,2-dimethoxypropane and 5 mol% PPTS in THF at around 40 °C (*e.g.*, for 4-5 hours).

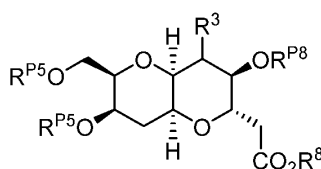
[00281] In certain embodiments, the compound of Formula (E-R-14), (E-R-8), (R-4-7), or (R-4-7B), or salt thereof, is purified by any combination of silica gel column chromatography, ODS (octadecylsilyl) column chromatography, and recrystallization.

[00282] As also shown in *Scheme 3B*, provided herein is a method of preparing a compound of Formula (R-4-5B):



(R-4-5B),

or a salt thereof, the method comprising reducing a compound of Formula (R-4-5A):



(R-4-5A),

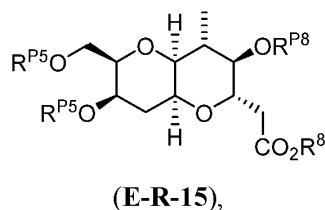
or a salt thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl;

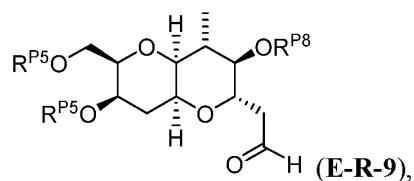
each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00283] In certain embodiments, the method comprises reducing a compound of Formula (**E-R-15**):



or a salt thereof, to yield a compound of Formula (**E-R-9**):



or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

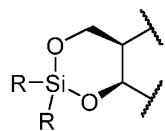
[00284] The step of reducing a compound of (**R-4-5A**), (**E-R-15**), or a salt thereof, converts the $-CO_2R^8$ moiety to an aldehyde. In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H^-) source. Any hydride source known in the art may be used in this transformation. Examples of hydride sources are provided herein. In certain

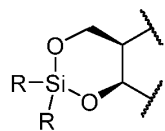
embodiments, the hydride source is diisobutylaluminum hydride (DIBAL). In certain embodiments, a stoichiometric or excess amount of DIBAL is used in the reaction.

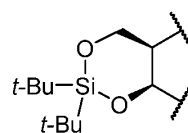
[00285] The step of reducing may optionally comprise reducing the $-\text{CO}_2\text{R}^8$ moiety to an alcohol, followed by oxidation of the resulting alcohol to an aldehyde to yield a compound of Formula (R-4-5B) or (E-R-9), or a salt thereof.

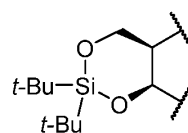
[00286] In certain embodiments, the step of reducing is carried out in the presence of DIBAL. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, DCM). In certain embodiments, the reaction is carried out at below room temperature. In certain embodiments, the reaction is carried out at around -78°C . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -70°C to approximately -78°C . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78°C to approximately 0°C . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78°C to approximately room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: DIBAL in DCM at -78°C (*e.g.*, for 1-2 hours). For example, in certain embodiments, the reaction is carried out under the following conditions: 2.3 equivalents DIBAL in DCM at -70°C to -78°C (*e.g.*, for 1-2 hours).

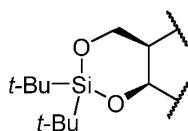
[00287] In certain embodiments, two $\text{R}^{\text{P}5}$ are joined with the intervening atoms to form a ring

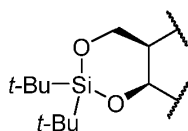


of the formula: ; $\text{R}^{\text{P}8}$ is optionally substituted benzyl or optionally substituted silyl protecting group; and R^8 is optionally substituted alkyl. In certain embodiments, two $\text{R}^{\text{P}5}$

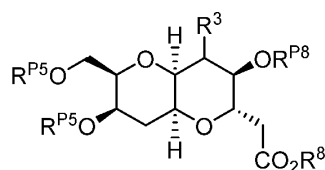


are joined with the intervening atoms to form a ring of the formula: ; $\text{R}^{\text{P}8}$ is MPM; and R^8 is methyl. In certain embodiments, two $\text{R}^{\text{P}5}$ are joined with the intervening



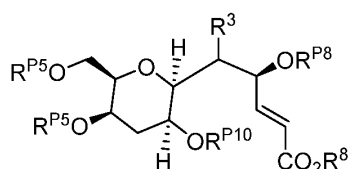
atoms to form a ring of the formula: ; $\text{R}^{\text{P}8}$ is TBS; and R^8 is methyl.

[00288] Also provided herein is a method of preparing a compound of Formula (R-4-5A):



(R-4-5A),

or a salt thereof, the method comprising cyclizing a compound of Formula (R-4-4):



(R-4-4),

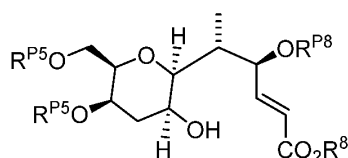
or a salt thereof, wherein:

R³ is hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P5}, R^{P8}, and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

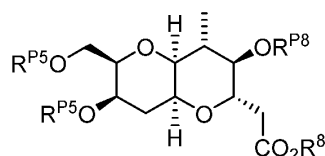
R⁸ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00289] In certain embodiments, the method comprises a step of cyclizing a compound of Formula (E-R-16):



(E-R-16),

or a salt thereof, to yield a compound of Formula (E-R-15):



(E-R-15),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

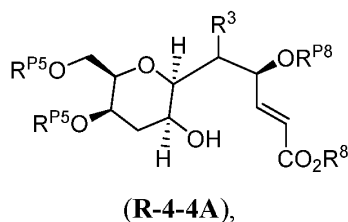
[00290] In certain embodiments, the step of cyclizing a compound of Formula (**R-4-4**) or (**E-R-16**), or a salt thereof, is carried out in the presence of a base. Any base may be used in this cyclization reaction. In certain embodiments, the base is a phosphate salt. In certain embodiments, the base is potassium phosphate (K_3PO_4). In certain embodiments, the base is present in 1 equivalent or less. In certain embodiments, the base is present in excess amount.

[00291] In certain embodiments, the step of cyclizing is carried out in the presence of one or more additional reagents, such as a metal chelator. In certain embodiments, the reaction is carried out in the presence of a crown ether (*e.g.*, 18-crown-6). In certain embodiments, the reaction is carried out in the presence of 18-crown-6. In certain embodiments, 1 equivalent or less of 18-crown-6 is used.

[00292] In certain embodiment, the reaction is carried out in the presence of a solvent. In certain embodiments, the solvent is toluene and/or MeOAc. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at from 0 °C to room temperature. In certain embodiments, the reaction is carried out at around room temperature.

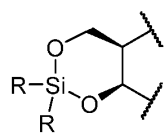
[00293] In certain embodiments, the step of cyclizing is carried out in the presence of a base and a crown ether. In certain embodiments, the reaction is carried out in the presence of K_3PO_4 and 18-crown-6. For example, in certain embodiments, the reaction is carried out under the following conditions: 1 equivalent K_3PO_4 , 3 equivalents 18-crown-6, in toluene at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 0.3 equivalents K_3PO_4 , 0.9 equivalents 18-crown-6, in toluene and MeOAc at around 3 °C (*e.g.*, for 1-2 hours).

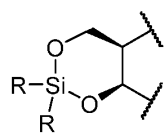
[00294] In certain embodiments, the compound of Formula (R-4-4) is a compound of Formula (R-4-4A):

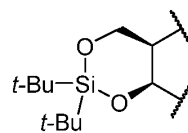


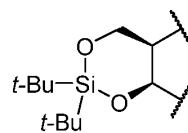
or a salt thereof.

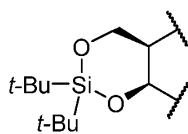
[00295] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

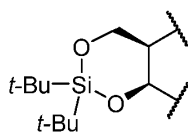


of the formula: ; R^{P8} is optionally substituted benzyl or optionally substituted silyl protecting group; and R⁸ is optionally substituted alkyl. In certain embodiments, two R^{P5}



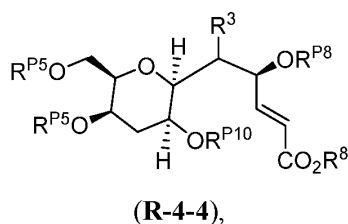
are joined with the intervening atoms to form a ring of the formula: ; R^{P8} is MPM; and R⁸ is methyl. In certain embodiments, two R^{P5} are joined with the intervening



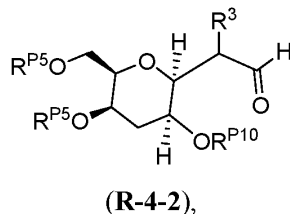
atoms to form a ring of the formula: ; R^{P8} is TBS; and R⁸ is methyl.

[00296] In certain embodiments, the compound of Formula (R-4-5A) or (E-R-15), or a salt thereof, is purified by silica gel column chromatography and/or recrystallization.

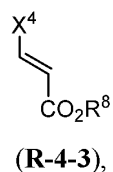
[00297] Also provided herein is a method of preparing a compound of Formula (R-4-4):



or a salt thereof, the method comprising coupling a compound of Formula (R-4-2):



or a salt thereof, with a compound of Formula (R-4-3):



or a salt thereof, wherein:

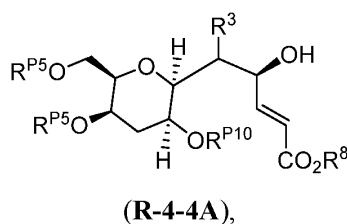
X^4 is halogen or a leaving group;

R^3 is hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

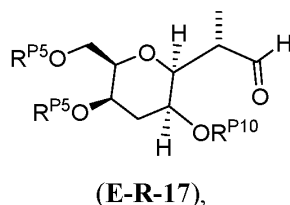
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00298] In certain embodiments, the coupling of a compound of Formula (R-4-2) and a compound of Formula (R-4-3) yields a compound of the Formula (R-4-4A):

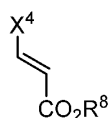


or a salt thereof, and the method of preparing a compound of (R-4-4), or a salt thereof, comprises protecting an oxygen atom of a compound of Formula (R-4-4A), or a salt thereof (*e.g.*, to introduce the group R^{P8}). The method may further comprise a step of deprotecting the compound to remove the protecting group R^{P10} .

[00299] In certain embodiments, the method comprises coupling a compound of Formula (E-R-17):

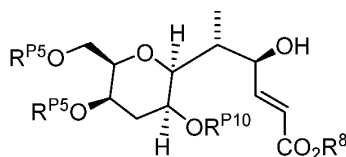


or a salt thereof, with a compound of Formula (R-4-3):



(R-4-3),

or a salt thereof, to yield a compound of Formula (E-R-18):



(E-R-18),

or a salt thereof, wherein:

X^4 is halogen or a leaving group;

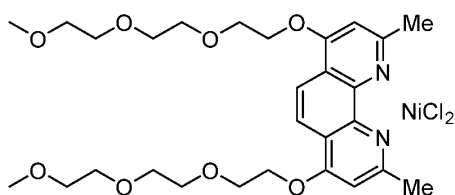
R^3 is hydrogen, halogen, or optionally substituted alkyl;

each instance of $\text{R}^{\text{P}5}$ and $\text{R}^{\text{P}10}$ is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two $\text{R}^{\text{P}5}$ groups are joined together with the intervening atoms to form an optionally substituted heterocycl ring; and

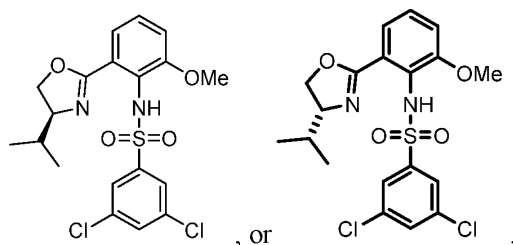
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocycl, optionally substituted aryl, optionally substituted heterocycl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00300] In certain embodiments, the method further comprising steps of (a) protecting the free hydroxyl group of the compound of Formula (E-R-18), or a salt thereof; and (b) deprotecting the resulting compound to remove the group $\text{R}^{\text{P}10}$.

[00301] In certain embodiments, the coupling of a compound of Formula (R-4-2) and a compound of Formula (R-4-3) to yield a compound of Formula (R-4-4) (or the coupling of a compound of Formula (E-R-17) and a compound of Formula (R-4-3) to yield a compound of Formula (E-R-18)) is a Ni/Cr-mediated coupling. The Ni/Cr-mediated reductive coupling is carried out in the presence of nickel and chromium. In certain embodiments, the nickel is a nickel complex. Examples of nickel complexes include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the nickel complex is $(\text{Me})_2\text{Phen}(\text{OMe})_2\bullet\text{NiCl}_2$. In certain embodiments, the nickel complex is present in a catalytic amount. In certain embodiments, the nickel complex is the following:



[00302] In certain embodiments, the chromium is a chromium complex. In certain embodiments, the chromium complex is prepared from a chromium salt and a chiral ligand. In certain embodiments, the chromium salt is CrCl_3 or CrCl_2 . In certain embodiments, the chiral ligand is a chiral sulfonamide. Examples of chiral ligands include, but are not limited to, those shown in *Figure 9B*. In certain embodiments, the chiral ligand is (*R*)-**4-E**. In certain embodiments, the chromium complex is present in a catalytic amount. In certain embodiments, the sulfonamide ligand is one of the following:



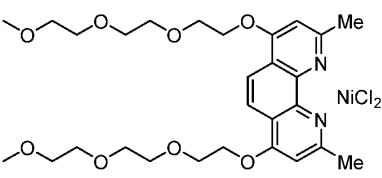
or a salt thereof.

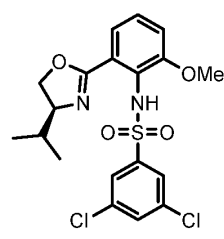
[00303] The Ni/Cr-mediated reductive coupling may be carried out in the presence of one or more additional reagents. In certain embodiments, the coupling is carried out in the presence of a lithium salt (*e.g.*, LiCl). In certain embodiments, the coupling is carried out in the presence of a reducing metal such as zinc or manganese (*e.g.*, zinc or manganese metal). In certain embodiments, the coupling is carried out in the presence of zirconium (*e.g.*, ZrCp_2Cl_2). In certain embodiments, the coupling is carried out in the presence of a base or proton scavenger (*e.g.*, 2,6-di-*tert*-butyl-4-methylpyridine or 2,6-lutidine). In certain embodiments, the coupling is carried out in the presence of proton sponge (*e.g.*, 1,8-bis(dimethylamino)naphthalene).

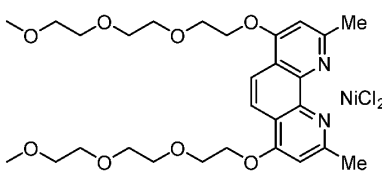
[00304] In certain embodiments, the reaction is carried out in a solvent (*e.g.*, MeCN). In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 100 °C. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out at around 30 °C.

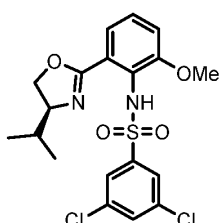
[00305] In certain embodiments, the Ni/Cr-mediated reductive coupling is carried out in the presence of a nickel complex, a chromium salt, a sulfonamide ligand, a lithium salt, a zirconium complex, a reducing metal, and a base or proton scavenger. In certain embodiments, the step of coupling is carried out in the presence of $(\text{Me})_2\text{Phen}(\text{OMe})_2\bullet\text{NiCl}_2$, CrCl_2 , ligand (*S*)-**4-E**, LiCl , manganese metal, 2,6-lutidine, and ZrCp_2Cl_2 . In certain embodiments, the reaction is carried out in a solvent (*e.g.*, MeCN). In certain embodiments, the reaction is carried out at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 2 mol% $(\text{Me})_2\text{Phen}(\text{OMe})_2\bullet\text{NiCl}_2$, 10 mol% CrCl_2 , 10 mol% ligand (*S*)-**4-E**, 2 equivalents LiCl , 1.1 equivalents Cp_2ZrCl_2 , 1 equivalent 2,6-lutidine, and excess manganese in MeCN at room temperature.

[00306] In certain embodiments, the coupling is carried out in the presence of: a nickel

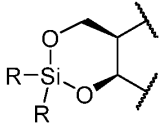
complex of the formula: , CrCl_2 , a sulfonamide ligand of

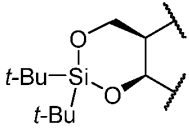
the formula: , Cp_2ZrCl_2 , manganese metal, and a base or proton scavenger (*e.g.*, 2,6-lutidine and/or proton sponge (*e.g.*, 1,8-bis(dimethylamino)naphthalene)). For example, in certain embodiments, the reaction is carried out under the following conditions:

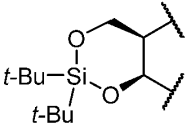
0.5 mol% of a nickel complex of the formula: , 20 ml%

CrCl_2 , 20 mol% of a sulfonamide ligand of the formula: , 1.1 equivalents Cp_2ZrCl_2 , 4 equivalents manganese metal, 2 equivalents 2,6-lutidine, and proton sponge in MeCN at around 30 °C (*e.g.*, for 2-3 hours).

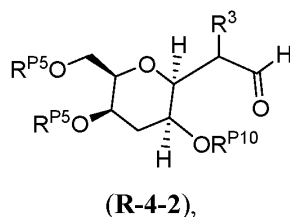
[00307] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

of the formula: ; R^{P8} is optionally substituted benzyl or optionally substituted silyl protecting group; R^8 is optionally substituted alkyl; and R^{P10} is a silyl protecting group. In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the

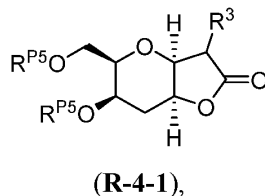
formula: ; R^{P8} is MPM; R^8 is methyl; and R^{P10} is TES. In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the formula:

; R^{P8} is TBS; R^8 is methyl; and R^{P10} is TES.

[00308] Provided herein a method of preparing a compound of Formula (R-4-2):



or a salt thereof, the method comprising reducing a compound of Formula (R-4-1):



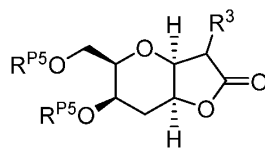
or a salt thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl; and

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocycl ring.

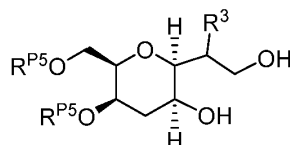
[00309] In certain embodiments, the method of preparing a compound of Formula (R-4-2), or a salt thereof, comprises the steps of:

(a) reducing a compound of Formula **(R-4-1)**:



(R-4-1),

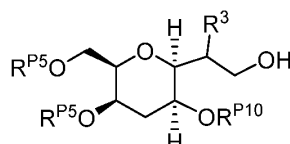
or a salt thereof, to yield a compound of Formula **(R-4-1A)**:



(R-4-1A),

or a salt thereof;

(b) protecting a compound of Formula **(R-4-1)**, or a salt thereof, to yield a compound of Formula **(R-4-1B)**:



(R-4-1B),

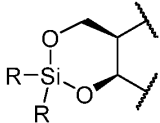
or a salt thereof; and

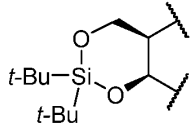
(c) oxidizing the compound of Formula **(R-4-1B)**, or a salt thereof, to yield a compound of Formula **(R-4-2)**, or a salt thereof.

[00310] The step of reducing a compound of **(R-4-2)**, or a salt thereof, reduces the lactone of the compound. In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H⁻) source. Any hydride source known in the art may be used in this transformation. Examples of hydride sources are provided herein. In certain embodiments, the hydride source is lithium borohydride (LiBH₄). In certain embodiments, the step of oxidizing (*i.e.*, step (c)) involves a Swern oxidation.

[00311] In certain embodiments, the step of reducing is carried out in the presence of LiBH₄. In certain embodiments, the reaction is carried out in a solvent such as diethyl ether. In certain embodiments, the reaction is carried out at approximately 0 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: LiBH₄ in diethyl ether at 0 °C.

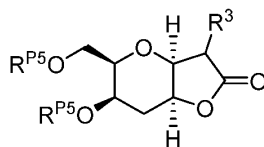
[00312] In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring

of the formula: ; and R^{P10} is a silyl protecting group. In certain embodiments,

two R^{P5} are joined with the intervening atoms to form a ring of the formula: ; and R^{P10} is TES.

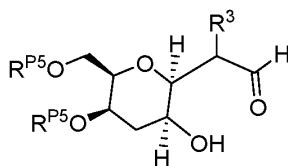
[00313] Also provided herein is an alternative method of preparing a compound of Formula (R-4-2), or a salt thereof, comprising:

(a) a step of reducing a compound of Formula (R-4-1):



(R-4-1),

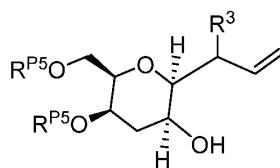
or a salt thereof, to yield a compound of Formula (R-4-2):



(R-4-2A),

or a salt thereof;

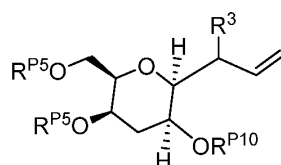
(b) a step of olefinating the compound of Formula (R-4-2A), or a salt thereof, to yield a compound of Formula (R-4-2B):



(R-4-2B),

or a salt thereof;

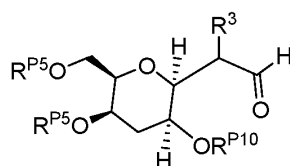
(c) a step of protecting the compound of Formula (R-4-2B), or salt thereof, to yield a compound of Formula (R-4-2C):



(R-4-2C),

or a salt thereof; and

(d) a step of oxidizing a compound of Formula (R-4-2C), or a salt thereof, to yield a compound of Formula (R-4-2):



(R-4-2),

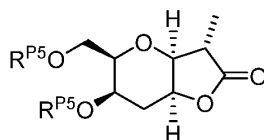
or a salt thereof, wherein:

R³ is hydrogen, halogen, or optionally substituted alkyl; and

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

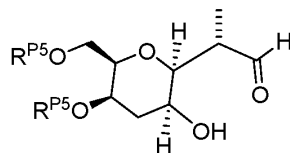
[00314] In certain embodiments, the method comprises:

(a) a step of reducing a compound of Formula (E-R-19):



(E-R-19),

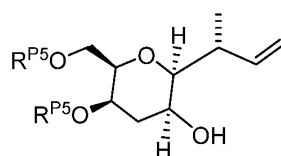
or a salt thereof, to yield a compound of Formula (E-R-20):



(E-R-20),

or a salt thereof;

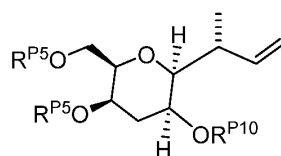
(b) a step of olefinating the compound of Formula (E-R-20), or a salt thereof, to yield a compound of Formula (E-R-21):



(E-R-21),

or a salt thereof,

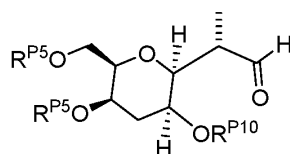
(c) a step of protecting the compound of Formula (E-R-21), or salt thereof, to yield a compound of Formula (E-R-22):



(E-R-22),

or a salt thereof; and

(d) a step of oxidizing a compound of Formula (E-R-22), or a salt thereof, to yield a compound of Formula (E-R-17):



(E-R-17),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00315] In certain embodiments, the step of reducing a compound of Formula (R-4-1), (E-R-19), or a salt thereof (*i.e.*, step (a)), is carried out in the presence of a hydride source.

Examples of hydride sources are provided herein. In certain embodiments, the hydride source is lithium borohydride ($LiBH_4$). In certain embodiments, the hydride source is diisobutylaluminum hydride (DIBAL). In certain embodiments, the reaction is carried out in a solvent (*e.g.*, toluene). In certain embodiments, the reaction is carried out in the presence of DIBAL in toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately room temperature to approximately $-78^\circ C$ to approximately $0^\circ C$. In certain embodiments, the reaction is carried out under the following conditions:

approximately 1.3 equivalents of DIBAL in toluene at from -78 to -60 °C (*e.g.*, for less than 1 hour).

[00316] In certain embodiments, the step of olefinating a compound of Formula (**R-4-2A**), (**E-R-20**), or a salt thereof (*i.e.*, step (b)), is carried out in the presence of an olefinating reagent and a base. In certain embodiments, the olefinating reagent is $\text{Ph}_3\text{PCH}_3\text{Br}$. In certain embodiments, the base is an alkoxide. In certain embodiments, the base is *t*-BuOK. In certain embodiments, the step of olefinating is carried out in the presence of $\text{Ph}_3\text{PCH}_3\text{Br}$ and *t*-BuOK. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, THF). In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately room temperature. In certain embodiments, the step of olefinating is carried out under the following conditions: 4 equivalents $\text{Ph}_3\text{PCH}_3\text{Br}$, 3 equivalents *t*-BuOK, in THF at from 0 to 10 °C (*e.g.*, for less than 1 hour).

[00317] In certain embodiments, R^{P10} is a silyl protecting group; and the step (c) of protecting is carried out in the presence of a silylating reagent and an amine base. In certain embodiments, R^{P10} is TES; and the silylating reagent is TESOTf. In certain embodiments, the amine base is triethylamine (TEA). In certain embodiments, the step of protecting is carried out in the presence of TESOTf and TEA. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately room temperature. In certain embodiments, the step of protecting is carried out in the presence of TESOTf and TEA in THF at from 0 to 10 °C (*e.g.*, for less than 1 hour).

[00318] In certain embodiments, the step of oxidizing a compound of Formula (**R-4-2C**), (**E-R-22**), or a salt thereof, is a *Johnson-Lemieux* oxidative cleavage. For example, in certain embodiments, the reaction is carried out in the presence of osmium tetroxide (OsO_4) or K_2OsO_4 ; and N-Methylmorpholine N-oxide (NMO). In certain embodiments, the reaction is carried out in the presence of sodium periodate (NaIO_4) or lead acetate $\text{Pb}(\text{OAc})_4$. In certain embodiments, the reaction is carried out in the presence of osmium tetroxide (OsO_4) and N-Methylmorpholine N-oxide (NMO), followed by sodium periodate (NaIO_4). In certain embodiments, the step of oxidizing is carried out in the presence of THF, acetone, and/or water. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. For example, in certain embodiments, the step of oxidizing is carried out under the following conditions: 25 equivalents OsO_4 and 3 equivalents NMO in THF/acetone/water at room temperature (*e.g.*, for 19 hours), followed by the addition of 3 equivalents NaIO_4 at room temperature (*e.g.*, for less than 1 hour).

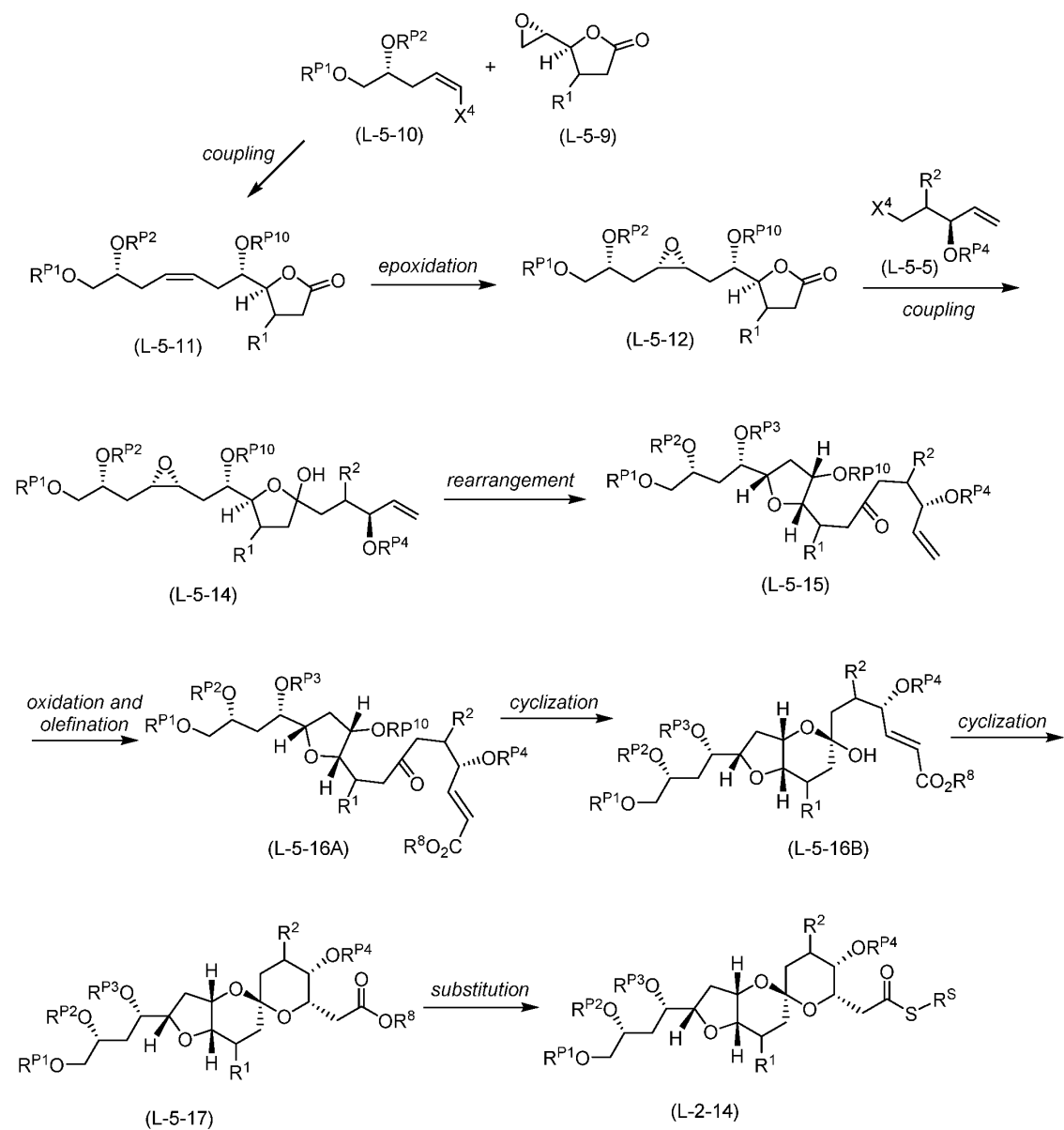
Preparation of Left Halves

[00319] As described herein, preparation of halichondrin natural products and analogs thereof may comprise a coupling of a “left half” fragment with a “right half” fragment. Methods useful in the preparation of right half building blocks are provided above. In another aspect, the present invention provides “left hand” building blocks, and methods useful in their preparation.

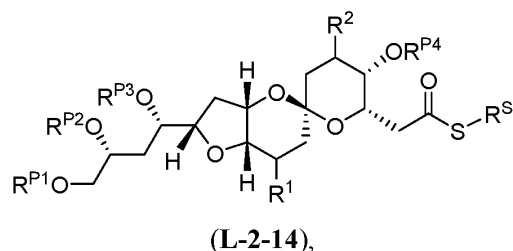
Preparation of Left Halves of Halichondrins

[00320] Provided herein are methods useful in the preparation of “left half” building blocks of halichondrins and analogs thereof. For example, left halves of compounds in the halichondrin series (*e.g.*, halichondrin A, B, C, and analogs thereof) can be prepared as shown in *Scheme 4A*. For example, a left half building block of Formula (L-2-14) can be prepared by thiolation of a compound of Formula (L-5-17), which can be prepared by cyclizing a compound of Formula (L-5-16B). To this end, a compound of Formula (L-5-16B) can be prepared by cyclization of a compound of Formula (L-5-16A), which can be prepared from an intermediate of Formula (L-5-15) via oxidation and olefination. As also shown in *Scheme 4A*, an intermediate of Formula (L-5-15) can be prepared by rearrangement of a compound of Formula (L-5-14). A compound of Formula (L-5-14) can be prepared by coupling a compound of Formula (L-5-12) with a compound of Formula (L-5-5). A compound of Formula (L-5-12) can be prepared by epoxidation of a compound of Formula (L-5-11), which may be prepared by coupling a compound of Formula (L-5-10) with a compound of Formula (L-5-9).

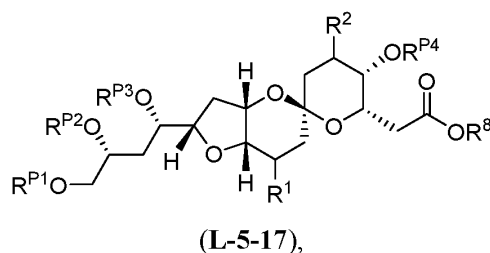
Scheme 4A



[00321] As shown in *Scheme 4A*, provided herein is a method of preparing a compound of Formula (L-2-14):



or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-17):



or a salt thereof, in the presence of a thiolating agent; wherein:

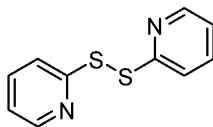
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P2} , R^{P3} , and R^{P4} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00322] As described herein, the step of forming a compound of Formula (L-2-14) comprises reacting a compound of Formula (L-5-17) in the presence of a thiolating agent. Any thiolating agent known in the art may be used to this end. In certain embodiments, the thiolating agent is a disulfide. In certain embodiments, the thiolating agent is of the formula $(R^S S)_2$. In certain



embodiments, the thiolating agent is: (Py-S)₂.

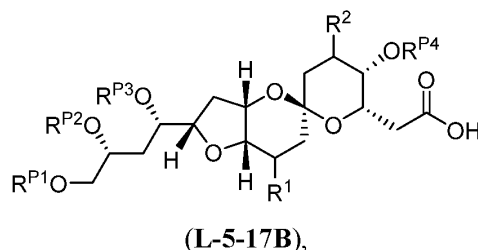
[00323] In certain embodiments, the step of thiolating a compound of Formula (L-5-17) is carried out in the presence of one of more additional reagents. In certain embodiments, the

step of thiolating is carried out in the presence of a phosphine reagent (*e.g.*, triphenylphosphine (Ph_3P)).

[00324] In certain embodiments, the step of thiolating is carried out in the presence of a disulfide and a phosphine. In certain embodiments, the reaction is carried out in the presence of $(\text{Py-S})_2$ and Ph_3P . In certain embodiments, the reaction is carried out in a solvent such as CH_2Cl_2 . In certain embodiments, the reaction is carried out at a temperature ranging from approximately $0\text{ }^\circ\text{C}$ to approximately $50\text{ }^\circ\text{C}$. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the step of thiolating is carried out under the following conditions: 1.4 equivalents of $(\text{Py-S})_2$, 1.2 equivalents of Ph_3P , in CH_2Cl_2 at room temperature (*e.g.*, for 10-20 hours).

[00325] In certain embodiments, the method of thiolating a compound of Formula (**L-5-17**), or a salt thereof, comprises the steps of:

- (a) deprotecting a compound of Formula (**L-5-17**), or a salt thereof, to yield a compound of Formula (**L-5-17B**):

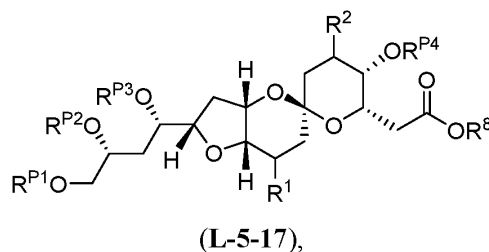


or a salt thereof; and

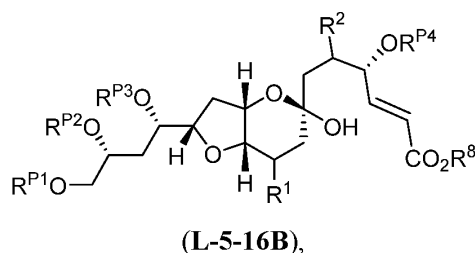
- (b) thiolating a compound of Formula (**L-5-17B**), or a salt thereof, to yield a compound of Formula (**L-2-14**), or a salt thereof.

[00326] In certain embodiments, R^{P1} , R^{P2} , R^{P3} , and R^{P4} are silyl protecting groups. In certain embodiments, R^{P1} and R^{P2} are TBS; and R^{P3} and R^{P4} are TES.

[00327] As also shown in *Scheme 4A*, provided herein is a method of preparing a compound of Formula (**L-5-17**):



or a salt thereof, the method comprising a step of cyclizing a compound of Formula (L-5-16B):



or a salt thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P2} , R^{P3} , and R^{P4} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00328] In certain embodiments, the step of cyclizing a compound of Formula (7-5-16B) is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amine or amide base. In certain embodiments, the base is an amidine or guanidine base. In certain embodiments, the base is an amidine base (*e.g.*, 1,8-diazabicyclo(5.4.0)undec-7-ene (DBU)). In certain embodiments, the step of cyclizing is carried out in the presence of an acid. In certain embodiments, the acid is a Lewis acid. In certain embodiments, the acid is a Brønsted acid.

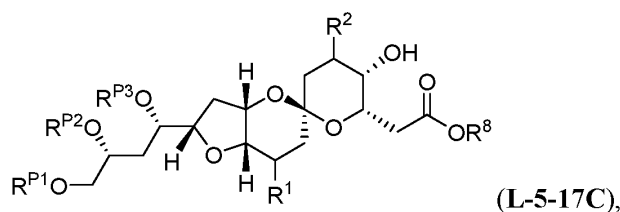
[00329] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt (*e.g.*, LiBr, LiCl). The step of cyclizing may be carried out in the presence of one or more additional reagents. In certain embodiments, the step of cyclizing is carried out in the presence of R^8 -OAc. In certain embodiments, the step of cyclizing is carried out in the presence of BnOAc.

[00330] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt, and a base. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr and DBU. In certain embodiments, the reaction is carried out in a solvent such as MeCN. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried

out under the following conditions: 10 equivalents LiBr, 5 equivalents DBU, and 10 equivalents BnOAc in MeCN at room temperature (*e.g.*, for 10-20 hours).

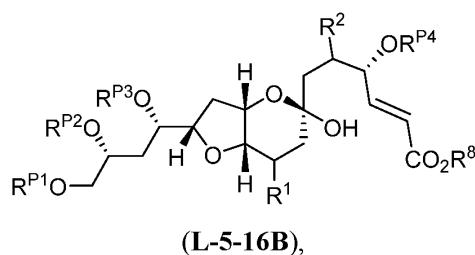
[00331] In certain embodiments, R^{P1} , R^{P2} , and R^{P3} are silyl protecting groups; and R^{P4} and R^{P8} are optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; R^{P3} is TES; R^{P4} is MPM; and R^8 is benzyl.

[00332] In certain embodiments, the compound of Formula (L-5-17), or a salt thereof, is deprotected to remove the group R^{P4} yield a compound of Formula (L-5-17C):



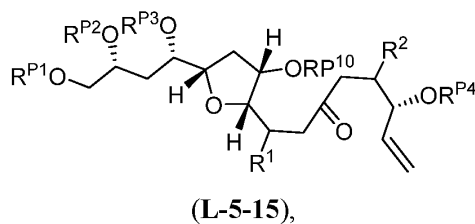
or a salt thereof; and optionally re-protecting (*i.e.*, to switch the group R^{P4} from, *e.g.*, a benzyl protecting group (*e.g.*, MPM) to a silyl protecting group (*e.g.*, trialkylsilyl such as triethylsilyl).

[00333] Provided herein is a method of preparing a compound of Formula (L-5-16B):

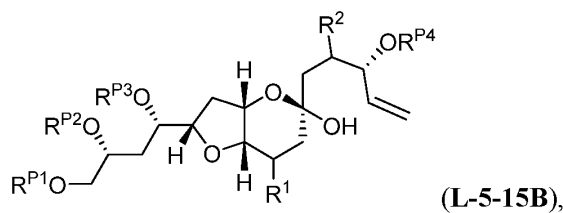


or a salt thereof, the method comprising the steps of:

(a) cyclizing a compound of Formula (L-5-15):



or a salt thereof, to give a compound of Formula (L-5-15B):

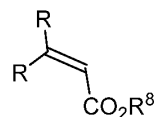


or a salt thereof; and

(b) reacting the compound of Formula (L-5-15B), or a salt thereof, in the presence of an olefin and an olefin metathesis catalyst to yield a compound of Formula (L-5-16B), wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

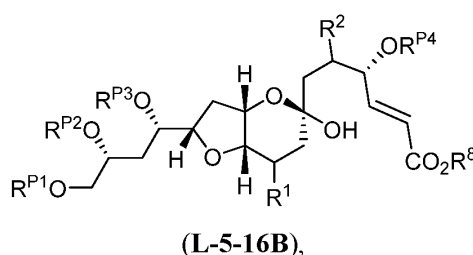
R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.



[00334] In certain embodiments, the olefin is of the formula: $R^2C=CH-CO_2R^8$. Furthermore, any olefin metathesis catalyst known in the art may be used in the metathesis reaction to furnish a compound of Formula (L-5-16B).

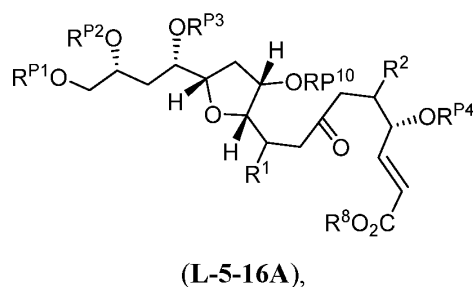
[00335] In certain embodiments, R^{P1} , R^{P2} , R^{P10} , and R^{P3} are silyl protecting groups; and R^{P4} is optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; and R^{P3} is TES; R^{P4} is MPM; and R^{P10} is TES.

[00336] Provided herein is a method of preparing a compound of Formula (L-5-16B):



or a salt thereof,

the method comprising a step of cyclizing a compound of Formula (L-5-16A):



or a salt thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

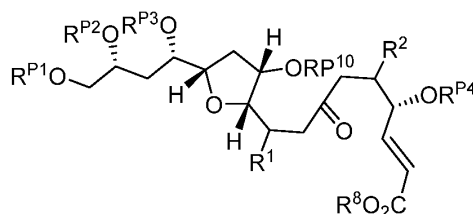
R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00337] In certain embodiments, the step of cyclizing a compound of Formula (L-5-16A), or a salt thereof, is carried out in the presence of a base. In certain embodiment, the step of cyclizing is carried out in the presence of an acid (*e.g.*, Lewis acid or Brønsted acid). In certain embodiments, the acid is a phosphoric acid. In certain embodiments, the acid is diphenylphosphate ((PhO)₂P(=O)OH). In certain embodiments, the acid is present in catalytic, stoichiometric, or excess amount relative to the compound of Formula (L-5-16A). In certain embodiments, the acid is present in catalytic amount (*e.g.*, approximately 5 mol%).

[00338] In certain embodiments, the step of cyclizing is carried out in the presence of diphenylphosphate. In certain embodiments, the step of cyclizing is carried out in a solvent such as THF, or a mixture of THF and H₂O. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 5 mol% diphenylphosphate in THF-H₂O at room temperature (*e.g.*, for approximately 24 hours).

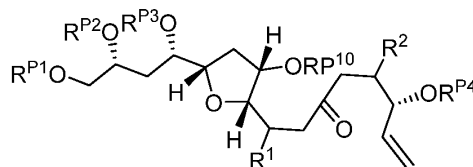
[00339] Also provided herein is a method of preparing a compound of Formula (L-5-16A):



(L-5-16A),

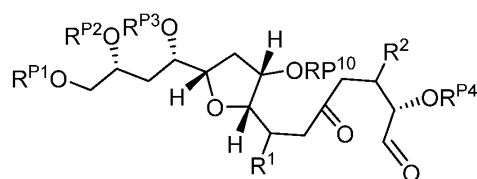
or a salt thereof, the method comprising the steps of:

(a) oxidizing a compound of Formula (L-5-15):

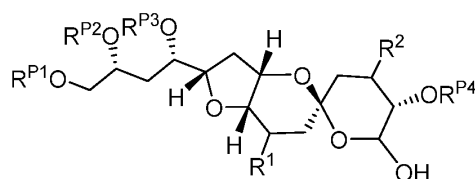


(L-5-15),

or a salt thereof, to yield a compound of Formula (L-5-15B) or (L-5-15BB):



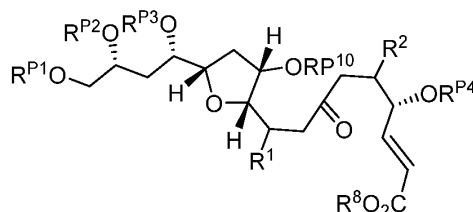
(L-5-15B),



(L-5-15BB),

or a salt thereof; and

(b) reacting the compound of Formula (L-5-15B) or (L-5-15BB), or a salt thereof, in the presence of a olefination reagent, to yield a compound of Formula (L-5-15C):



(L-5-15C),

or a salt thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00340] The reaction in step (a) above is an oxidative cleavage; the reaction in step (b) is an olefination reaction. In certain embodiments, the oxidative cleavage is carried out via ozonolysis (*e.g.*, in the presence of O_3). In certain embodiments, the cleavage is carried out in the presence of one or more reagents capable of dihydroxylating a double bond (*e.g.*, osmium tetroxide (OsO_4), N-methylmorpholine N-oxide (NMMO)), followed by a transition metal (*e.g.*, a lead complex such as $Pb(OAc)_4$). In certain embodiments, the double bond is dihydroxylated by treatment with OsO_4 , NMMO, and water. In certain embodiments, the reaction is carried out in the presence of a solvent such as acetone. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the double bond is dihydroxylated under the following conditions: 10 mol% OsO_4 , 2 equivalents NMMO, and water, in acetone at room temperature (*e.g.*, for

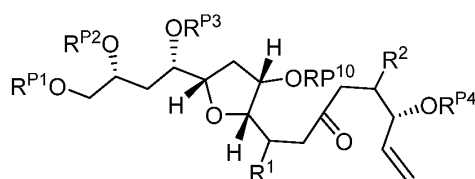
20-25 hours). The resulting compound is then treated, in certain embodiments, with $\text{Pb}(\text{OAc})_4$ and K_2CO_3 to yield the aldehyde or hemiacetal. For example, in certain embodiments, this step is carried out under the following conditions: 1.2 equivalents $\text{Pb}(\text{OAc})_4$, 3 equivalents K_2CO_3 , in CH_2Cl_2 at room temperature (*e.g.*, for approximately 1 hour).

[00341] In certain embodiments, the olefination is carried out in the presence of a Wittig or Horner-Wadsworth Emmons reagent. In certain embodiments, the olefination is carried out in the presence of a reagent of the formula: $(\text{RO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$. In certain embodiments, the reagent is of the formula: $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$ (*e.g.*, $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$). In certain embodiments, the olefination is carried out in the presence of a base (*e.g.*, a phosphate salt such as K_3PO_4).

[00342] In certain embodiments, the olefination is carried out in the presence of an olefination reagent of the formula: $(\text{RO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$, and a base. In certain embodiments, the olefination is carried out in the presence of $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$ and K_3PO_4 . In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 4 equivalents $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$, 3 equivalents K_3PO_4 , in toluene at room temperature (*e.g.*, for about 20-25 hours).

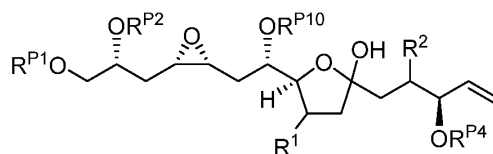
[00343] In certain embodiments, R^{P1} , R^{P2} , R^{P3} , and R^{P10} are silyl protecting groups; and R^{P4} and R^8 are optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; R^{P3} and R^{P10} are TES; R^{P4} is MPM; and R^8 is benzyl.

[00344] Provided herein is a method of preparing a compound of Formula (L-5-15):



(L-5-15),

or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-14):



(L-5-14),

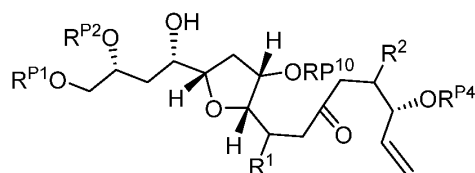
or a salt thereof, in the presence of an acid or a base, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00345] As described above, the method of forming a compound of Formula (L-5-15), or a salt thereof, involves reacting a step of reacting a compound of Formula (L-5-14), or a salt thereof, in the presence of an acid or a base. In certain embodiments, an acid is used. The acid may be a Lewis acid or a Brønsted acid. In certain embodiments, the acid is a Brønsted acid. In certain embodiments, the acid is a phosphoric acid (*e.g.*, phosphoric acid, diphenylphosphate). In certain embodiments, the acid is diphenylphosphate ((PhO)₂P(=O)OH). In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out from approximately 0° C to room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 5 mol% (PhO)₂P(=O)OH in toluene from 0° C to room temperature (*e.g.*, over 10-15 hours).

[00346] In certain embodiments, the compound of Formula (L-5-15) is of the Formula (L-5-15A):

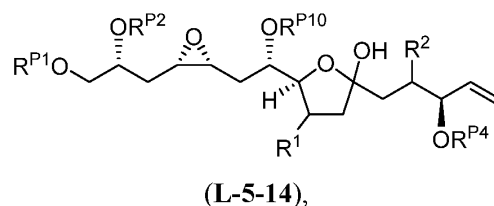


(L-5-15A),

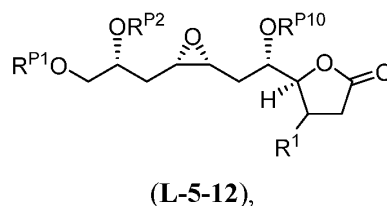
or a salt thereof, and the method further comprises a step of protecting the compound of Formula (L-5-15A), or a salt thereof, to yield a compound of Formula (L-5-15) (*e.g.*, to install the group R^{P3} , wherein the group R^{P3} is an oxygen protecting group).

[00347] In certain embodiments, R^{P1} , R^{P2} , and R^{P10} are silyl protecting groups; and R^{P4} is optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; R^{P10} is TES; and R^{P4} is MPM.

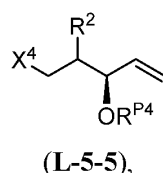
[00348] As shown in *Scheme 4A*, also provided herein is a method of preparing a compound of Formula (L-5-14):



or a salt thereof, the method comprising a step of coupling a compound of Formula (L-5-12):



or a salt thereof, with a compound of Formula (L-5-5):



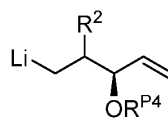
or a salt thereof, wherein:

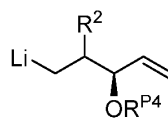
X⁴ is halogen or a leaving group;

R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1}, R^{P2}, R^{P4}, and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00349] In certain embodiments, the coupling of a compound of Formula (L-5-12) with a compound of Formula (L-5-5) is carried out in the presence of an organometallic reagent (*e.g.*, to convert X⁴ to a metal for addition to the compound of Formula (L-5-12)). In certain embodiments, the organometallic reagent is a lithium reagent (*e.g.*, to convert the compound

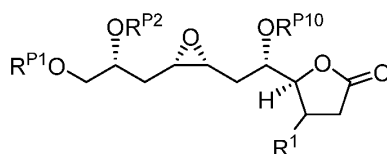


of the Formula (L-5-5) to a compound of the formula:  for addition to the compound of Formula (L-5-12)). In certain embodiments, lithium reagent is an organolithium (*e.g.*, *n*-butyllithium, *tert*-butyllithium, *sec*-butyllithium). In certain embodiments, the lithium reagent is LiHMDS or LDA. In certain embodiments, the reaction is carried out in the presence of *tert*-butyllithium. In certain embodiments, the reaction is performed in a solvent such as THF. In certain embodiments, the reaction is carried out at a temperature ranging

from approximately -78 °C to approximately 0 °C. In certain embodiments, the reaction is carried out at -78° C to room temperature. For example, in certain embodiments, the reaction is carried out with 2.6 equivalents of *tert*-butyllithium in THF from -78° C to room temperature (*e.g.*, over less than 1 hour).

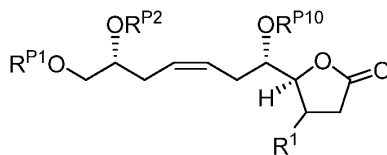
[00350] In certain embodiments, R^{P1}, R^{P2}, and R^{P10} are silyl protecting groups; and R^{P4} is optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; R^{P10} is TES; and R^{P4} is MPM.

[00351] Also provided herein is a method of preparing a compound of Formula (L-5-12):



(L-5-12),

or a salt thereof, the method comprising a step of epoxidizing a compound of Formula (L-5-11):



(L-5-11),

or a salt thereof, wherein:

R¹ is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1}, R^{P2}, and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

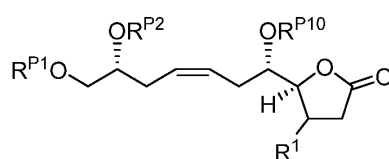
[00352] Any epoxidation reagent may be used in the step of epoxidizing described above. In certain embodiments, the epoxidation reagent is a peracid (*e.g.*, *m*-CPBA). In certain embodiments, the epoxidation reagent is an organometallic reagent. In certain embodiments, the epoxidation reagent is a titanium reagent (*e.g.*, Ti(Oi-Pr)₄). In certain embodiments, the epoxidation reagent is a vanadium reagent (*e.g.*, VO(TMHD)₂). In certain embodiments, the epoxidation is a *Sharpless epoxidation*. In certain embodiments, the step of epoxidizing is carried out in the presence of one or more additional reagents. In certain embodiments, epoxidation is carried out in the presence of a peroxide (*e.g.*, *t*-BuOOH).

[00353] In certain embodiments, the step of epoxidizing is carried out in the presence of a vanadium reagent and a peroxide. In certain embodiments, the reaction is carried out in the

presence of $\text{VO}(\text{TMHD})_2$ and *t*-BuOOH. In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 5 mol% $\text{VO}(\text{TMHD})_2$ and 2 equivalents *t*-BuOOH in toluene at room temperature (*e.g.*, for 1-10 hours).

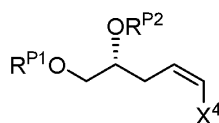
[00354] In certain embodiments, R^{P1} , R^{P2} , and R^{P10} are silyl protecting groups; and R^{P4} is optionally substituted benzyl. In certain embodiments, R^{P1} and R^{P2} are TBS; and R^{P10} is TES.

[00355] Also provided herein is a method of preparing a compound of Formula (L-5-11):



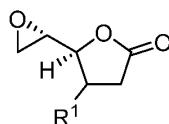
(L-5-11),

or a salt thereof, the method comprising a step of coupling a compound of Formula (L-5-10):



(L-5-10),

or a salt thereof, with a compound of Formula (L-5-9):



(L-5-9),

or a salt thereof, wherein:

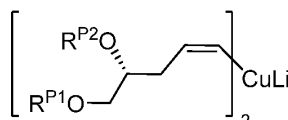
X^4 is halogen or a leaving group;

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P2} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00356] In certain embodiments, the coupling of a compound of Formula (L-5-10) with a compound of Formula (L-5-9) is carried out in the presence of a metal or organometallic reagent (*e.g.*, to convert X^4 to a metal for addition to the compound of Formula (L-5-9)). In certain embodiments, the reaction is carried out in the presence of copper. In certain embodiments, the copper is a copper complex or copper salt. In a particular embodiment, the

copper source is Li(thienylCuCN). In certain embodiments, the reaction is carried out in the presence of a lithium reagent. In certain embodiments, lithium reagent is an organolithium (*e.g.*, *n*-butyllithium, *tert*-butyllithium, *sec*-butyllithium). In certain embodiments, the lithium reagent is LiHMDS or LDA. In certain embodiments, the reactions is carried out in the presence of a lithium reagent and a copper reagent (*e.g.*, to convert the compound of the



Formula (L-5-10) to a compound of the formula: for addition to the compound of Formula (L-5-9)). The reaction may also be carried out in the presence of a Lewis acid (*e.g.*, $\text{BF}_3 \bullet \text{Et}_2\text{O}$).

[00357] In certain embodiments, the step of coupling is carried out in the presence of a copper source, an organometallic, and a Lewis acid. In certain embodiments, the reaction is carried out in the presence of Li(thienylCuCN), *n*-butyllithium, and $\text{BF}_3 \bullet \text{Et}_2\text{O}$. In certain embodiments, the reaction is carried out in a solvent such as Et_2O . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78°C to approximately 0°C . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78°C to approximately room temperature. In certain embodiments, the reaction is carried out at around -78°C . For example, in certain embodiments, the reaction is carried out under the following conditions: 2 equivalents Li(thienylCuCN), 1.75 equivalents *n*-butyllithium, and 1.6 equivalents $\text{BF}_3 \bullet \text{Et}_2\text{O}$, in Et_2O at -78°C (*e.g.*, for 1 hour).

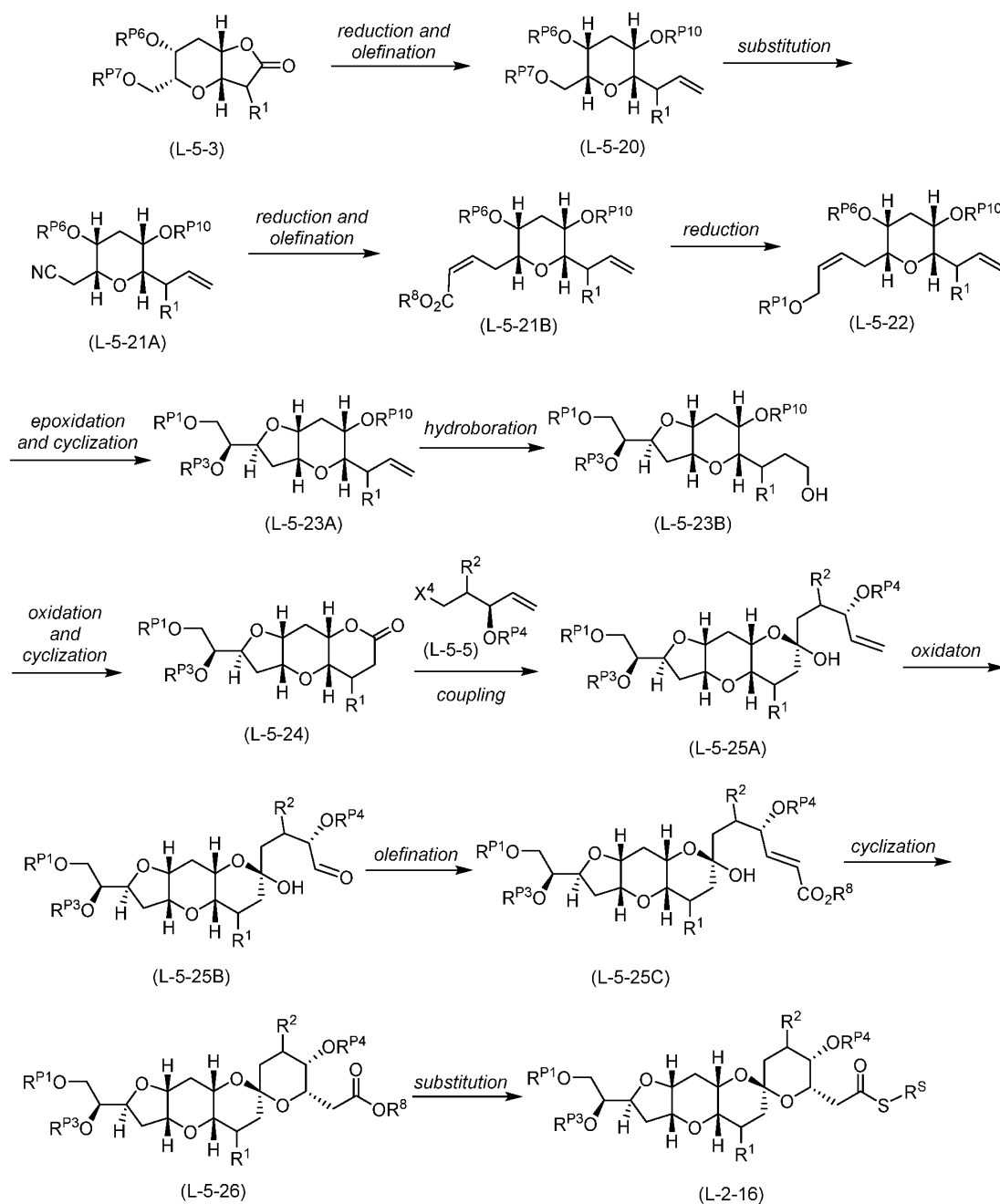
[00358] In certain embodiments, R^{P1} and R^{P2} are silyl protecting groups; In certain embodiments, R^{P1} and R^{P2} are TBS.

Preparation of Left Halves of Homohalichondrins

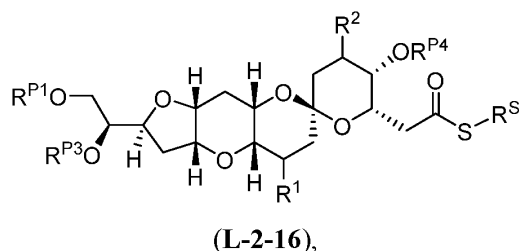
[00359] Also provided herein are “left hand” building blocks of homohalichondrins (*e.g.*, homohalichondrin A, B, C), and analogs thereof, such as compounds of Formula (L-2-16). Methods useful in the preparation of left hand building blocks of homohalichondrins (*e.g.*, compounds of Formula (L-2-16)) are outlined in *Scheme 4B*. For instance, a compound of Formula (L-2-16) can be prepared by thiolating a compound of Formula (L-5-26), which can be prepared via cyclization of a compound of Formula (L-5-25C). To this end, a compound of Formula (L-5-25C) can be prepared by oxidation and olefination of a compound of Formula (L-5-25A). As also shown in *Scheme 4B*, coupling of a compound of Formula (L-5-

24) with a compound of Formula (**L-5-5**) can provide a compound of Formula (**L-5-25A**). Furthermore, a compound of Formula (**L-5-24**) can be prepared by hydroboration, oxidation, and cyclization of a compound of Formula (**L-5-23A**), which can be prepared by epoxidizing the internal olefin of a compound of Formula (**L-5-22**), followed by cyclization. A compound of Formula (**L-5-22**) can be prepared by reducing a compound of Formula (**L-5-21B**), which may be prepared by reduction and olefination of a nitrile of Formula (**L-5-21A**). The nitrile can be prepared by reduction and olefination of a compound of Formula (**L-5-3**), followed by substitution of a compound of Formula (**L-5-20**) (*i.e.*, to convert the group $-\text{OR}^{\text{P7}}$ to $-\text{CN}$).

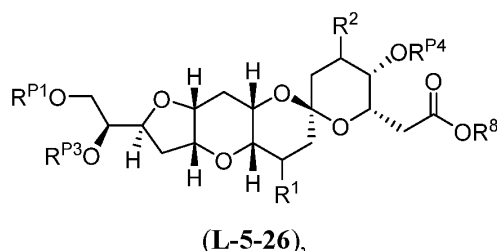
Scheme 4B



[00360] As shown in *Scheme 4B*, provided herein is a method of preparing a compound of Formula (L-2-16):



or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-26):



or a salt thereof, in the presence of a thiolating agent; wherein:

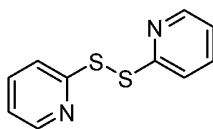
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00361] As described herein, the step of forming a compound of Formula (L-2-16) comprises reacting a compound of Formula (L-5-26) in the presence of a thiolating agent. Any thiolating agent known in the art may be used to this end. In certain embodiments, the thiolating agent is a disulfide. In certain embodiments, the thiolating agent is of the formula $(R^S S)_2$. In certain



embodiments, the thiolating agent is:

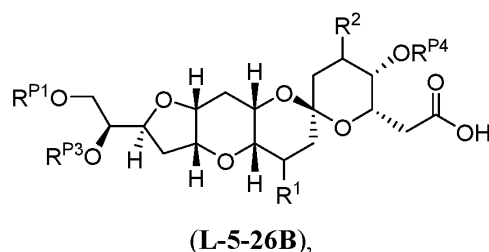
[00362] In certain embodiments, the step of thiolating a compound of Formula (L-5-26) is carried out in the presence of one or more additional reagents. In certain embodiments, the

step of thiolating is carried out in the presence of a phosphine reagent (*e.g.*, triphenylphosphine (Ph_3P)).

[00363] In certain embodiments, the step of thiolating is carried out in the presence of a disulfide and a phosphine. In certain embodiments, the reaction is carried out in the presence of $(\text{Py-S})_2$ and Ph_3P . In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the step of thiolating is carried out under the following conditions: 1.2 equivalents of $(\text{Py-S})_2$, 3 equivalents of Ph_3P , in toluene at room temperature (*e.g.*, for 10-20 hours).

[00364] In certain embodiments, the method of thiolating a compound of Formula (**L-5-26**), or a salt thereof, comprises the steps of:

- (a) deprotecting a compound of Formula (**L-5-26**), or a salt thereof, to yield a compound of Formula (**L-5-26B**):

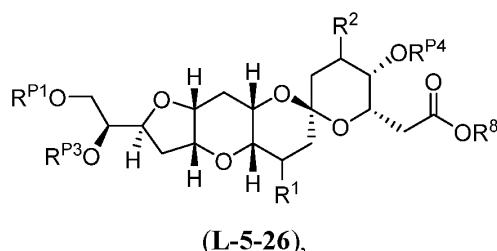


or a salt thereof; and

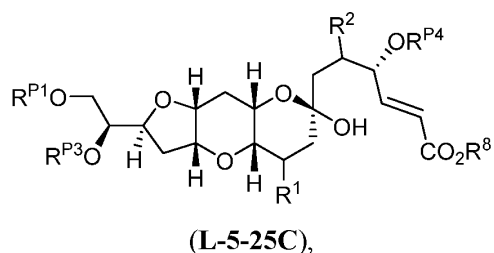
- (b) thiolating a compound of Formula (**L-5-26B**), or a salt thereof, to yield a compound of Formula (**L-2-6**), or a salt thereof.

[00365] In certain embodiments, R^{P1} , R^{P3} , and R^{P4} are silyl protecting groups. In certain embodiments, R^{P1} is TBS; and R^{P3} and R^{P4} are TES.

[00366] As also shown in *Scheme 4B*, provided herein is a method of preparing a compound of Formula (**L-5-26**):



or a salt thereof, the method comprising a step of cyclizing a compound of Formula (L-5-25C):



or a salt thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

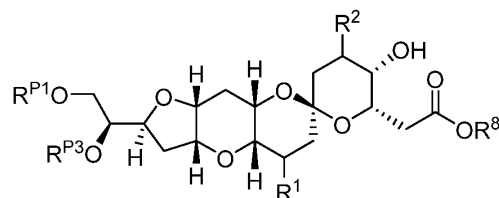
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00367] In certain embodiments, the step of cyclizing a compound of Formula (7-5-25C) is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amidine, guanidine base. In certain embodiments, the base is an amine or amide base. In certain embodiments, the base is an amidine base (*e.g.*, 1,8-diazabicyclo(5.4.0)undec-7-ene (DBU)). In certain embodiments, the step of cyclizing is carried out in the presence of an acid. In certain embodiments, the acid is a Lewis acid. In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt (*e.g.*, LiBr, LiCl). The step of cyclizing may be carried out in the presence of one or more additional reagents. In certain embodiments, the step of cyclizing is carried out in the presence of R^8 -OAc. In certain embodiments, the step of cyclizing is carried out in the presence of BnOAc.

[00368] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt, and a base. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr and DBU. In certain embodiments, the reaction is carried out in a solvent such as MeCN. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 10 equivalents LiBr and 20 equivalents DBU in MeCN at room temperature (*e.g.*, for 10-20 hours).

[00369] In certain embodiments, R^{P1} and R^{P3} are silyl protecting groups; R^{P4} is optionally substituted benzyl; and R^8 is optionally substituted benzyl. In certain embodiments, R^{P1} is TBS; R^{P3} is TES; R^{P4} is MPM; and R^8 is benzyl.

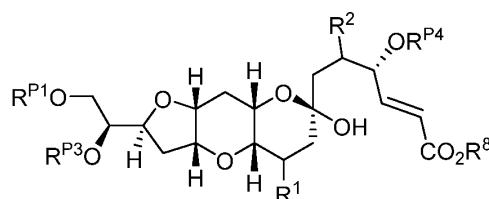
[00370] In certain embodiments, the compound of Formula (L-5-26), or a salt thereof, is deprotected to remove the group R^{P4} yield a compound of Formula (L-5-26B):



(L-5-26B),

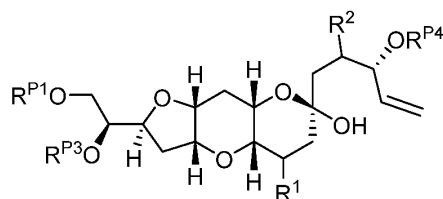
or a salt thereof; and optionally re-protected (*i.e.*, to switch the group R^{P4} from, *e.g.*, a benzyl protecting group (*e.g.*, MPM) to a silyl protecting group (*e.g.*, trialkylsilyl such as triethylsilyl).

[00371] Also provided herein is a method of preparing a compound of Formula (L-5-25C):



(L-5-25C),

or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-25A):



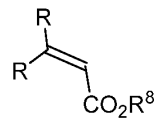
(L-5-25A),

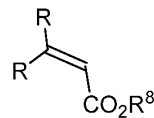
or a salt thereof, in the presence of an olefin and an olefin metathesis catalyst; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

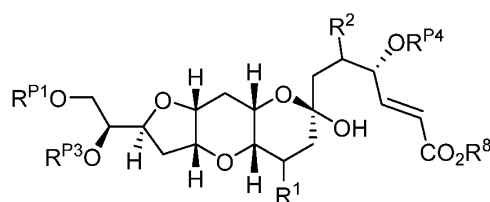
R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.



[00372] In certain embodiments, the olefin is of the formula: . Further, any olefin metathesis known in the art may be used in the metathesis reaction to furnish a compound of Formula (L-5-25C).

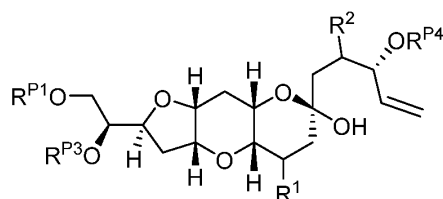
[00373] Also provided herein is an alternative method of preparing a compound of Formula (L-5-25C):



(L-5-25C),

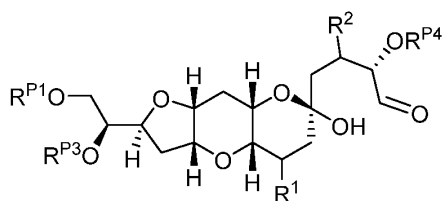
or a salt thereof, the method comprising the steps of:

(a) oxidizing a compound of Formula (L-5-25A):

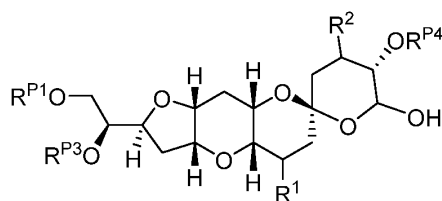


(L-5-25A),

or a salt thereof, to yield a compound of Formula (L-5-25B) or (L-5-25BB):



(L-5-25B),



(L-5-25BB),

or a salt thereof; and

(c) reacting the compound of Formula (L-5-25B) or (L-5-25BB), or a salt thereof, in the presence of an olefination reagent, to yield a compound of Formula (L-5-25C), or a salt thereof.

[00374] The reaction in step (a) above is an oxidative cleavage; the reaction in step (b) is an olefination reaction. In certain embodiments, the oxidative cleavage is carried out via ozonolysis (*e.g.*, in the presence of O₃). In certain embodiments, the cleavage is carried out in the presence of reagents capable of dihydroxylating a double bond (*e.g.*, osmium tetroxide (OsO₄), N-methylmorpholine N-oxide (NMMO)), followed by a transition metal (*e.g.*, a lead complex such as Pb(OAc)₄).

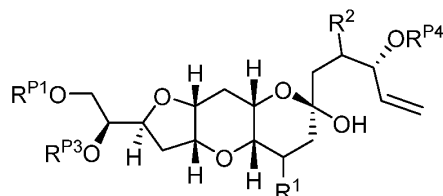
[00375] In certain embodiments, the double bond is dihydroxylated by treatment with OsO₄, NMMO, and water. In certain embodiments, the reaction is carried out in the presence of a solvent such as acetone. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the double bond is dihydroxylated under the following conditions: 10 mol% OsO₄, 2 equivalents NMMO, and water, in acetone at room temperature (*e.g.*, for 1-5 hours). The resulting compound is then treated, in certain embodiments, with Pb(OAc)₄ and K₂CO₃ to yield the aldehyde or hemiacetal. For example, in certain embodiments, this step is carried out under the following conditions: 1.5 equivalents Pb(OAc)₄, 10 equivalents K₂CO₃, in CH₂Cl₂ at room temperature (*e.g.*, for under 1 hour).

[00376] In certain embodiments, the olefination is carried out in the presence of a Wittig or Horner-Wadsworth Emmons reagent. In certain embodiments, the olefination is carried out in the presence of a reagent of the formula: (RO)₂P(O)CH₂CO₂R⁸. In certain embodiments, the reagent is of the formula: (MeO)₂P(O)CH₂CO₂R⁸ (*e.g.*, (MeO)₂P(O)CH₂CO₂Bn). In certain embodiments, the olefination is carried out in the presence of a base (*e.g.*, a phosphate salt such as K₃PO₄, or a hydride such as NaH).

[00377] In certain embodiments, the olefination is carried out in the presence of an olefination reagent of the formula: (RO)₂P(O)CH₂CO₂R⁸, and a base. In certain embodiments, the olefination is carried out in the presence of (MeO)₂P(O)CH₂CO₂Bn and NaH. In certain embodiments, the reaction is carried out in a solvent such as THF. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at 0 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: 5 equivalents (MeO)₂P(O)CH₂CO₂Bn, 4 equivalents NaH, in THF at 0 °C (*e.g.*, for about 1-5 hours).

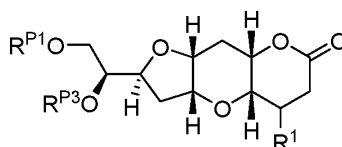
[00378] In certain embodiments, R^{P1} and R^{P3} are silyl protecting groups; R^{P4} is optionally substituted benzyl; and R^8 is optionally substituted benzyl. In certain embodiments, R^{P1} is TBS; R^{P3} is TES; R^{P4} is MPM; and R^8 is benzyl.

[00379] Also provided herein is a method of preparing a compound of Formula (L-5-25A):



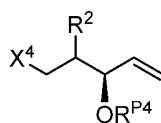
(L-5-25A),

or a salt thereof, the method comprising a step of coupling a compound of Formula (L-5-24):



(L-5-24),

or a salt thereof, with a compound of Formula (L-5-5):



(L-5-5),

or a salt thereof, wherein:

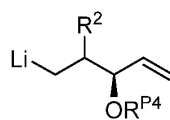
X^4 is halogen or a leaving group;

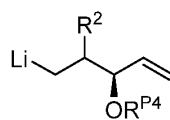
R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00380] In certain embodiments, the coupling of a compound of Formula (L-5-24) with a compound of Formula (L-5-5) is carried out in the presence of an organometallic reagent (*e.g.*, to convert X^4 to a metal for addition to the compound of Formula (L-5-24)). In certain embodiments, the organometallic reagent is a lithium reagent (*e.g.*, to convert the compound

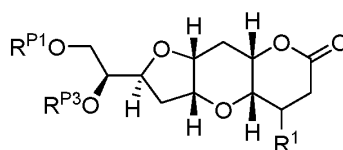


of the Formula (L-5-5) to a compound of the formula:  for addition to the compound of Formula (L-5-24)). In certain embodiments, lithium reagent is an organolithium (e.g., *n*-butyllithium, *tert*-butyllithium, *sec*-butyllithium). In certain embodiments, the lithium reagent is LiHMDS or LDA.

[00381] In certain embodiments, the reaction is carried out in the presence of *tert*-butyllithium. In certain embodiments, the reaction is performed in a solvent such as THF. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at approximately -78° C. For example, in certain embodiments, the reaction is carried out with 2.5 equivalents of *tert*-butyllithium in THF at -78° C (e.g., over less than 1 hour).

[00382] In certain embodiments, R^{P1} and R^{P3} are silyl protecting groups; and R^{P4} is optionally substituted benzyl. In certain embodiments, R^{P1} is TBS; R^{P3} is TES; and R^{P4} is MPM.

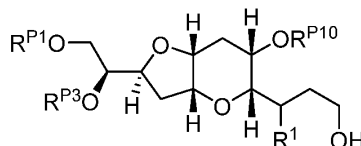
[00383] Provided herein is a method of preparing a compound of Formula (L-5-24):



(L-5-24),

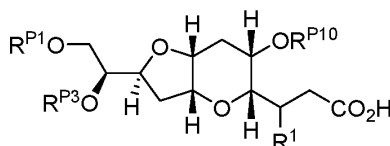
or a salt thereof, the method comprising the steps of:

(a) oxidizing a compound of Formula (L-5-23B):



(L-5-23B),

or a salt thereof, to yield a compound of Formula (L-5-23C):



(L-5-23C),

or a salt thereof; and

(b) cyclizing a compound of Formula (L-5-23C), or a salt thereof, to yield a compound of Formula (L-5-24), or a salt thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

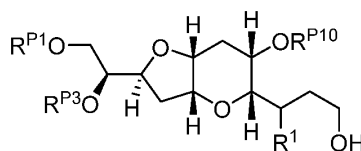
R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00384] The step of oxidizing a compound of Formula (L-5-23B) is carried out in the presence of an oxidant. In certain embodiments, the oxidant is a hypervalent iodine reagent. In certain embodiments, the oxidant is a periodinane (*e.g.*, Dess-Martin periodinane). In certain embodiments, the oxidant is (Diacetoxyiodo)benzene ($\text{PhI}(\text{OAc})_2$). In certain embodiments, the oxidation is carried out in the presence of one or more addition reagents. In certain embodiments, the oxidation is carried out in the presence of (2,2,6,6-tetramethylpiperidin-1-yl)oxyl (TEMPO). In certain embodiments, the oxidation is carried out in the presence of TEMPO and hypervalent iodine. In certain embodiments, the oxidation in step (a) and the cyclization in step (b) are carried out in the same step, or in subsequent steps. In certain embodiments, the cyclization in step (b) is carried out in a separate step, and in the presence of an acid (*e.g.*, Lewis acid or Brønsted acid) or a base.

[00385] In certain embodiments, the step of oxidizing is carried out in the presence of $\text{PhI}(\text{OAc})_2$ and TEMPO. In certain embodiments, the step of oxidizing is carried out in a solvent such as CH_2Cl_2 . In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the step of oxidizing is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 20 mol% TEMPO, 3 equivalents $\text{PhI}(\text{OAc})_2$, in CH_2Cl_2 at room temperature (*e.g.*, over 24-48 hours).

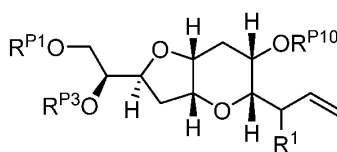
[00386] In certain embodiments, R^{P1} and R^{P3} are silyl protecting groups; and R^{P10} is hydrogen. In certain embodiments, R^{P1} is TBS; R^{P3} is TES; and R^{P10} is hydrogen.

[00387] Provided herein is a method of preparing a compound of Formula (L-5-23B):



(L-5-23B),

or a salt thereof, the method comprising a step of hydrating a compound of Formula (L-5-23A):



(L-5-23A),

or a salt thereof, wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

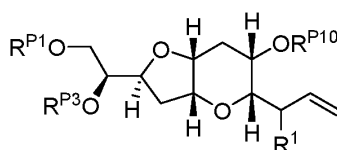
R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00388] In certain embodiments, the step of hydrating the compound of Formula (L-5-23A) is a hydroboration reaction. Any reagents or conditions to effect hydroboration may be used. For example, the reaction can be carried out in the presence of a borane (*e.g.*, BH_3 or 9-BBN), followed by a peroxide (*e.g.*, H_2O_2) or a perborate (*e.g.*, sodium perborate ($NaBO_3$)). In certain embodiments, the reaction is carried out in the presence of 9-BBN. In certain embodiments, the reaction involves addition of $NaBO_3 \cdot H_2O$.

[00389] In certain embodiments, the step of hydrating is carried out in the presence of 9-BBN, followed by $NaBO_3 \cdot H_2O$. In certain embodiments, the reaction is carried out in a solvent such as THF. In certain embodiments, the reaction is carried out at 0 °C to room temperature. In certain embodiments, the reaction is carried out under the following conditions: 3 equivalents 9-BBN in THF, from 0 °C to room temperature (*e.g.*, over 1 hour) followed by the addition of aqueous $NaBO_3 \cdot H_2O$.

[00390] In certain embodiments, R^{P1} and R^{P3} are silyl protecting groups; and R^{P10} is hydrogen. In certain embodiments, R^{P1} is TBS; R^{P3} is TES; and R^{P10} is hydrogen.

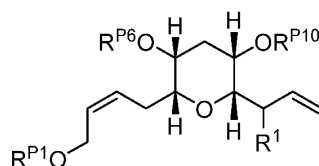
[00391] Provided herein is a method of preparing a compound of Formula (L-5-23A):



(L-5-23A),

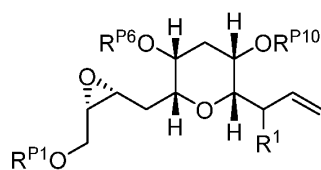
or a salt thereof, the method comprising the steps of:

(a) epoxidizing a compound of Formula (L-5-22):



(L-5-22),

or a salt thereof, to yield a compound of Formula (L-5-22A):



(L-5-22A),

or a salt thereof, and

(b) cyclizing a compound of Formula (L-5-22A), or a salt thereof, to yield a compound of Formula (L-5-23A), or a salt thereof.

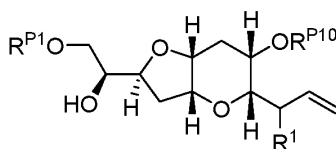
[00392] Any epoxidation reagent may be used in the step of epoxidizing described above. In certain embodiments, the epoxidation reagent is a peracid (*e.g.*, *m*-CPBA). In certain embodiments, the epoxidation reagent is an organometallic reagent. In certain embodiments, the epoxidation reagent is a titanium reagent (*e.g.*, $\text{Ti}(\text{Oi-Pr})_4$). In certain embodiments, the epoxidation reagent is a vanadium reagent (*e.g.*, $\text{VO}(\text{TMHD})_2$). In certain embodiments, the epoxidation is a Sharpless epoxidation. In certain embodiments, the epoxidation is an asymmetric epoxidation (*e.g.*, Sharpless asymmetric epoxidation). In certain embodiments, the epoxidation is carried out in the presence of one or more chiral ligands (*e.g.*, (+)- or (-)-DET, (+)- or (-)-DIPT; wherein DET = diethyltartrate and DIPT = diisopropyltartrate). In certain embodiments, the step of epoxidizing is carried out in the presence of one or more additional reagents. In certain embodiments, epoxidation is carried out in the presence of a peroxide (*e.g.*, *t*-BuOOH).

[00393] In certain embodiments, the step of epoxidizing is carried out in the presence of a titanium complex, a tartrate ligand, and a peroxide. In certain embodiments, the reaction is carried out in the presence of $\text{Ti}(\text{Oi-Pr})_4$, (+)-DET, and *t*-BuOOH. In certain embodiments, the reaction is carried out in the presence of molecular sieves. In certain embodiments, the reaction is carried out in the presence of a solvent such as CH_2Cl_2 . In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at around -10 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: 15 mol% $\text{Ti}(\text{Oi-Pr})_4$, 20 mol% (+)-DET, 1.5 equivalents *t*-BuOOH, and 4 Å molecular sieves in CH_2Cl_2 at -10 °C (*e.g.*, for 10-20 hours).

[00394] In certain embodiments, R^{P6} and R^{P10} are silyl protecting groups; and R^{P1} is hydrogen. In certain embodiments, R^{P6} and R^{P10} are TBS; and R^{P1} is hydrogen. In certain

embodiments, R^{P6} is deprotected before the step of cyclizing a compound of Formula (L-5-22A).

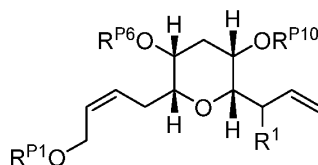
[00395] In certain embodiments, the epoxidation/cyclization provides a compound of Formula (L-5-22B):



(L-5-22B),

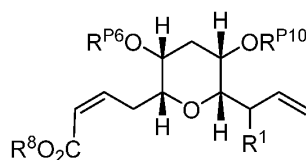
or a salt thereof, which can then be protected to yield a compound of Formula (L-5-23A), or a salt thereof (*e.g.*, to install the group R^{P3} , wherein R^{P3} is an oxygen protecting group).

[00396] As shown in *Scheme 4B*, provided herein is a method of preparing a compound of Formula (L-5-22):



(L-5-22),

or a salt thereof, the method comprising a step of reducing a compound of Formula (L-5-21B):



(L-5-21B),

or a salt thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P6} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

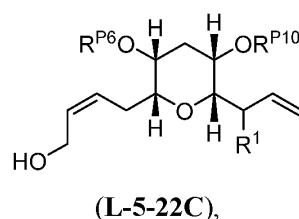
[00397] The step of reducing a compound of (L-5-21B), or a salt thereof, converts the $-CO_2R^8$ moiety to an $-OR^{P1}$ group (*i.e.*, $-OH$). In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H^-) source. Any hydride source known in the art

may be used in this transformation. Examples of hydride sources include, but are not limited to, lithium aluminum hydride, sodium borohydride, lithium borohydride, and diisobutylaluminum hydride. In certain embodiments, the hydride source is diisobutylaluminum hydride (DIBAL).

[00398] In certain embodiments, the step of reducing is carried out in the presence of DIBAL. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, THF). In certain embodiments, the reaction is carried out at below room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at around -78 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: 4 equivalents of DIBAL in THF at -78 °C (*e.g.*, for under 1 hour).

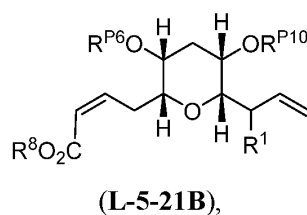
[00399] In certain embodiments, R^{P6} and R^{P10} are silyl protecting groups; and R⁸ is optionally substituted alkyl. In certain embodiments, R^{P6} and R^{P10} are TBS; and R⁸ is methyl.

[00400] In certain embodiments, the compound of Formula (L-5-22) is of Formula (L-5-22-C):



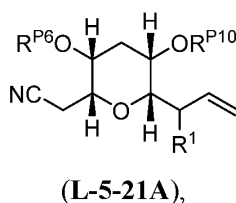
or a salt thereof.

[00401] Also provided herein is a method of preparing a compound of Formula (L-5-21B):

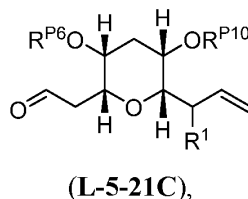


or a salt thereof, the method comprising the steps of:

(a) reducing a compound of Formula (L-5-21A):



or a salt thereof, to yield a compound of Formula (L-5-21C):



or a salt thereof; and

(b) reacting a compound of Formula (L-5-21C), or a salt thereof, in the presence of an olefinating reagent to yield a compound of Formula (L-5-21B), or a salt thereof.

[00402] The step of reducing a compound of (L-5-21A), or a salt thereof (*i.e.*, step (a) above), converts the $-CN$ moiety to an aldehyde group (*i.e.*, $-CHO$). In certain embodiments, the step of reducing is carried out in the presence of a hydride (*i.e.*, H^-) source. Any hydride source known in the art may be used in this transformation. Examples of hydride sources include, but are not limited to, lithium aluminum hydride, sodium borohydride, lithium borohydride, and diisobutylaluminum hydride. In certain embodiments, the hydride source is diisobutylaluminum hydride (DIBAL). The step of reducing may optionally comprise reducing the $-CN$ moiety to an alcohol, followed by oxidation of the resulting alcohol to an aldehyde to yield a compound of Formula (L-5-21C), or a salt thereof.

[00403] In certain embodiments, the step of reducing is carried out in the presence of DIBAL. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, hexanes, CH_2Cl_2). In certain embodiments, the reaction is carried out at below room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately $-78^\circ C$ to approximately room temperature. In certain embodiments, the reaction is carried out at around $-78^\circ C$. For example, in certain embodiments, the reaction is carried out under the following conditions: 1.1 equivalents of DIBAL in hexanes- CH_2Cl_2 at $-78^\circ C$ (*e.g.*, for under 1 hour).

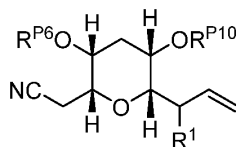
[00404] In certain embodiments, the olefination of a compound of Formula (L-5-21C), or a salt thereof (*i.e.*, step (b) above), is carried out in the presence of a Wittig or Horner-Wadsworth Emmons reagent. In certain embodiments, the olefination is carried out in the presence of a reagent of the formula: $(RO)_2P(O)CH_2CO_2R^8$. In certain embodiments, the reagent is of the formula: $(MeO)_2P(O)CH_2CO_2R^8$ (*e.g.*, $(MeO)_2P(O)CH_2CO_2Bn$). In certain embodiments, the reagent is of the formula: $(CF_3CH_2O)_2P(O)CH_2CO_2R^8$ (*e.g.*, $(CF_3CH_2O)_2P(O)CH_2CO_2Me$). In certain embodiments, the olefination is carried out in the presence of a base. In certain embodiments, the base is a phosphate salt such as K_3PO_4 . In

certain embodiments, the base is an amide base. In certain embodiments, the base is a diisopropyl amide base (*e.g.*, LDA). In certain embodiments, the base is a hexamethyldisilazide base (*e.g.*, LiHMDS, NaHMDS, KHMDS). In certain embodiments, the olefination is carried out in the presence of one or more additional reagents. In certain embodiments, the olefination is carried out in the presence of a crown ether (*e.g.*, 18-crown-6).

[00405] In certain embodiments, the olefination is carried out in the presence of a reagent of the formula $(RO)_2P(O)CH_2CO_2R^8$, a base. In certain embodiments, the reaction is carried out in the presence of $(CF_3CH_2O)_2P(O)CH_2CO_2Me$ and KHMDS. In certain embodiments, 18-crown-6 is present. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, THF). In certain embodiments, the reaction is carried out at a temperature ranging from approximately $-78\text{ }^{\circ}\text{C}$ to approximately room temperature. In certain embodiments, the reaction is carried out at $-78\text{ }^{\circ}\text{C}$. For example, in certain embodiments, the reaction is carried out under the following conditions: 1.5 equivalents $(CF_3CH_2O)_2P(O)CH_2CO_2Me$, 1.5 equivalents KHMDS, 8 equivalents 18-crown-6, in THF at $-78\text{ }^{\circ}\text{C}$ (*e.g.*, for under 1 hour).

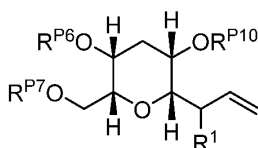
[00406] In certain embodiments, R^{P6} and R^{P10} are silyl protecting groups; and R^8 is optionally substituted alkyl. In certain embodiments, R^{P6} and R^{P10} are TBS; and R^8 is methyl.

[00407] Also provided herein is a method of preparing a compound of Formula (L-5-21A):



(L-5-21A),

or a salt thereof, the method comprising reacting a compound of Formula (L-5-20):



(L-5-20),

or a salt thereof, in the presence of cyanide; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl;

R^{P6} , R^{P7} and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and wherein $-OR^{P7}$ is a leaving group.

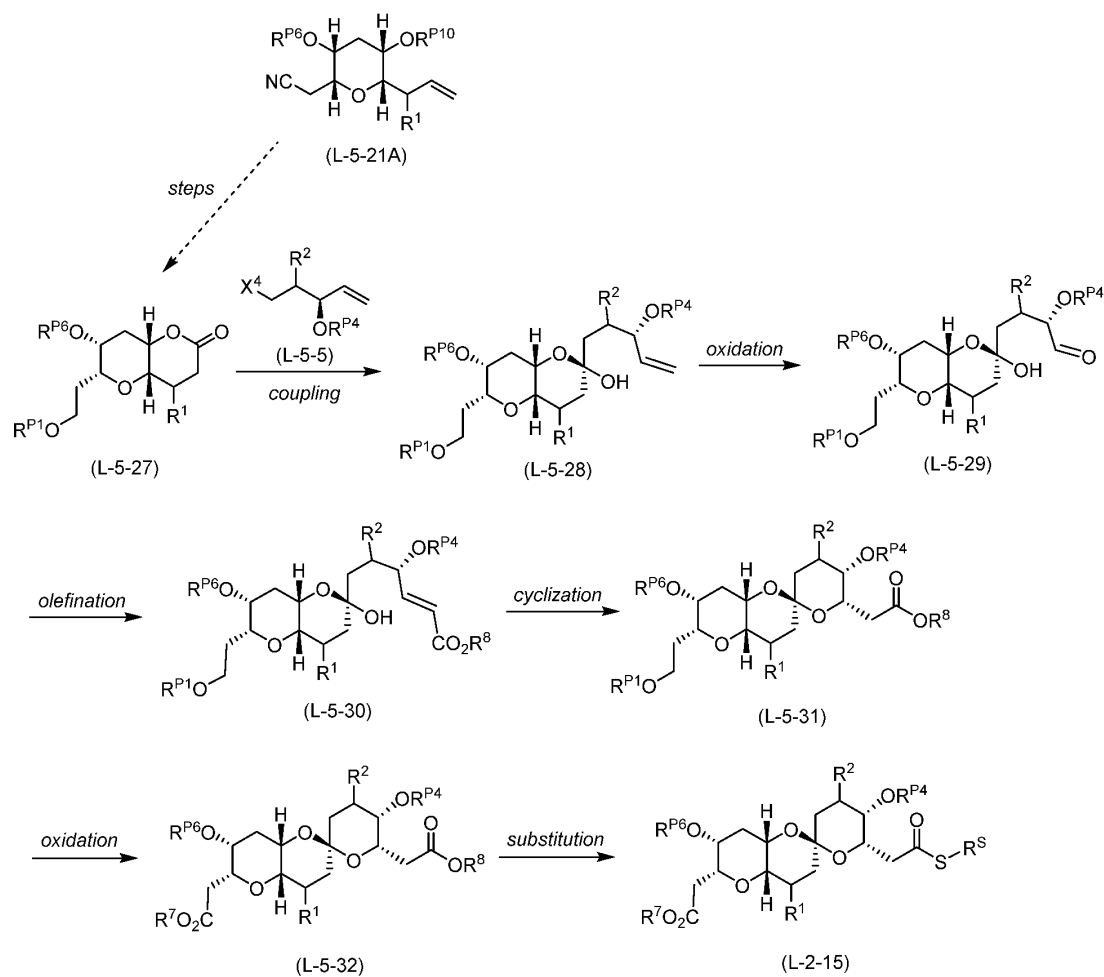
[00408] The method of preparing a compound of Formula (L-5-21A), or a salt thereof, comprises reacting a compound of Formula (L-5-20), or a salt thereof, in the presence of cyanide. In certain embodiments, the cyanide is a cyanide salt (e.g., NaCN, KCN, LiCN). In certain embodiments, the cyanide salt is sodium cyanide (NaCN). The reaction may be carried out in the presence of one or more additional reagents (e.g., a crown ether). In certain embodiments, the reaction is carried out in the presence of NaCN, in a solvent such as DMSO. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 20 equivalents NaCN in DMSO at room temperature (e.g., for 1 hour).

[00409] In certain embodiments, R^{P6} and R^{P10} are silyl protecting groups. In certain embodiments, R^{P6} and R^{P10} are TBS.

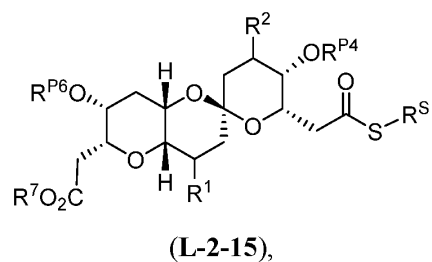
Preparation of Left Halves of Norhalichondrins

[00410] Provided herein are method of preparing “left half” building blocks of compounds in the norhalichondrin series (e.g., norhalichondrin A, B, C, and analogs thereof). For example, as shown in *Scheme 4C*, left half building blocks of Formula (L-2-15) can be prepared by converting the ester group (i.e., -CO₂R⁸) of a compound of Formula (L-5-32) to a thioester moiety (i.e., -C(O)SR^S). To this end, a compound of Formula (L-5-32) can be prepared by oxidizing a compound of Formula (L-5-31), which can be prepared by cyclizing a compound of Formula (L-5-30). A compound of Formula (L-5-30) can be prepared via oxidative cleavage and olefination of a compound of Formula (L-5-28), which can be obtained by coupling a compound of Formula (L-5-27) with a compound of Formula (L-5-5). A compound of Formula (L-5-27) can be obtained from an intermediate of Formula (L-5-21A), as described herein.

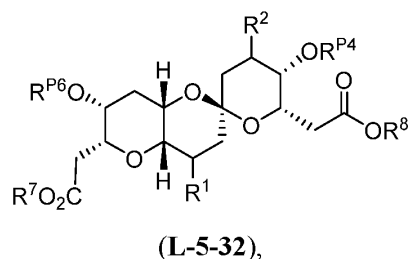
Scheme 4C



[00411] As shown in *Scheme 4C*, provided herein is a method of preparing a compound of Formula (L-2-15):



or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-32):



or a salt thereof, in the presence of a thiolating agent; wherein:

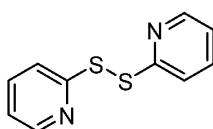
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^7 and R^8 are independently hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl.

[00412] As described herein, the step of forming a compound of Formula (L-2-15) comprises reacting a compound of Formula (L-5-32) in the presence of a thiolating agent. Any thiolating agent known in the art may be used to this end. In certain embodiments, the thiolating agent is a disulfide. In certain embodiments, the thiolating agent is of the formula $(R^S S)_2$. In certain embodiments, the thiolating agent is of the formula (pyridine-S)₂. In certain



embodiments, the thiolating agent is:

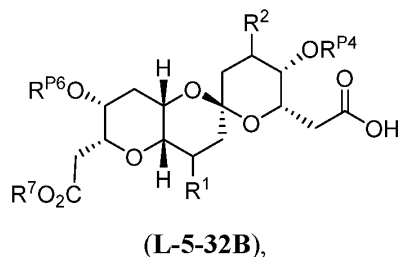
[00413] In certain embodiments, the step of thiolating a compound of Formula (L-5-32) is carried out in the presence of one or more additional reagents. In certain embodiments, the step of thiolating is carried out in the presence of a phosphine reagent (*e.g.*, triphenylphosphine (Ph_3P)).

[00414] In certain embodiments, the step of thiolating is carried out in the presence of a disulfide and a phosphine. In certain embodiments, the reaction is carried out in the presence of $(Py-S)_2$ and Ph_3P . In certain embodiments, the reaction is carried out in a solvent such as toluene or CH_2Cl_2 . In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the step of

thiolating is carried out under the following conditions: 1.4 equivalents of (Py-S)₂, 1.2 equivalents of Ph₃P, in toluene at room temperature (*e.g.*, for 10-20 hours).

[00415] In certain embodiments, the method of thiolating a compound of Formula (L-5-32), or a salt thereof, comprises the steps of:

- (a) deprotecting a compound of Formula (L-5-32), or a salt thereof, to yield a compound of Formula (L-5-32B):

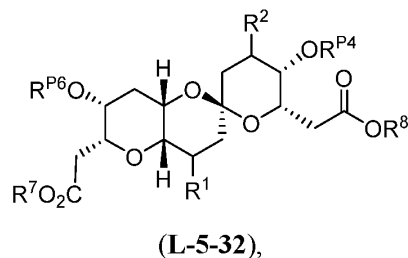


or a salt thereof; and

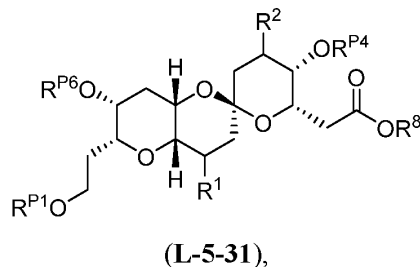
- (b) thiolating a compound of Formula (L-5-32B), or a salt thereof, to yield a compound of Formula (L-2-15), or a salt thereof.

[00416] In certain embodiments, R⁷ is optionally substituted alkyl; and R^{P6} and R^{P4} are silyl protecting groups. In certain embodiments, R⁷ is optionally substituted alkyl; R^{P6} and R^{P4} are TES.

[00417] Also provided herein is a method of preparing a compound of Formula (L-5-32):



or a salt thereof, the method comprising oxidizing a compound of Formula (L-5-31):



or a salt thereof; wherein:

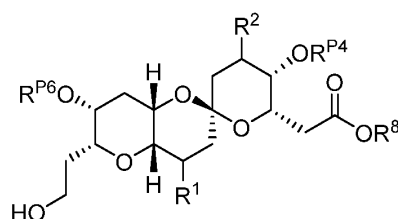
- R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P4} , and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

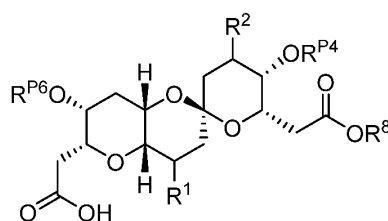
[00418] In certain embodiments, the method of preparing a compound of Formula (L-5-32), or a salt thereof, comprises the steps of:

(a) Oxidizing a compound of Formula (L-5-31B):



(L-5-31B),

or a salt thereof, to yield a compound of Formula (L-5-32C):



(L-5-32C),

or a salt thereof; and

(b) protecting a compound of Formula (L-5-32C), or a salt thereof, to yield a compound of Formula (L-5-32), or a salt thereof.

[00419] Any method can be used in the step of oxidizing a compound of Formula (L-5-31) or (L-5-31B). In certain embodiments, the oxidation is carried out in the presence of a periodinane (*e.g.*, Dess-Martin periodinane (DMP)). In certain embodiments, the oxidation involves a Swern oxidation. In certain embodiments, the oxidation is carried out in the presence of a chromium reagent (*e.g.*, pyridinium chlorochromate (PCC)). In certain embodiments, the step of oxidizing involves a Pinnick oxidation, *e.g.*, treatment of the reaction mixture with a chlorite (*e.g.*, sodium chlorite (NaClO₂)). In certain embodiments, the oxidation involves carrying out the reaction in the presence of a periodinane (*e.g.*, DMP) followed by a chlorite (*e.g.*, NaClO₂). In certain embodiments, the oxidation is carried out in the presence of DMP and NaHCO₃ in a solvent (*e.g.*, CH₂Cl₂), followed by NaClO₂ and

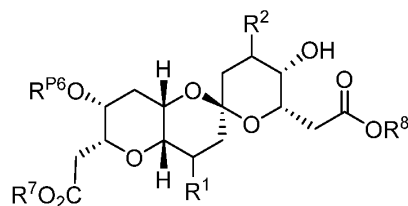
NaH₂PO₄ in a solvent (*e.g.*, *t*-BuOH/H₂O). In certain embodiments, the reactions are carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reactions are carried out at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: (a) 2 equivalents DMP, 10 equivalents NaHCO₃ in CH₂Cl₂ at room temperature (*e.g.*, for under 1 hour); followed by (b) 3 equivalents NaClO₂, 4 equivalents NaH₂PO₄, with 2-methyl-2-butene in *t*-BuOH and water at room temperature (*e.g.*, for under 1 hour).

[00420] In certain embodiments, the step of protecting a compound of Formula (L-5-32C) involves treating the compound with an alkylating agent. In certain embodiments, the alkylating agent is an alkyl halide or a reagent of the structure: alkyl-leaving group. In certain embodiments, the alkylating agent is a methyl transfer reagent (*e.g.*, diazomethane, trimethylsilyldiazomethane (TMSCH₂N₂)).

[00421] In certain embodiments, the step of protecting is carried out in the presence of TMSCH₂N₂. In certain embodiments, the reaction is carried out in a solvent (*e.g.*, benzene/MeOH). In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 3 equivalents TMSCH₂N₂ in benzene/MeOH at room temperature (*e.g.*, for 5 min).

[00422] In certain embodiments, R⁷ is optionally substituted alkyl; and R^{P6} is a silyl protecting group; R^{P4} is optionally substituted benzyl; and R⁸ is optionally substituted benzyl. In certain embodiments, R⁷ is methyl; R^{P6} is TES; R^{P4} is MPM; and R⁸ is benzyl.

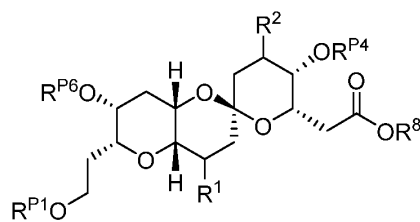
[00423] In certain embodiments, the compound of Formula (L-5-32), or a salt thereof, is deprotected to remove the group R^{P4} yield a compound of Formula (L-5-32D):



(L-5-32D),

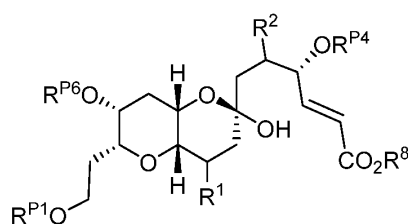
or a salt thereof; and optionally re-protected (*i.e.*, to switch the group R^{P4} from, *e.g.*, a benzyl protecting group (*e.g.*, MPM) to a silyl protecting group (*e.g.*, trialkylsilyl such as triethylsilyl).

[00424] As shown in *Scheme 4C*, provided herein is a method of preparing a compound of Formula (L-5-31):



(L-5-31),

or a salt thereof, the method comprising a step of cyclizing a compound of Formula (L-5-32A):



(L-5-30),

or a salt thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

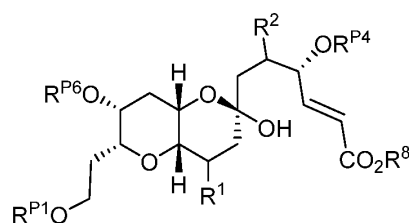
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00425] In certain embodiments, the step of cyclizing a compound of Formula (7-5-30) is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amidine or guanidine base. In certain embodiments, the base is an amine or an amide. In certain embodiments, the base is an amidine base (*e.g.*, 1,8-diazabicyclo(5.4.0)undec-7-ene (DBU)). In certain embodiments, the step of cyclizing is carried out in the presence of an acid. In certain embodiments, the acid is a Lewis acid. In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt (*e.g.*, LiBr, LiCl). The step of cyclizing may be carried out in the presence of one or more additional reagents. In certain embodiments, the step of cyclizing is carried out in the presence of BnOAc.

[00426] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt, and a base. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr and DBU. In certain embodiments, the reaction is carried out in a solvent such as MeCN. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 10 equivalents LiBr, 5 equivalents DBU, and 2 equivalents BnOAc in MeCN at room temperature (*e.g.*, for 10-20 hours).

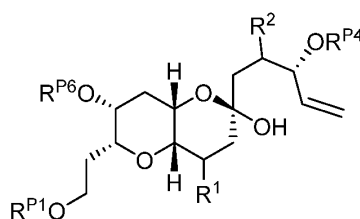
[00427] In certain embodiments, R^{P1} is a silyl; and R^{P6} is a silyl protecting group; R^{P4} is optionally substituted benzyl; and R^8 is optionally substituted benzyl. In certain embodiments, R^{P1} is TES; R^{P6} is TES; R^{P4} is MPM; and R^8 is benzyl.

[00428] Also provided herein is a method of preparing a compound of Formula (L-5-30):



(L-5-30),

or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-28):

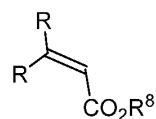


(L-5-28),

or a salt thereof, in the presence of an olefin and an olefin metathesis catalyst; wherein:

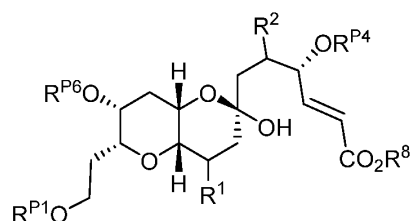
R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocycl.



[00429] In certain embodiments, the olefin is of the formula: $\text{R}^2\text{C}=\text{CH}-\text{CO}_2\text{R}^8$. Further, any olefin metathesis known in the art may be used in the metathesis reaction to furnish a compound of Formula (L-5-30).

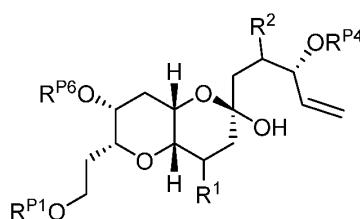
[00430] Also provided herein is an alternative method of preparing a compound of Formula (L-5-30):



(L-5-30),

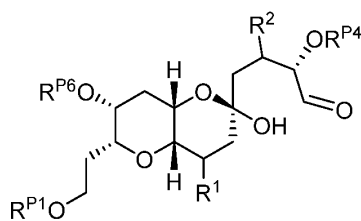
or a salt thereof, the method comprising the steps of:

(a) oxidizing a compound of Formula (L-5-28):

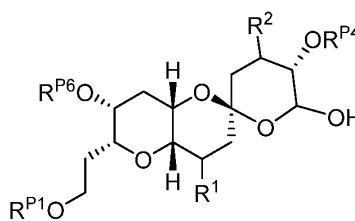


(L-5-28),

or a salt thereof, to yield a compound of Formula (L-5-29) or (L-5-29B):



(L-5-29),



(L-5-29B),

or a salt thereof; and

(b) reacting the compound of Formula (L-5-29) or (L-5-29B), or a salt thereof, in the presence of a olefination reagent, to yield a compound of Formula (L-5-30), or a salt thereof. The reaction in step (a) above is an oxidative cleavage; the reaction in step (b) is an olefination reaction. In certain embodiments, the oxidative cleavage is carried out via ozonolysis (*e.g.*, in the presence of O_3). In certain embodiments, the cleavage is carried out in

the presence of reagents capable of dihydroxylating a double bond (*e.g.*, osmium tetroxide (OsO_4), N-methylmorpholine N-oxide (NMMO)), followed by a transition metal (*e.g.*, a lead complex such as $\text{Pb}(\text{OAc})_4$).

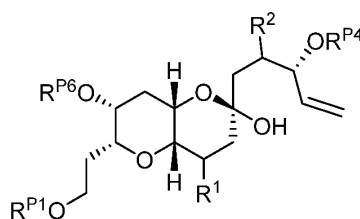
[00431] In certain embodiments, the double bond is dihydroxylated by treatment with OsO_4 , NMMO, and water. In certain embodiments, the reaction is carried out in the presence of a solvent such as acetone. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the double bond is dihydroxylated under the following conditions: 5 mol% OsO_4 , 2 equivalents NMMO, and water, in acetone at room temperature (*e.g.*, for 10-20 hours). The resulting compound is then treated, in certain embodiments, with $\text{Pb}(\text{OAc})_4$ and K_2CO_3 to yield the aldehyde or hemiacetal. For example, in certain embodiments, this step is carried out under the following conditions: 2 equivalents $\text{Pb}(\text{OAc})_4$, 10 equivalents K_2CO_3 , in CH_2Cl_2 at room temperature (*e.g.*, for under 1 hour).

[00432] In certain embodiments, the olefination is carried out in the presence of a Wittig or Horner-Wadsworth Emmons reagent. In certain embodiments, the olefination is carried out in the presence of a reagent of the formula: $(\text{RO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$. In certain embodiments, the reagent is of the formula: $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$ (*e.g.*, $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$). In certain embodiments, the olefination is carried out in the presence of a base (*e.g.*, a phosphate salt such as K_3PO_4).

[00433] In certain embodiments, the olefination is carried out in the presence of an olefination reagent of the formula: $(\text{RO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{R}^8$, and a base. In certain embodiments, the olefination is carried out in the presence of $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$ and K_3PO_4 . In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: 4 equivalents $(\text{MeO})_2\text{P}(\text{O})\text{CH}_2\text{CO}_2\text{Bn}$, 3 equivalents K_3PO_4 , in a solvent at room temperature (*e.g.*, for 24-48 hours).

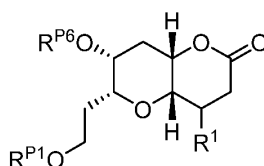
[00434] In certain embodiments, $\text{R}^{\text{P}1}$ is a silyl; and $\text{R}^{\text{P}6}$ is a silyl protecting group; $\text{R}^{\text{P}4}$ is optionally substituted benzyl; and R^8 is optionally substituted benzyl. In certain embodiments, $\text{R}^{\text{P}1}$ is TES; $\text{R}^{\text{P}6}$ is TES; $\text{R}^{\text{P}4}$ is MPM; and R^8 is benzyl.

[00435] Also provided herein is a method of preparing a compound of Formula (L-5-28):



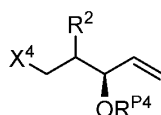
(L-5-28),

or a salt thereof, the method comprising a step of coupling a compound of Formula (L-5-27):



(L-5-27),

or a salt thereof, with a compound of Formula (L-5-5):



(L-5-5),

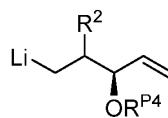
or a salt thereof, wherein:

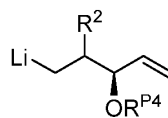
X⁴ is halogen or a leaving group;

R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl; and

each instance of R^{P1}, R^{P4}, and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00436] In certain embodiments, the coupling of a compound of Formula (L-5-4) with a compound of Formula (L-5-5) is carried out in the presence of an organometallic reagent (*e.g.*, to convert X⁴ to a metal for addition to the compound of Formula (L-5-4)). In certain embodiments, the organometallic reagent is a lithium reagent (*e.g.*, to convert the compound



of the Formula (L-5-5) to a compound of the formula:  for addition to the compound of Formula (L-5-4)). In certain embodiments, lithium reagent is an organolithium (*e.g.*, *n*-butyllithium, *tert*-butyllithium, *sec*-butyllithium). In certain embodiments, the lithium reagent is LiHMDS or LDA.

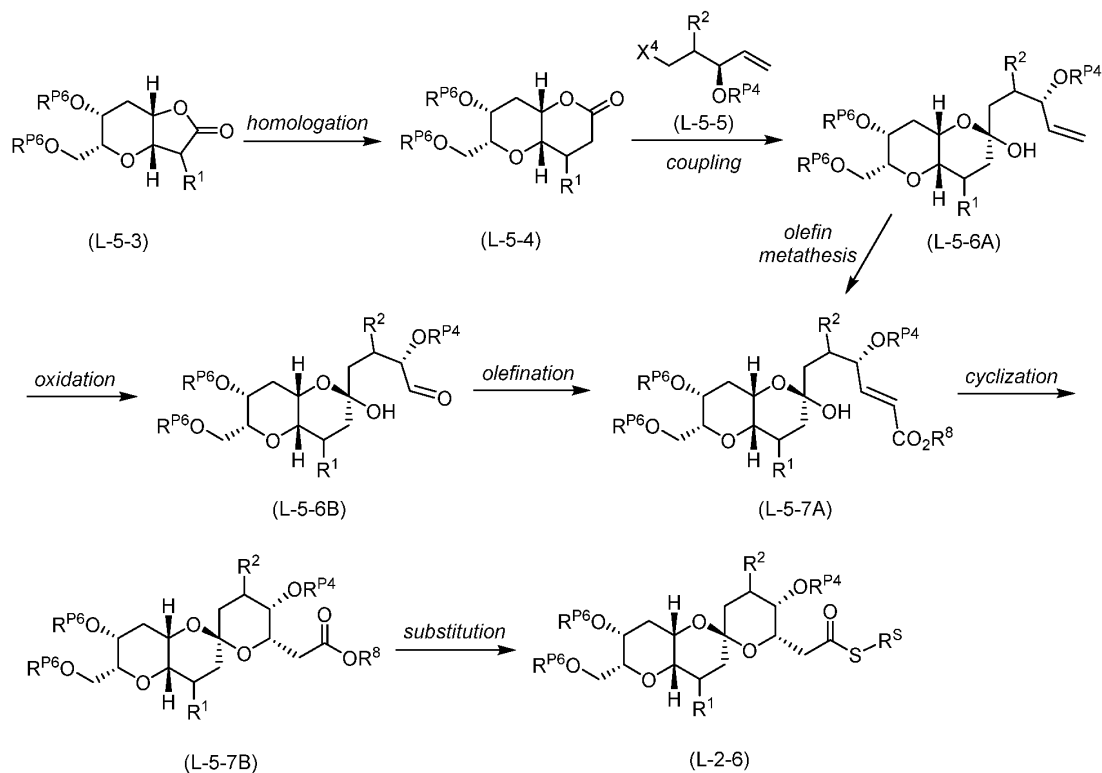
[00437] In certain embodiments, the reaction is carried out in the presence of *tert*-butyllithium. In certain embodiments, the reaction is performed in a solvent such as toluene, THF, Et₂O, or a combination thereof. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at -78° C. For example, in certain embodiments, the reaction is carried out with 2.2 equivalents of *tert*-butyllithium in toluene and Et₂O at -78° C (*e.g.*, for less than 1 hour).

[00438] In certain embodiments, R^{P1} is a silyl; and R^{P6} is a silyl protecting group; R^{P4} is optionally substituted benzyl; and R⁸ is optionally substituted benzyl. In certain embodiments, R^{P1} is TES; R^{P6} is TES; R^{P4} is MPM; and R⁸ is benzyl.

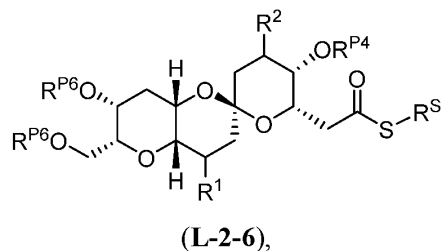
Preparation Left Halves of Halichondrin Analogs

[00439] Provided herein are methods useful in the preparation of “left half” building blocks of other halichondrin analogs (*e.g.*, compounds of Formula (H3-2-1)). For example, as shown in Scheme 4D, left half building blocks of Formula (L-2-6) can be prepared by converting the ester group (*i.e.*, -CO₂R⁸) of a compound of Formula (L-5-7B) to a thioester moiety (*i.e.*, -C(O)SR^S). A compound of Formula (L-5-7B) can be prepared by cyclizing a compound of Formula (L-5-7A), which may be prepared by oxidative cleavage and olefination of a compound of Formula (L-5-6A). A compound of Formula (L-5-6A) can be prepared by coupling a compound of Formula (L-5-4) with a compound of Formula (L-5-5). As also shown in scheme 4D, a compound of Formula (L-5-4) can be prepared via homologation of a lactone of Formula (L-5-3).

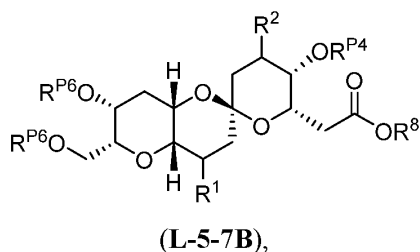
Scheme 4D



[00440] As shown in *Scheme 4D*, provided herein is a method of preparing a compound of Formula (L-2-6):



or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-7B):



or a salt thereof, in the presence of a thiolating agent; wherein:

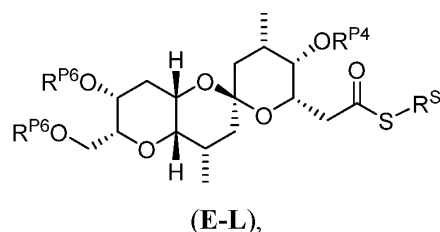
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

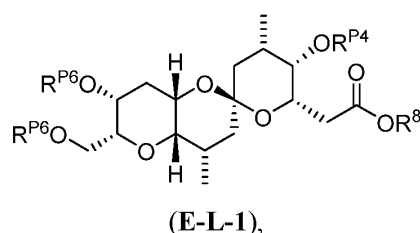
each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00441] In certain embodiments, the method is a method of preparing a compound of Formula (E-L):



or a salt thereof, the method comprising a step of reacting a compound of Formula (E-L-1):



or a salt thereof, in the presence of a thiolating agent; wherein:

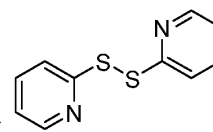
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00442] As described herein, the step of forming a compound of Formula (L-2-6), (E-L), or a salt thereof, comprises reacting a compound of Formula (L-5-7B), (E-L-1), or a salt thereof, in the presence of a thiolating agent. Any thiolating agent known in the art may be used to

this end. In certain embodiments, the thiolating agent is a disulfide. In certain embodiments, the thiolating agent is of the formula $(R^S S)_2$. In certain embodiments, the thiolating agent is of



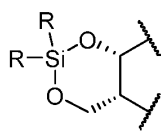
the formula (pyridine-S)₂. In certain embodiments, the thiolating agent is: (2,2'-dipyridyl sulfide). In certain embodiments, the thiolating reagent is present in stoichiometric or excess amounts (*e.g.*, 1-2 equivalents)

[00443] In certain embodiments, the step of thiolating is carried out in the presence of one of more additional reagents. In certain embodiments, the step of thiolating is carried out in the presence of a phosphine reagent. In certain embodiments, the phosphine is a trialkyl phosphine. In certain embodiments, the phosphine is a triaryl phosphine. In certain embodiments, the phosphine is PPh₃. In certain embodiments, the phosphine is polymer-bound PPh₃. In certain embodiments, the phosphine is present in stoichiometric or excess amounts (*e.g.*, 1-3 equivalents).

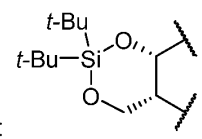
[00444] In certain embodiments, the step of thiolating is carried out in the presence of a disulfide and a phosphine. In certain embodiments, the reaction is carried out in the presence of 2,2'-dipyridyl sulfide and Ph₃P. In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is DCM. In certain embodiments, the solvent is acetonitrile. In certain embodiments, the reaction is carried out at from 0° C to room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. In certain embodiments, the reaction is carried out in the presence of 2,2'-dipyridyl sulfide and Ph₃P in MeCN at from 0° C to room temperature.

[00445] For example, in certain embodiments, the step of thiolating is carried out under the following conditions: 1.4 equivalents of 2,2'-dipyridyl sulfide, 1.2 equivalents of Ph₃P, in DCM at room temperature (*e.g.*, for 10-20 hours). For example, in certain embodiments, the step of thiolating is carried out under the following conditions: 1.2 equivalents of 2,2'-dipyridyl sulfide, 2.3 equivalents of Ph₃P, in MeCN at from 0 °C to room temperature (*e.g.*, for 10-20 hours).

[00446] In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring



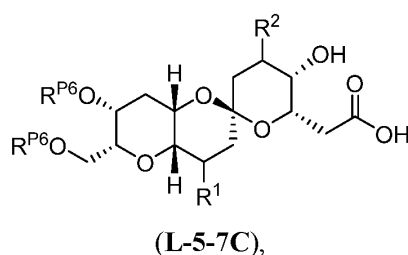
of the formula: ; and R^{P4} is a silyl protecting group. In certain embodiments,



two R^{P6} are joined with the intervening atoms to form a ring of the formula:
and R^{P4} is TES.

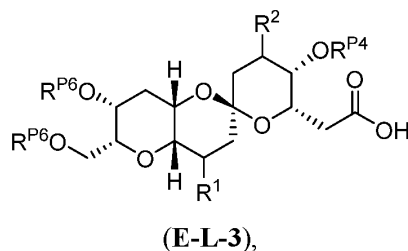
[00447] In certain embodiments, the method of thiolating a compound of Formula (L-5-7B), or a salt thereof, comprises:

(a) a step of deprotecting a compound of Formula (L-5-7B), or a salt thereof, under conditions sufficient to remove the R^{P4} and R^8 groups, to yield a compound of Formula (L-5-7C):



or a salt thereof; and

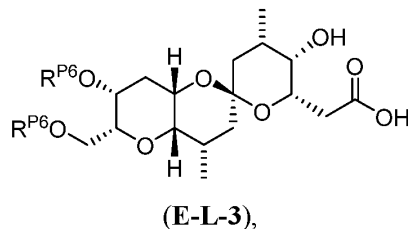
(b) a step of protecting a compound of Formula (L-5-7C), or a salt thereof, to yield a compound of Formula (L-5-7D):



or a salt thereof.

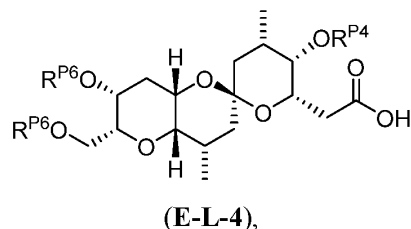
[00448] In certain embodiments, the method comprises:

(a) a step of deprotecting a compound of Formula (E-L-1), or a salt thereof, under conditions sufficient to remove the R^{P4} and R^8 groups, to yield a compound of Formula (E-L-3):



or a salt thereof; and

(b) a step of protecting a compound of Formula (E-L-3), or a salt thereof, to yield a compound of Formula (E-L-4):



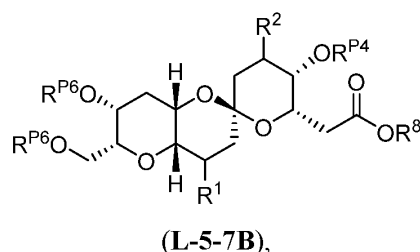
or a salt thereof.

[00449] In certain embodiments, with respect to the compounds of Formula (L-5-7B), (E-L-1), or a salt thereof, R^{P4} and R^8 are optionally substituted benzyl protecting groups; and the step of deprotecting (*i.e.*, step (a)) is carried out in the presence of H_2 and Pd/C. In certain embodiments, R^{P4} is MPM and R^8 is benzyl (Bn); and the step of deprotecting is carried out in the presence of H_2 and Pd/C. In certain embodiments, the step of deprotecting is carried out in the presence of H_2 and Pd/C in *i*-PrOAc.

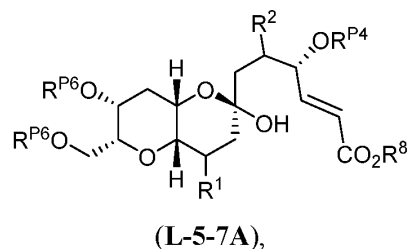
[00450] In certain embodiments, with respect to the compound of Formula (E-L-4), (L-5-7D), or salt thereof, R^{P4} is a silyl protecting group; and the step of protecting (*i.e.*, step (b)) is carried out in the presence of a silylating agent and base. In certain embodiments, R^{P4} is TES; and the silylating reagent is TESCl. In certain embodiments, the base is imidazole. In certain embodiments, the step of protecting is carried out in the presence of TESCl and imidazole. In certain embodiments, the step of protecting is carried out in the presence of TESCl and imidazole in DMF.

[00451] In certain embodiments, the compounds of Formulae (E-L-4), (L-5-7D), or salts thereof, are purified by silica gel chromatography and/or purification.

[00452] As also shown in *Scheme 4D*, provided herein is a method of preparing a compound of Formula (L-5-7B):



or a salt thereof, the method comprising cyclizing a compound of Formula (L-5-7A):



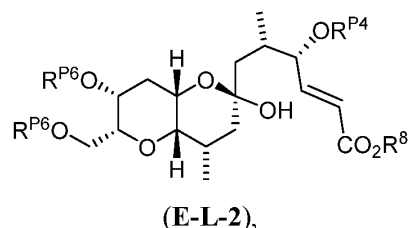
or a salt thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

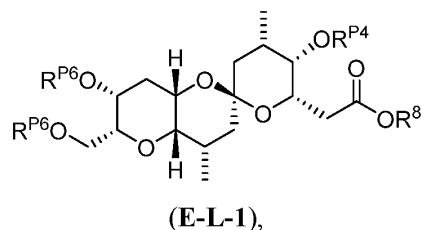
each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00453] In certain embodiments, the method comprises cyclizing a compound of Formula (E-L-2):



or a salt thereof, to yield a compound of Formula (E-L-1):



or a salt thereof, wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00454] In certain embodiments, the step of cyclizing a compound of Formula (7-5-7A), (E-L-2), or a salt thereof, is carried out in the presence of a base. In certain embodiments, the base is a nitrogen base. In certain embodiments, the base is an amidine, guanidine base. In certain embodiments, the base is an amine or amide base. In certain embodiments, the base is an amidine base (*e.g.*, 1,8-diazabicyclo(5.4.0)undec-7-ene (DBU)). In certain embodiments, the base is DBU. In certain embodiments, the base is used in an excess amount.

[00455] In certain embodiments, the step of cyclizing is carried out in the presence of an acid. In certain embodiments, the acid is a Lewis acid.

[00456] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt (*e.g.*, LiBr, LiCl). In certain embodiments, the reaction is carried out in the presence of LiBr. In certain embodiments, the base is used in an excess amount.

[00457] The step of cyclizing may be carried out in the presence of one or more additional reagents. In certain embodiments, the step of cyclizing is carried out in the presence of a reagent of the formula: R^8OAc . In certain embodiments, the step of cyclizing is carried out in the presence of $BnOAc$. In certain embodiments, the reagent is present in an excess amount.

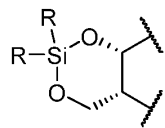
[00458] In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the solvent is MeCN. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. In certain embodiments, the reaction is carried out at around 30 °C.

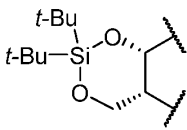
[00459] In certain embodiments, the step of cyclizing is carried out in the presence of a lithium salt and a base. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr and DBU. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr, DBU, and R^8OAc . In certain embodiments, the step of cyclizing is carried out in the presence of LiBr, DBU, and $BnOAc$. In certain embodiments, the step of cyclizing is carried out in the presence of LiBr, DBU, and $BnOAc$ in MeCN from room temperature to around 30 °C.

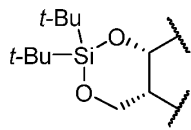
[00460] For example, in certain embodiments, the reaction is carried out under the following conditions: 10 equivalents LiBr, 5 equivalents DBU, and 10 equivalents $BnOAc$ in MeCN at room temperature (*e.g.*, for 10-20 hours). For example, in certain embodiments, the reaction

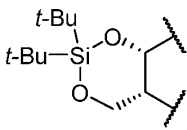
is carried out under the following conditions: 10 equivalents LiBr, 5 equivalents DBU, and 5 equivalents BnOAc in MeCN at room temperature to around 30 °C (*e.g.*, for around 24 hours).

[00461] In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring

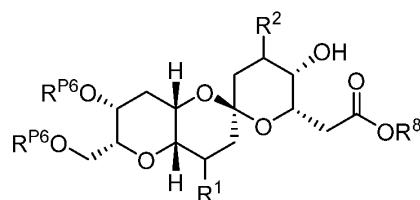


of the formula: ; and R^{P4} and R^{P8} are optionally substituted benzyl groups. In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring of the



formula: ; R^{P4} is MPM; and R^{P8} is benzyl.

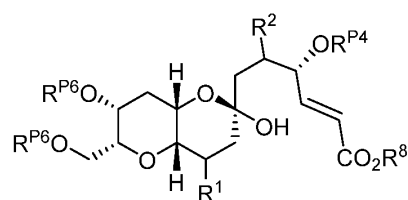
[00462] In certain embodiments, the compound of Formula (L-5-7B), or a salt thereof, is deprotected to remove the group R^{P4} yield a compound of Formula (L-5-7D):



(L-5-7D),

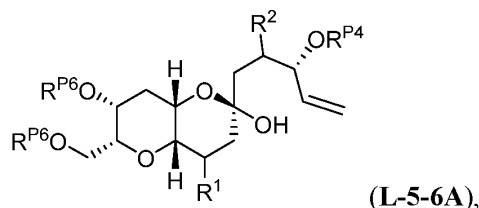
or a salt thereof; and optionally re-protected (*i.e.*, to switch the group R^{P4} from, *e.g.*, a benzyl protecting group (*e.g.*, MPM) to a silyl protecting group (*e.g.*, trialkylsilyl such as triethylsilyl).

[00463] Also provided herein is a method of preparing a compound of Formula (L-5-7A):



(L-5-7A),

or a salt thereof, the method comprising a step of reacting a compound of Formula (L-5-6A):

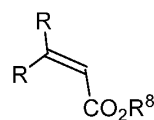


or a salt thereof, in the presence of an olefin and an olefin metathesis catalyst; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

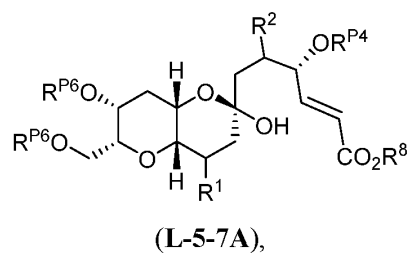
each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.



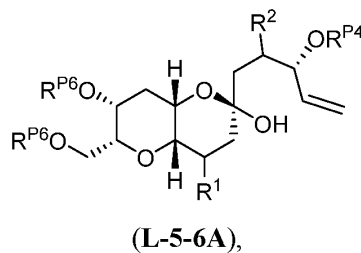
[00464] In certain embodiments, the olefin is of the formula: $\text{R}-\text{C}(\text{R})=\text{CH}-\text{CO}_2\text{R}^8$. Further, any olefin metathesis known in the art may be used in the metathesis reaction to furnish a compound of Formula (L-5-7A).

[00465] Also provided herein is an alternative method of preparing a compound of Formula (L-5-7A):

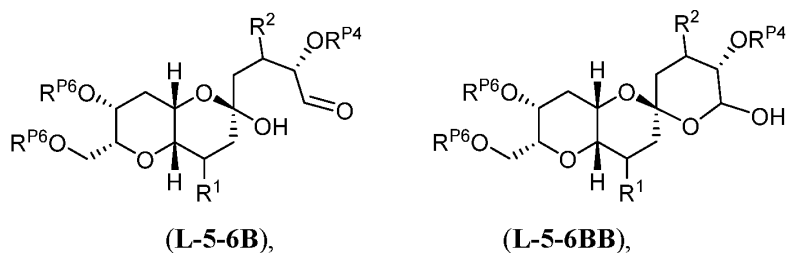


or a salt thereof, the method comprising the steps of:

(a) oxidizing a compound of Formula (L-5-6A):



or a salt thereof, to yield a compound of Formula (L-5-6B) and/or (L-5-6BB):

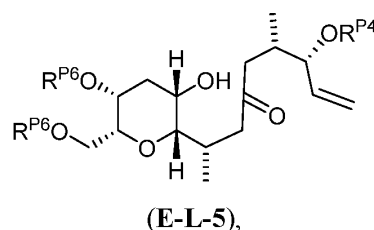


or a salt thereof; and

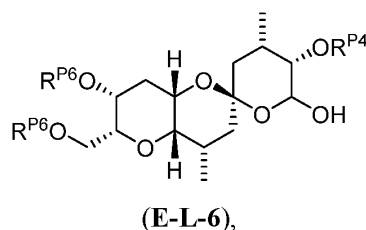
- (b) reacting the compound of Formula (L-5-6B) and/or (L-5-6BB), or a salt thereof, in the presence of a olefination reagent, to yield a compound of Formula (L-5-7A), or a salt thereof.

[00466] In certain embodiments, the method comprises the steps of:

- (a) oxidizing a compound of Formula (E-L-5):

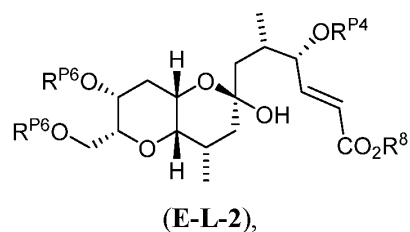


or a salt thereof, to yield a compound of Formula (E-L-6):



or a salt thereof; and

- (b) reacting the compound of Formula (E-L-6), or a salt thereof, in the presence of a olefination reagent, to yield a compound of Formula (E-L-2):



or a salt thereof, wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00467] The oxidation of a compound of Formula (L-5-6A), (E-L-5), or a salt thereof (*i.e.*, step (a)) above is an oxidative cleavage. In certain embodiments, the oxidative cleavage is carried out via ozonolysis (*e.g.*, in the presence of O_3). In certain embodiments, the oxidizing cleavage is a Johnson-Lemieux oxidative cleavage. For example, In certain embodiments, the cleavage is carried out in the presence of reagents capable of dihydroxylating a double bond (*e.g.*, osmium tetroxide (OsO_4) and N-methylmorpholine N-oxide (NMO); or potassium osmate (VI) dehydrate (K_2OsO_4) and NMO), followed by a transition metal (*e.g.*, a lead complex such as $Pb(OAc)_4$). In certain embodiments, the cleavage is carried out in the presence of reagents capable of dihydroxylating a double bond (*e.g.*, osmium tetroxide (OsO_4) and N-methylmorpholine N-oxide (NMO); or potassium osmate (VI) dehydrate (K_2OsO_4) and NMO), followed by sodium periodate ($NaIO_4$).

[00468] In certain embodiments, the double bond is dihydroxylated by treatment with OsO_4 , NMO, and water. In certain embodiments, the reaction is carried out in the presence of a solvent such as acetone. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. For example, in certain embodiments, the double bond is dihydroxylated under the following conditions: 10 mol% OsO_4 , 2 equivalents NMO, and water, in acetone at room temperature (*e.g.*, for 20-25 hours). The resulting compound is then treated, in certain embodiments, with $Pb(OAc)_4$ and K_2CO_3 to yield the aldehyde and/or hemiacetal. For example, in certain embodiments, this step is carried out under the following conditions: 1.2 equivalents $Pb(OAc)_4$, 3 equivalents K_2CO_3 , in CH_2Cl_2 at room temperature (*e.g.*, for approximately 1 hour).

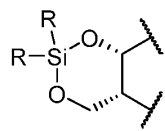
[00469] In certain embodiments, the step of oxidizing is carried out in the presence of osmium tetroxide (OsO_4) or potassium osmate (VI) dehydrate (K_2OsO_4), and NMO; followed by $NaIO_4$. In certain embodiments, the step of oxidizing is carried out in the presence of potassium osmate (VI) dehydrate (K_2OsO_4) and NMO, followed by $NaIO_4$. In certain embodiments, the reaction is carried out in a solvent. In certain embodiments, the reaction is

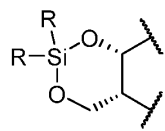
carried out in acetone and water. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at around room temperature. In certain embodiments, the reaction is carried out in the presence of K_2OsO_4 and NMO, followed by $NaIO_4$, in acetone and water, at around room temperature. For example, in certain embodiments, the reaction is carried out under the following conditions: $K_2OsO_4 \cdot 2H_2O$ and NMO, followed by $NaIO_4$, in acetone and water, at around room temperature.

[00470] In certain embodiments, the olefination in step (b) is carried out in the presence of a Wittig or Horner-Wadsworth Emmons reagent. In certain embodiments, the olefination is carried out in the presence of a reagent of the formula: $(RO)_2P(O)CH_2CO_2R^8$. In certain embodiments, the reagent is of the formula: $(MeO)_2P(O)CH_2CO_2R^8$ (*e.g.*, $(MeO)_2P(O)CH_2CO_2Bn$). In certain embodiments, the olefination is carried out in the presence of a base (*e.g.*, a phosphate salt such as K_3PO_4).

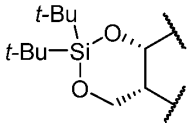
[00471] In certain embodiments, the olefination is carried out in the presence of an olefination reagent of the formula: $(RO)_2P(O)CH_2CO_2R^8$, and a base. In certain embodiments, the olefination is carried out in the presence of $(MeO)_2P(O)CH_2CO_2Bn$ and K_3PO_4 . In certain embodiments, the reaction is carried out in a solvent such as toluene. In certain embodiments, the reaction is carried out at a temperature ranging from approximately 0 °C to approximately 50 °C. In certain embodiments, the reaction is carried out at room temperature. In certain embodiments, the reaction is carried out at around 30 °C. In certain embodiments, the olefination is carried out in the presence of $(MeO)_2P(O)CH_2CO_2Bn$ and K_3PO_4 , in toluene at around 30 °C. For example, in certain embodiments, the reaction is carried out under the following conditions: 4 equivalents $(MeO)_2P(O)CH_2CO_2Bn$, 3 equivalents K_3PO_4 at room temperature (*e.g.*, for about 20-25 hours). For example, in certain embodiments, the reaction is carried out under the following conditions: 5 equivalents $(MeO)_2P(O)CH_2CO_2Bn$, 4 equivalents K_3PO_4 at around 30 °C (*e.g.*, for about 1-3 days).

[00472] In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring

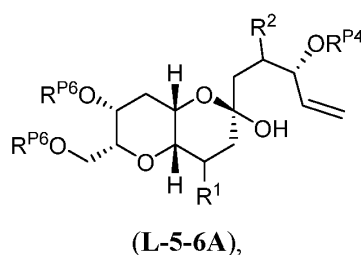


of the formula: ; and R^{P4} and R^{P8} are optionally substituted benzyl groups. In

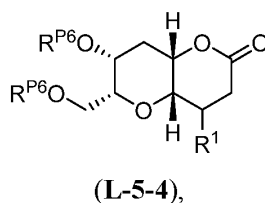
certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring of the

formula: ; R^{P4} is MPM; and R^{P8} is benzyl.

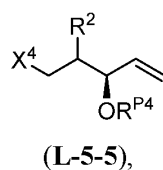
[00473] Also provided herein is a method of preparing a compound of Formula (L-5-6A):



or a salt thereof, the method comprising a step of coupling a compound of Formula (L-5-4):



or a salt thereof, with a compound of Formula (L-5-5):



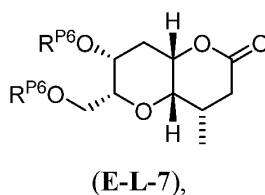
or a salt thereof, wherein:

X^4 is halogen or a leaving group;

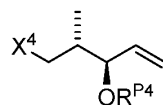
R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocycl.

[00474] In certain embodiments, the method comprises comprising a step of coupling a compound of Formula (E-L-7):

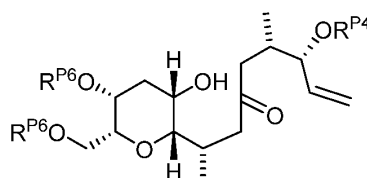


or a salt thereof, with a compound of Formula (E-L-8):



(E-L-8),

or a salt thereof, to yield a compound of Formula (E-L-5):



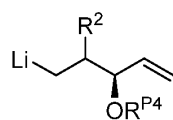
(E-L-5),

or a salt thereof, wherein:

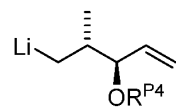
X^4 is halogen or a leaving group; and

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocycl.

[00475] In certain embodiments, the coupling of a compound of Formula (L-5-4) with a compound of Formula (L-5-5) (or a compound of the formula (E-L-7) and (E-L-8)) is carried out in the presence of an organometallic reagent (*e.g.*, to convert X^4 to a metal for addition to the compound of Formula (L-5-4) or (E-L-7)). In certain embodiments, the organometallic reagent is a lithium reagent (*e.g.*, to convert the compound of the Formula (L-5-5) to a



compound of the formula: for addition to the compound of Formula (L-5-4); *e.g.*, to convert the compound of the Formula (E-L-8) to a compound of the formula:



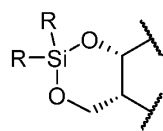
for addition to the compound of Formula (E-L-7)). In certain embodiments, lithium reagent is an organolithium (*e.g.*, *n*-butyllithium, *tert*-butyllithium, *sec*-butyllithium). In certain embodiments, the lithium reagent is LiHMDS or LDA. In certain embodiments, the lithium reagent is *sec*-butyllithium.

[00476] In certain embodiments, the reaction is carried out in the presence of *tert*-butyllithium. In certain embodiments, the reaction is performed in a solvent such as THF. In certain embodiments, the reaction is carried out at a temperature ranging from approximately

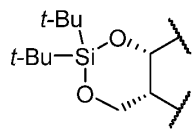
-78° C to approximately room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately 0 °C. For example, in certain embodiments, the reaction is carried out with 2.6 equivalents of *tert*-butyllithium in THF from -78° C to room temperature (*e.g.*, over less than 1 hour).

[00477] In certain embodiments, the reaction is carried out in the presence of *sec*-butyllithium. In certain embodiments, the reaction is performed in THF. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately room temperature. In certain embodiments, the reaction is carried out at a temperature ranging from approximately -78 °C to approximately 0 °C. In certain embodiments, the reaction is carried out with *sec*-butyllithium in THF at around -78° C to room temperature. For example, in certain embodiments, the reaction is carried out with about 2 equivalents of *sec*-butyllithium in THF from -78° C to room temperature (*e.g.*, over less than 1 hour).

[00478] In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring



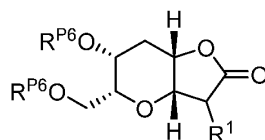
of the formula: ; and R^{P4} is optionally substituted benzyl. In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring of the formula:



; and R^{P4} is MPM.

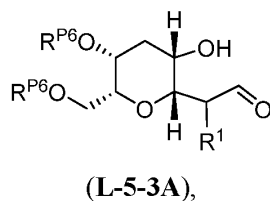
[00479] As shown in *Scheme 4D*, provided herein is a method of preparing a compound of Formula (L-5-4) from a compound of Formula (L-5-3). In certain embodiments, the method comprises the steps of:

(a) reducing a compound of Formula (L-5-3):



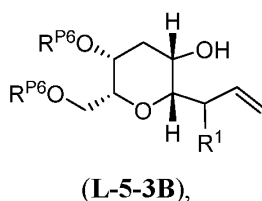
(L-5-3),

or a salt thereof, to yield a compound of Formula (L-5-3A):



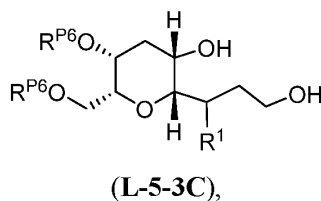
or a salt thereof;

(b) olefinating a compound of Formula (L-5-3A), or a salt thereof, to yield a compound of Formula (L-5-3B):



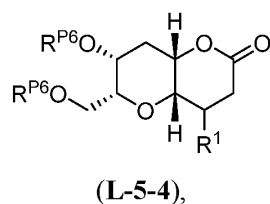
or a salt thereof;

(c) hydrating a compound of Formula (L-5-3B), or a salt thereof, to yield a compound of Formula (L-5-3C):



or a salt thereof; and

(d) oxidizing and cyclizing a compound of Formula (L-5-3C), or a salt thereof, to yield a compound of Formula (L-5-4):

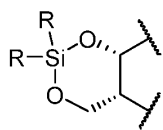


or a salt thereof.

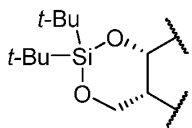
[00480] The step of reducing in step (a) above may be carried out in the presence of a hydride source. In certain embodiments, the hydride source is DIBAL. In certain embodiments, the step of olefination in step (b) above may be carried out in the presence of an olefination reagent (*e.g.*, MePPh₃Br). In certain embodiments, the step of olefination is carried out in the presence of a base (*e.g.*, an alkoxide such as *t*-BuOK). In certain

embodiments, the step of hydrating in step (c) above is a hydroboration reaction. In certain embodiments, the step of hydroboration involves treatment with 9-BBN followed by $\text{NaBO}_3 \cdot \text{H}_2\text{O}$. The steps of oxidizing and cyclizing in step (d) above may be carried out in the same step or subsequent steps. The step of oxidizing may be carried out in the presence of any oxidizing agents. In certain embodiments, the step of oxidizing is carried out in the presence of TEMPO and $\text{PhI}(\text{OAc})_2$. In certain embodiments, the step of oxidizing is carried out in the presence of NaHCO_3 .

[00481] In certain embodiments, two R^{P6} are joined with the intervening atoms to form a ring



of the formula: . In certain embodiments, two R^{P6} are joined with the



intervening atoms to form a ring of the formula:

General Reaction Parameters

[00482] The following embodiments apply to all synthetic methods described above and herein.

[00483] The reactions provided and described herein may involve one or more reagents. In certain embodiments, a reagent may be present in a catalytic amount. In certain embodiments, a catalytic amount is from 0-1 mol%, 0-5 mol%, 0-10 mol%, 1-5 mol%, 1-10 mol%, 5-10 mol%, 10-20 mol%, 20-30 mol%, 30-40 mol%, 40-50 mol%, 50-60 mol%, 60-70 mol%, 70-80 mol%, 80-90 mol%, or 90-99 mol%. In certain embodiments, a reagent may be present in a stoichiometric amount (*i.e.*, about 1 equivalent). In certain embodiments, a reagent may be present in excess amount (*i.e.*, greater than 1 equivalent). In certain embodiments, the excess amount is about 1.1, 1.2, 1.3, 1.4, 1.5, 2.0, 2.5, 3.0, 3.5, 4.0, 4.5, 5.0, 5.5, 6.0, 6.5, 7.0, 7.5, 8.0, 8.5, 9.0, 9.5, 10, 15, or 20 equivalents. In certain embodiments, the excess amount is from about 1.1-2, 2-3, 3-4, 4-5, 1.1-5, 5-10, 10-15, 15-20, or 10-20 equivalents. In certain embodiments, the excess amount is greater than 20 equivalents.

[00484] A reaction described herein may be carried out at any temperature. In certain embodiments, a reaction is carried out at or around room temperature (rt) (21 °C or 70 °F). In certain embodiments, a reaction is carried out at below room temperature (*e.g.*, from -100 °C to 21 °C). In certain embodiments, a reaction is carried out at or around -78 °C. In certain

embodiments, a reaction is carried out at or around -10 °C. In certain embodiments, a reaction is carried out at around 0 °C. In certain embodiments, a reaction is carried out at above room temperature. In certain embodiment, a reaction is carried out at 30, 40, 50, 60, 70, 80, 110, 120, 130, 140, or 150 °C. In certain embodiments, a reaction is carried out at above 150 °C.

[00485] A reaction described herein may be carried out in a solvent, or a mixture of solvents (*i.e.*, cosolvents). Solvents can be polar or non-polar, protic or aprotic. Any solvent may be used in the reactions described herein, and the reactions are not limited to particular solvents or combinations of solvents. Common organic solvents useful in the methods described herein include, but are not limited to, acetone, acetonitrile, benzene, benzonitrile, 1-butanol, 2-butanone, butyl acetate, *tert*-butyl methyl ether, carbon disulfide carbon tetrachloride, chlorobenzene, 1-chlorobutane, chloroform, cyclohexane, cyclopentane, 1,2-dichlorobenzene, 1,2-dichloroethane, dichloromethane (DCM), *N,N*-dimethylacetamide *N,N*-dimethylformamide (DMF), 1,3-dimethyl-3,4,5,6-tetrahydro-2-pyrimidinone (DMPU), 1,4-dioxane, 1,3-dioxane, diethylether, 2-ethoxyethyl ether, ethyl acetate, ethyl alcohol, ethylene glycol, dimethyl ether, heptane, *n*-hexane, hexanes, hexamethylphosphoramide (HMPA), 2-methoxyethanol, 2-methoxyethyl acetate, methyl alcohol, 2-methylbutane, 4-methyl-2-pentanone, 2-methyl-1-propanol, 2-methyl-2-propanol, 1-methyl-2-pyrrolidinone, dimethylsulfoxide (DMSO), nitromethane, 1-octanol, pentane, 3-pentanone, 1-propanol, 2-propanol, pyridine, tetrachloroethylene, tetrahyrdofuran (THF), 2-methyltetrahydrofuran, toluene, trichlorobenzene, 1,1,2-trichlorotrifluoroethane, 2,2,4-trimethylpentane, trimethylamine, triethylamine, *N,N*-diisopropylethylamine, diisopropylamine, water, *o*-xylene, *p*-xylene.

[00486] A reaction described herein may be carried out over any amount of time. In certain embodiments, a reaction is allowed to run for seconds, minutes, hours, or days.

[00487] Methods described herein can be used to prepare compounds in any chemical yield. In certain embodiments, a compound is produced in from 1-10%, 10-20% 20-30%, 30-40%, 40-50%, 50-60%, 60-70%, 70-80%, 80-90%, or 90-100% yield. In certain embodiments, the yield is the percent yield after one synthetic step. In certain embodiments, the yield is the percent yield after more than one synthetic step (*e.g.*, 2, 3, 4, or 5 synthetic steps).

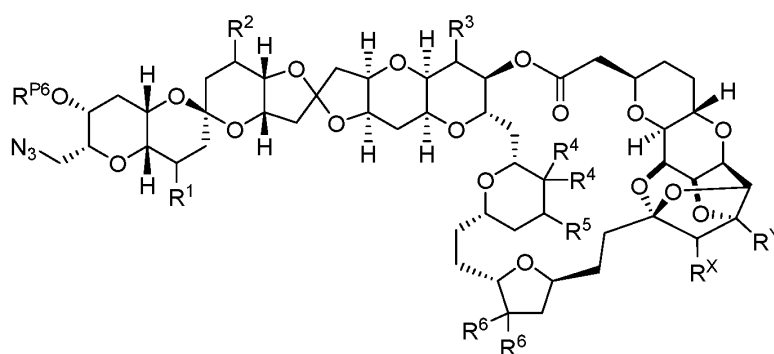
[00488] Methods described herein may further comprise one or more purification steps. For example, in certain embodiments, a compound produced by a method described herein may be purified by chromatography, extraction, filtration, precipitation, crystallization, or any other method known in the art. In certain embodiments, a compound or mixture is carried forward to the next synthetic step without purification (*i.e.*, crude).

[00489] The synthetic method provided herein can be carried out on any scale (*i.e.*, to yield any amount of product). In certain embodiments, the methods are applicable to small-scale synthesis or larger-scale process manufacture. In certain embodiments, a reaction provided herein is carried out to yield less than 1 g of product. In certain embodiments, a reaction provided herein is carried out to yield greater than 1 g, 2 g, 5 g, 10 g, 15 g, 20 g, 25 g, 30 g, 40 g, 50 g, 100 g, 200 g, 500 g, or 1 kg of product.

Compounds

[00490] The present invention also provides novel compounds. The compounds are useful in the preparation of halichondrins, analogs thereof, and intermediates thereto. In certain embodiments, the compounds provided herein are useful in the synthesis of compounds of Formula (H3-A), such as Compound (1), or intermediates thereto.

[00491] Provided herein are compounds of Formula (H3-N3):

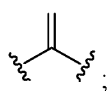


(H3-N3),

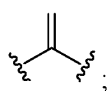
and salts thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

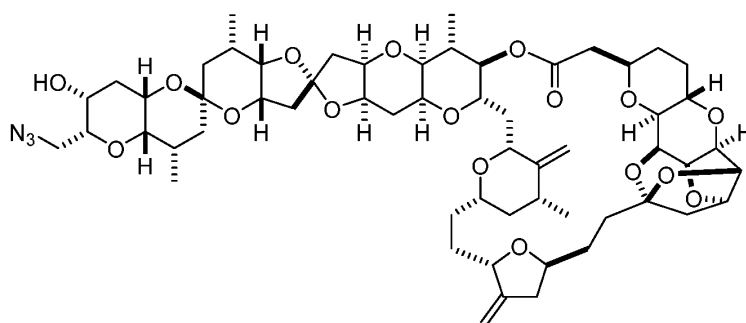
R^{P6} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

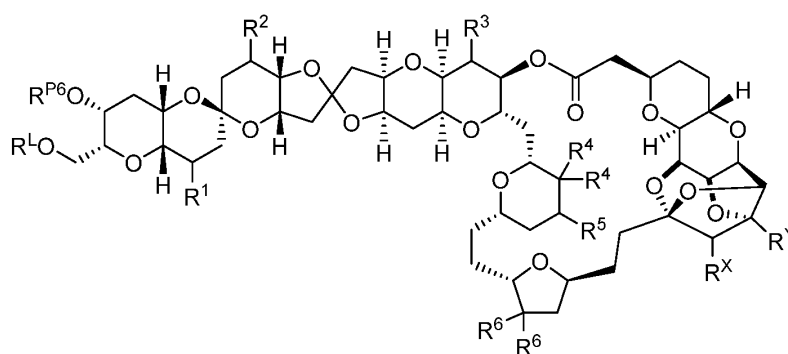
[00492] In certain embodiments, the compound is of the following formula:



Compound (B),

or a salt thereof.

[00493] Provided herein are compounds of Formula (H3-L):



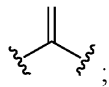
(H3-L),

and salts thereof, wherein:

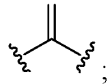
R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

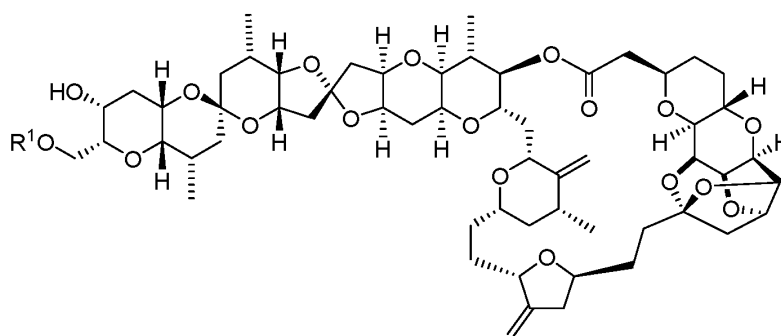
R^{P6} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00494] In certain embodiments, the compound is of the formula:

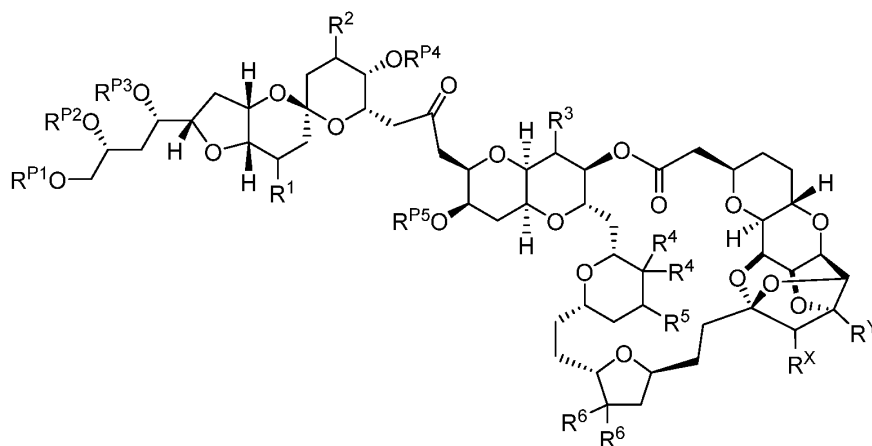


Compound (A),

or a salt thereof, wherein:

R^1 is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

[00495] Provided herein are compounds of Formula (H-2-II):

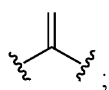


(H-2-II),

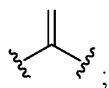
and salts thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

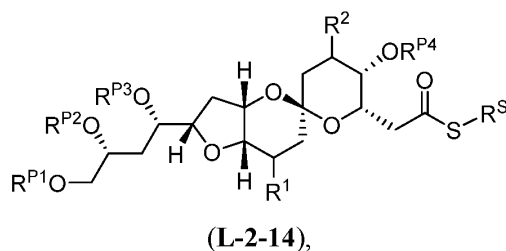
R^{P1} , R^{P2} , R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00496] Provided herein are compounds of Formula (L-2-14):



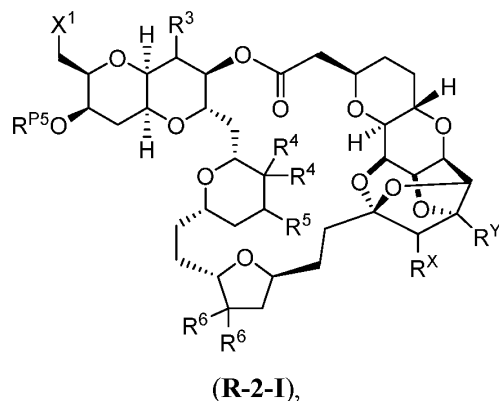
and salts thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R¹ and R² are each independently hydrogen, halogen, or optionally substituted alkyl;
and

R^{P1}, R^{P2}, R^{P3}, and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00497] Provided herein are compounds of Formula (R-2-I):



and salts thereof, wherein:

X¹ is halogen or a leaving group;

R³ and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl; each instance of R⁴ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁴ groups are taken together to form: ;

each instance of R⁶ is independently hydrogen, halogen, or optionally substituted

alkyl, or two R⁶ groups are taken together to form: ;

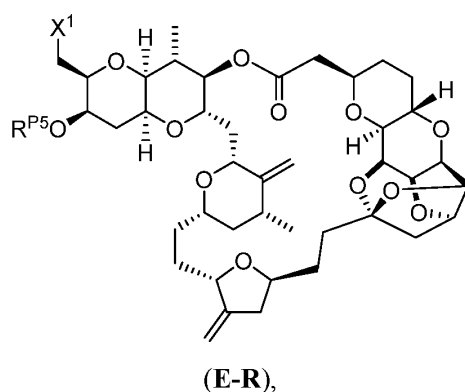
R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocycl.

[00498] In certain embodiments, the compound is of Formula (E-R):

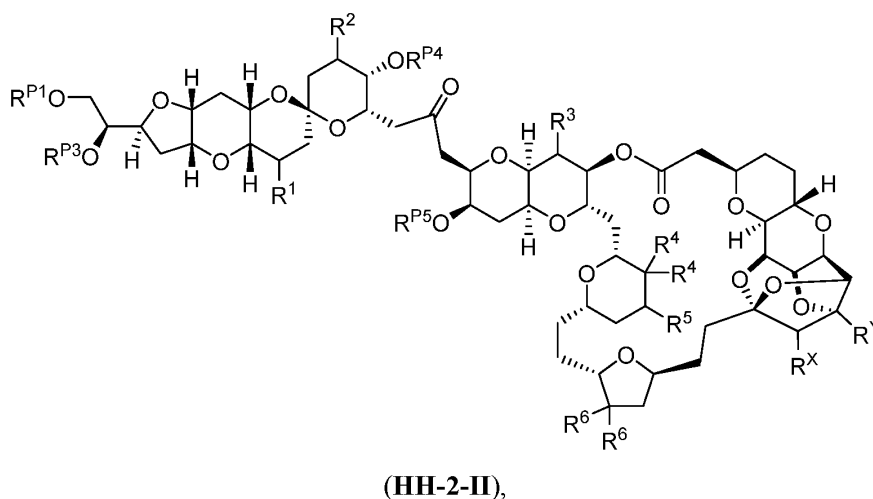


or a salt thereof, wherein:

X^1 is halogen or a leaving group; and

R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

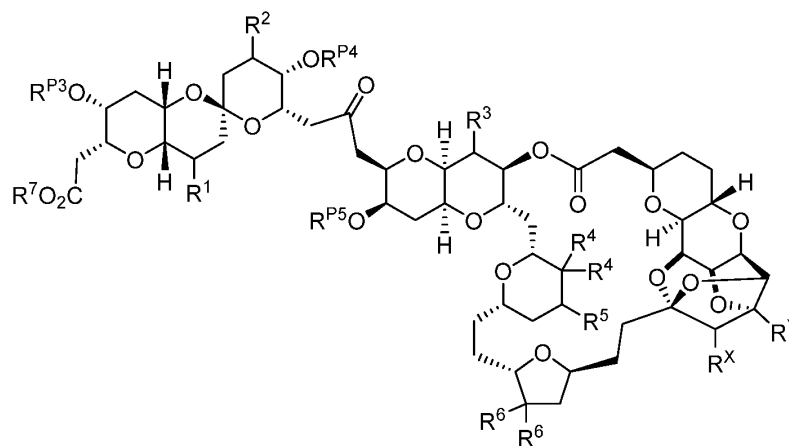
[00499] Provided herein are compounds of Formula (HH-2-II):



and salts thereof, wherein:

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00501] Provided herein are compounds of Formula (NH-2-II):

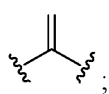


(NH-2-II),

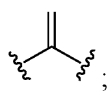
and salts thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

R^{P3} , R^{P4} , and R^{P5} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

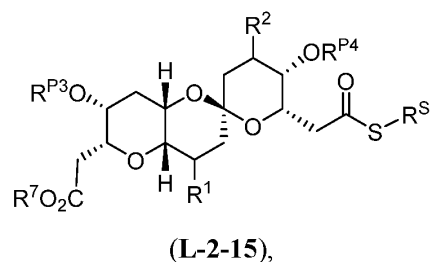
R^7 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00502] Provided herein are compounds of Formula (L-2-15):



and salts thereof, wherein:

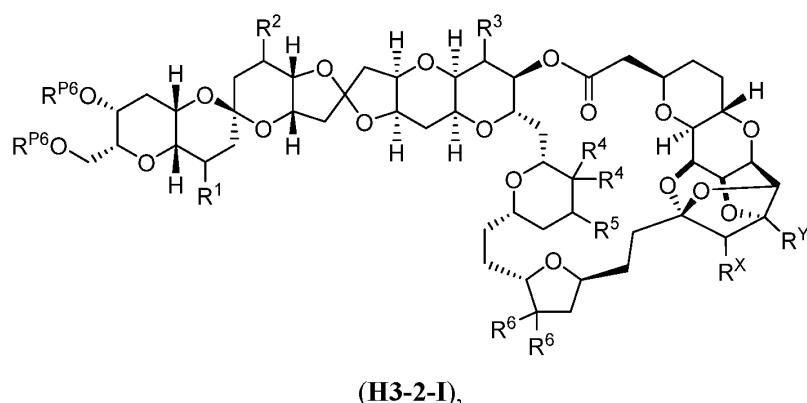
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P3} and R^{P4} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^7 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00503] Provided herein are compounds of Formula (H3-2-I):



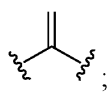
and salts thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

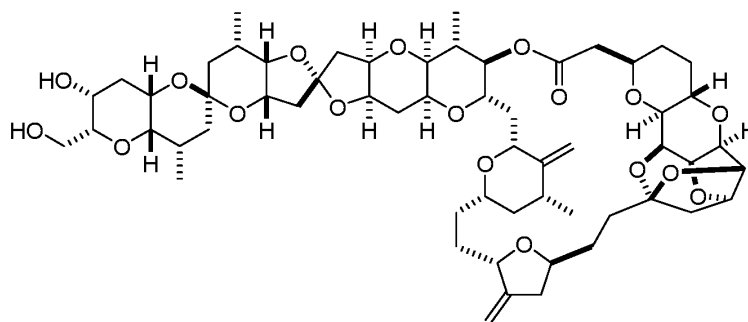
each instance of R^{P6} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

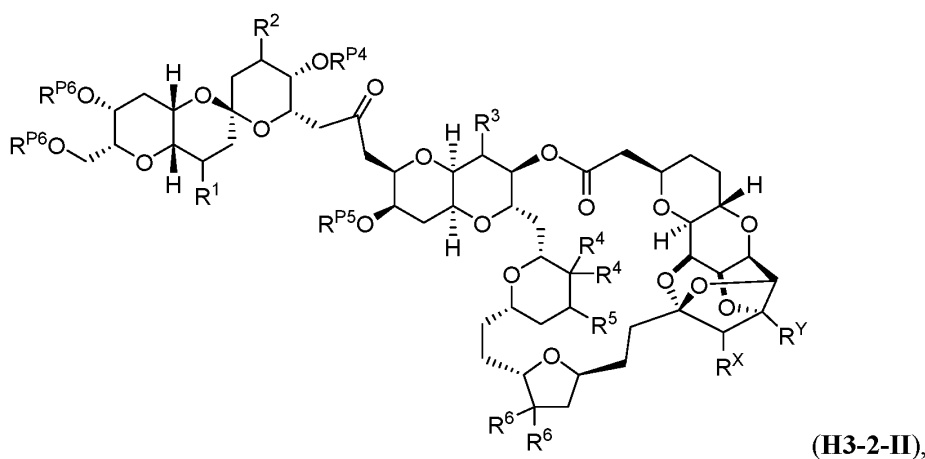
[00504] In certain embodiments, the compound is of the formula:



Compound (2),

or a salt thereof.

[00505] Provided herein are compounds of Formula **(H3-2-II)**:

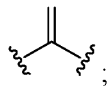


(H3-2-II),

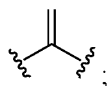
and salts thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

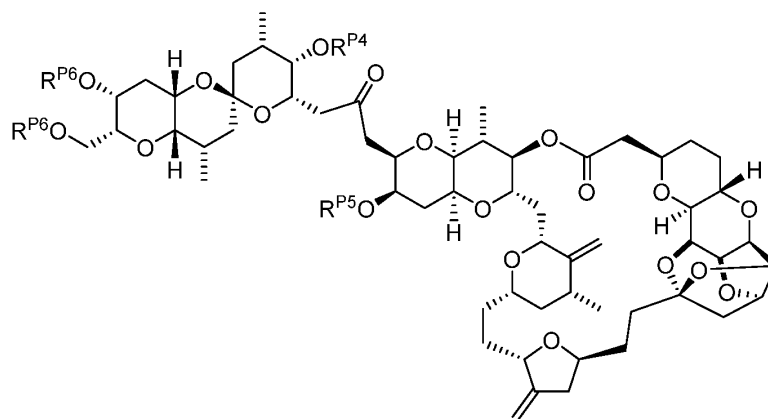
R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00506] In certain embodiments, the compound is of the Formula (E-1):

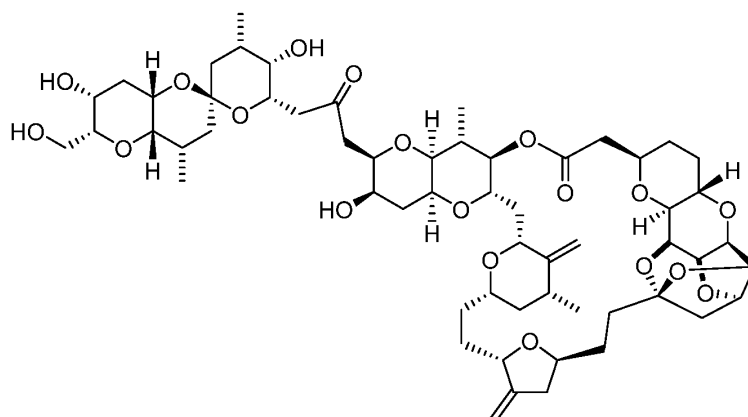


(E-1),

or a salt thereof, wherein:

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

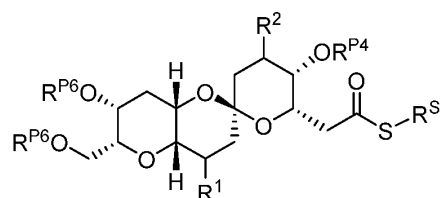
[00507] In certain embodiments, the compound is of the formula:



Compound (C),

or a salt thereof.

[00508] Provided herein are compounds of Formula (L-2-6):



(L-2-6),

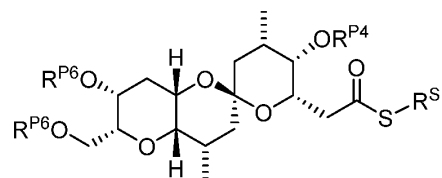
and salts thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P4} and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00509] In certain embodiments, the compound is of Formula (E-L):

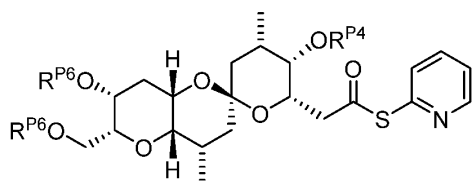


(E-L),

or a salt thereof, wherein:

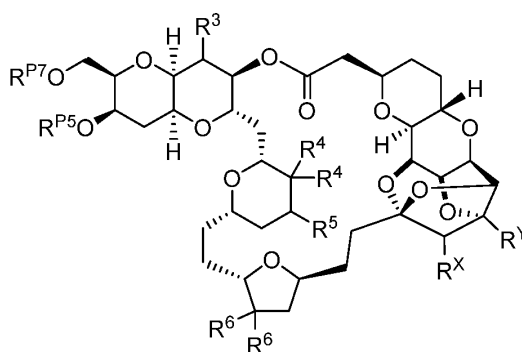
R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl; each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00510] In certain embodiments, the compound is of the formula:



or a salt thereof.

[00511] Provided herein are compounds of Formula (R-4-11B):

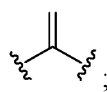


(R-4-11B),

and salts thereof, wherein:

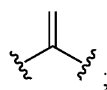
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl; each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form:



each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form:



R^{P5} and R^{P7} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group;

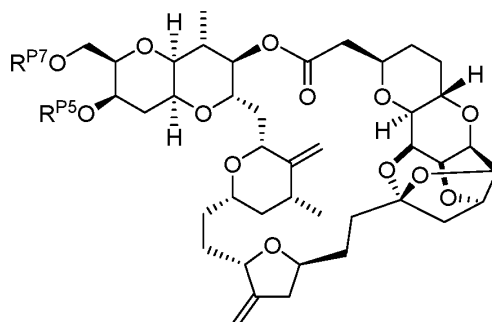
optionally wherein R^{P5} and R^{P7} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00512] In certain embodiments, the compound is of the Formula **(E-R-1)**:



(E-R-1),

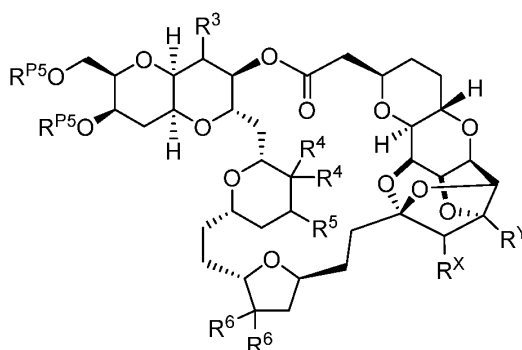
or a salt thereof, wherein:

R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group; and

optionally wherein R^{P5} and R^{P7} are joined with the intervening atoms to form optionally substituted heterocyclyl.

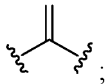
[00513] Provided herein are compounds of Formula **(R-4-11A)**:



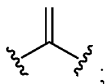
(R-4-11A),

and salts thereof, wherein:

R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, optionally substituted alkyl,

or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, optionally substituted alkyl,

or two R^6 groups are taken together to form: ;

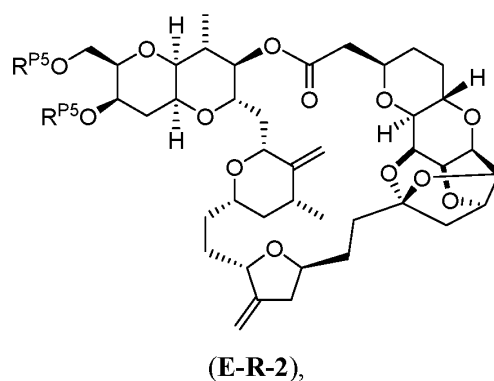
each instance of R^{P5} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

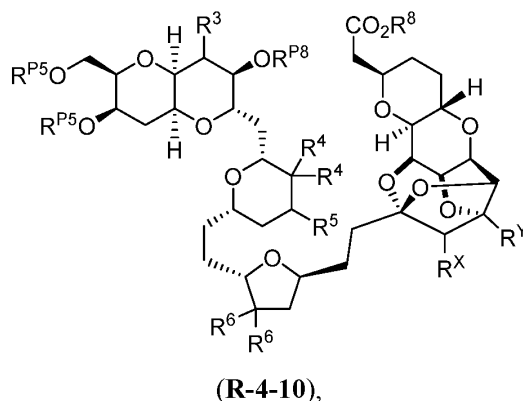
[00514] In certain embodiments, the compound is of the Formula (**E-R-2**):



or a salt thereof, wherein:

each instance of R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form optionally substituted heterocyclyl ring.

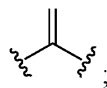
[00515] Provided herein are compounds of Formula (R-4-10):



and salts thereof, wherein:

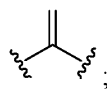
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form:



each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form:



each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring;

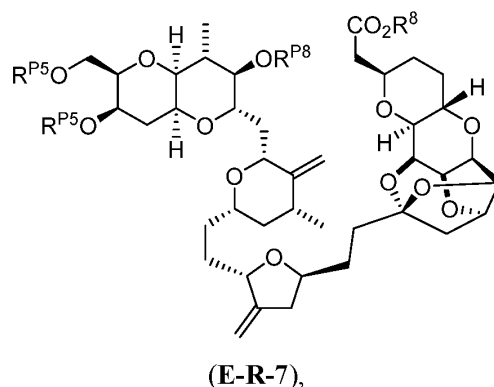
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00516] In certain embodiments, the compound is of the Formula (E-R-7):

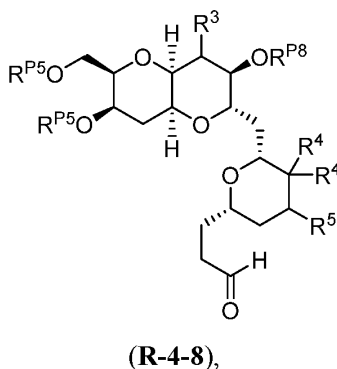


or a salt thereof, wherein:

each instance of R^{P5} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00517] Provided herein are compounds of Formula (R-4-8):



and salts thereof, wherein:

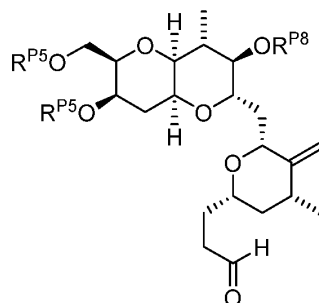
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl; each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ; and

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups

are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00518] In certain embodiments, the compound is of the Formula (E-R-4):

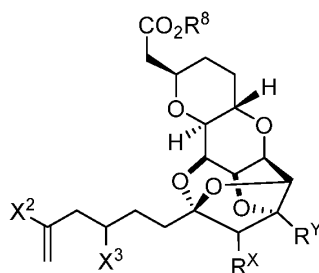


(E-R-4),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00519] Provided herein are compounds of Formula (R-4-9):



(R-4-9),

and salts thereof, wherein:

X^3 and X^2 are each independently halogen or a leaving group;

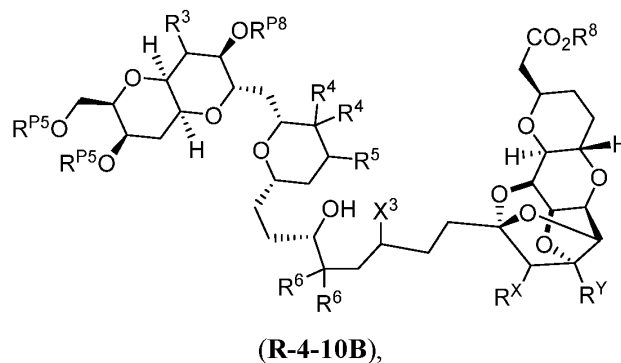
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

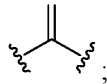
[00520] Provided herein are compounds of Formula (R-4-10B):



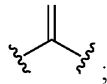
and salts thereof, wherein:

X^3 is halogen or a leaving group;

R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;
each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^6 groups are taken together to form: ;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring;

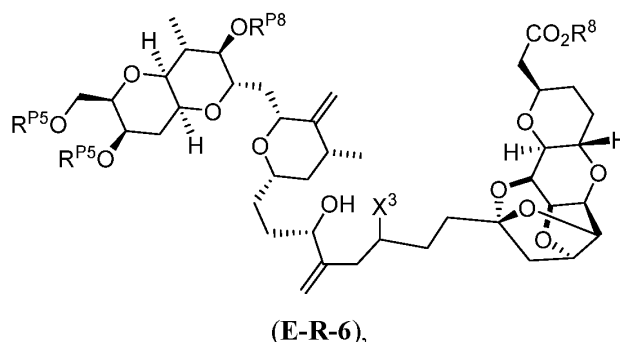
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

[00521] In certain embodiments, the compound is of the Formula (E-R-6):



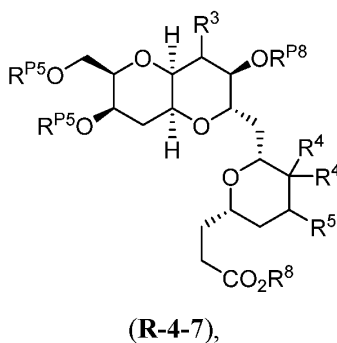
or a salt thereof, wherein:

X^3 is halogen or a leaving group;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclcyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclcyl, optionally substituted aryl, optionally substituted heterocyclcyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

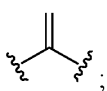
[00522] Provided herein are compounds of Formula (R-4-7):



and salts thereof, wherein:

R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, optionally substituted alkyl,

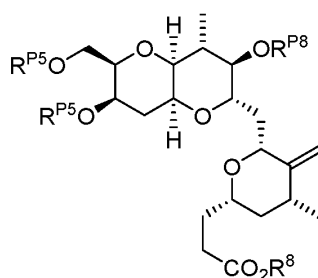
or two R^4 groups are taken together to form: ;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups

are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00523] In certain embodiments, the compound is of the Formula (E-R-8):



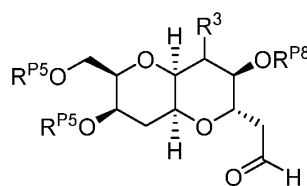
(E-R-8),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00524] Provided herein are compounds of Formula (R-4-5B):



(R-4-5B),

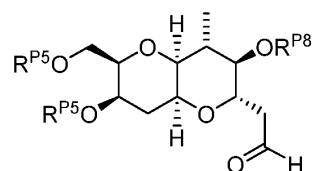
and salts thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P5} and R^{P8} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00525] In certain embodiments, the compound is of the Formula (E-R-9):

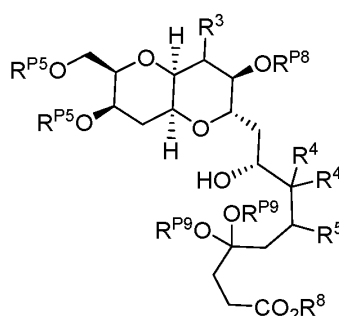


(E-R-9),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

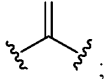
[00526] Provided herein are compounds of Formula (R-4-7A):



(R-4-7A),

and salts thereof; wherein:

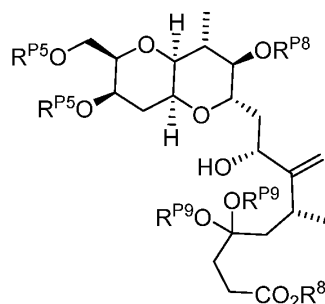
R^3 and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl; each instance of R^4 is independently hydrogen, halogen, or optionally substituted

alkyl, or two R^4 groups are taken together to form: ;

each instance of R^{P5} , R^{P8} , and R^{P9} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and optionally wherein two R^{P9} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00527] In certain embodiments, the compound is of the Formula (E-R-11):

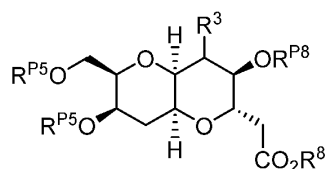


(E-R-11),

or a salt thereof, wherein:

each instance of R^{P5} , R^{P8} , and R^{P9} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; optionally wherein two R^{P9} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00528] Provided herein are compounds of Formula (R-4-5A):



(R-4-5A),

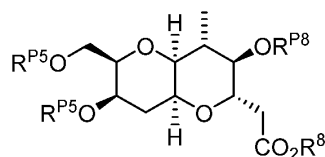
and salts thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00529] In certain embodiments, the compound is of Formula (E-R-15):



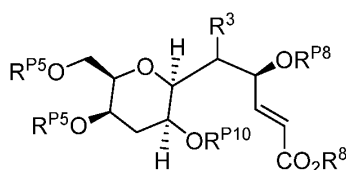
(E-R-15),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00530] Provided herein are compounds of Formula (R-4-4):



(R-4-4),

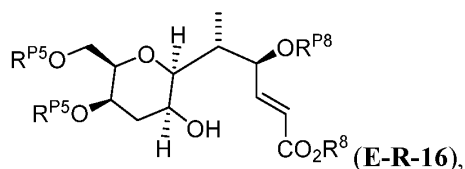
and salts thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P5} , R^{P8} , and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00531] In certain embodiments, the compound is of Formula (E-R-16):



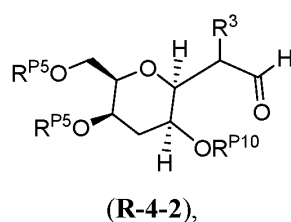
(E-R-16),

or a salt thereof, wherein:

each instance of R^{P5} and R^{P8} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00532] Provided herein are compounds of Formula (R-4-2):

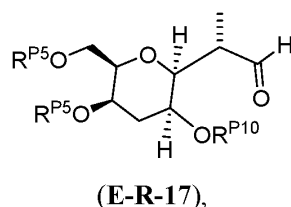


and salts thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl; and

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

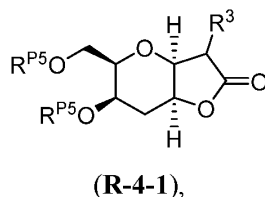
[00533] In certain embodiments, the compound is of Formula (E-R-17):



or a salt thereof, wherein:

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00534] Provided herein are compounds of Formula (R-4-1):

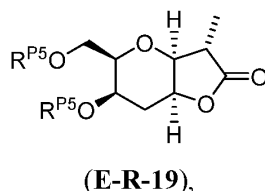


and salts thereof, wherein:

R^3 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P5} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

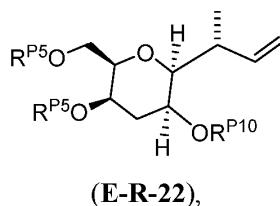
[00535] In certain embodiments, the compound is of Formula (E-R-19):



or a salt thereof, wherein:

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

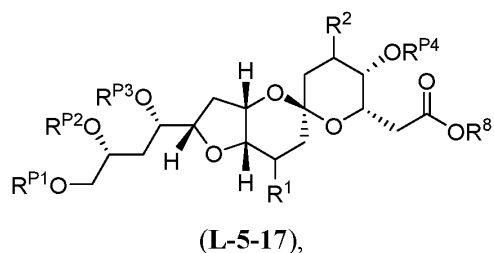
[00536] Provided herein are compounds of the compound of Formula (E-R-22):



and salts thereof, wherein:

each instance of R^{P5} and R^{P10} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} groups are joined together with the intervening atoms to form an optionally substituted heterocyclyl ring.

[00537] Provided herein are compounds of Formula (L-5-17):



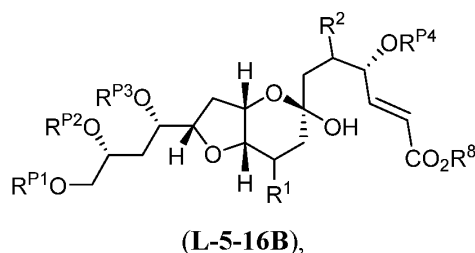
and salts thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P2} , R^{P3} , and R^{P4} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00538] Also provided herein are compounds of Formula (L-5-16B):



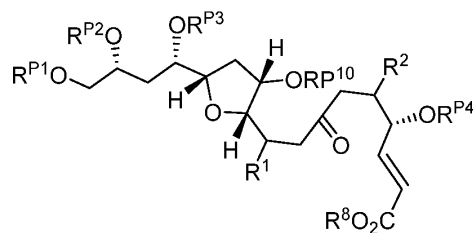
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P2} , R^{P3} , and R^{P4} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00539] Provided herein are compounds of Formula (L-5-16A):



(L-5-16A),

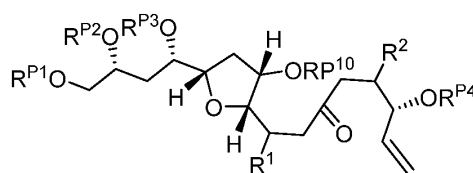
and salts thereof, wherein:

R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1}, R^{P2}, R^{P3}, R^{P4}, and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R⁸ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00540] Provided herein are compounds of Formula (L-5-15):



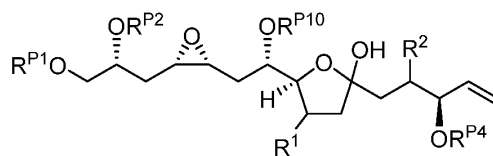
(L-5-15),

and salts thereof, wherein:

R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1}, R^{P2}, R^{P3}, R^{P4}, and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00541] Also provided herein are compounds of Formula (L-5-14):



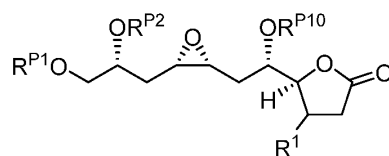
(L-5-14),

and salts thereof, in the presence of an acid, wherein:

R¹ and R² are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1}, R^{P2}, R^{P4}, and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00542] Provided herein are compounds of Formula (L-5-12):



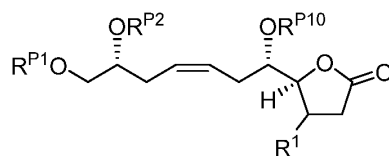
(L-5-12),

and salts thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P2} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00543] Provided herein are compounds of Formula (L-5-11):



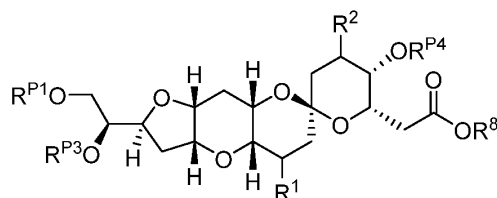
(L-5-11),

and salts thereof, wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P2} , and R^{P10} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00544] Also provided herein are compounds of Formula (L-5-26):



(L-5-26),

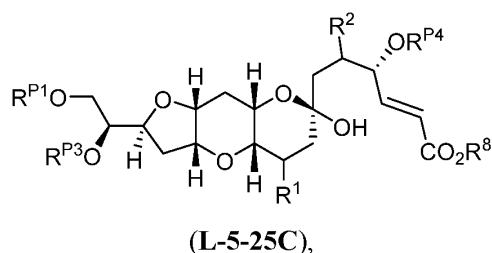
and salts thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00545] Also provided herein are compounds of Formula (L-5-25C):



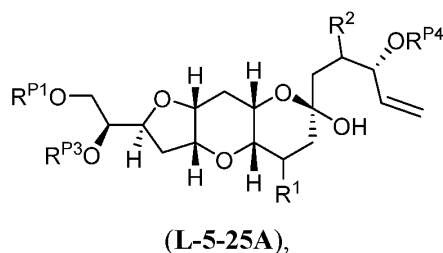
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00546] Also provided herein are compounds of Formula (L-5-25A):

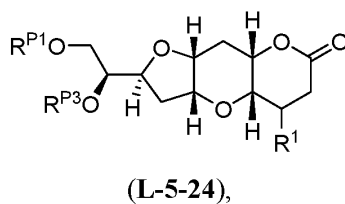


and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P4} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00547] Provided herein are compounds of Formula (L-5-24):

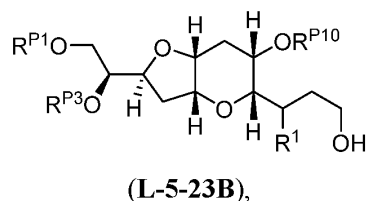


and salts thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} and R^{P3} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00548] Provided herein are compounds of Formula (L-5-23B):

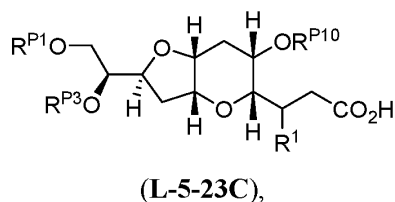


and salts thereof, wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00549] Also provided herein are compound of Formula (L-5-23C):

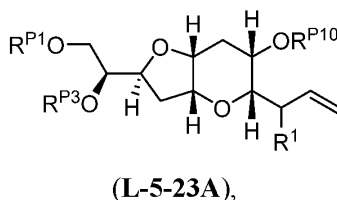


and salts thereof, wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00550] Provided herein are compounds of Formula (L-5-23A):

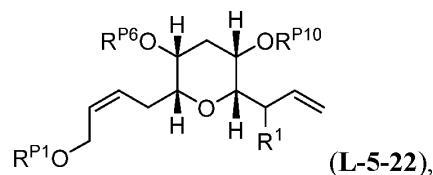


and salts thereof, wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00551] Provided herein are compounds of Formula (L-5-22):

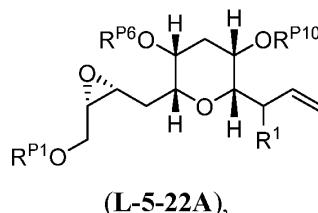


and salts thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00552] Also provided herein are compounds of Formula (L-5-22A):

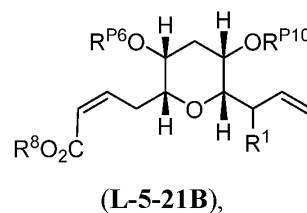


and salt thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P3} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00553] Provided herein are compounds of Formula (L-5-21B):



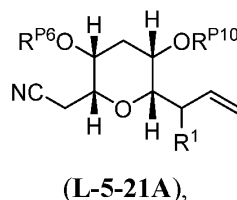
and salts thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P6} , and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00554] Provided herein are compounds of Formula (L-5-21A):

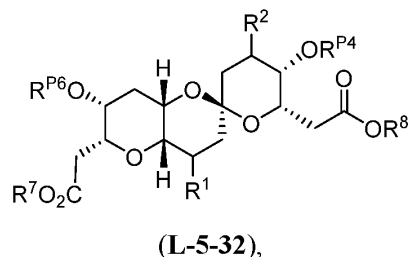


and salts thereof; wherein:

R^1 is hydrogen, halogen, or optionally substituted alkyl; and

R^{P6} and R^{P10} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00555] Provided herein are compounds of Formula (L-5-32):



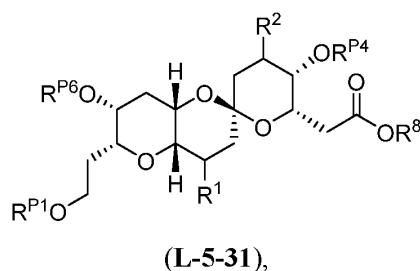
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^7 and R^8 are independently hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00556] Also provided herein are compounds of Formula (L-5-31):



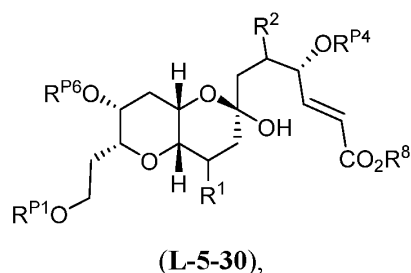
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P4} , and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00557] Also provided herein are compounds of Formula (L-5-32A):



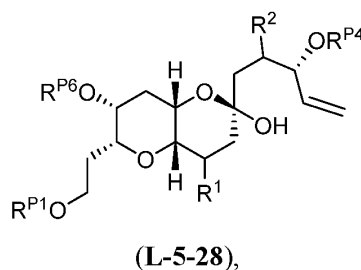
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

R^{P1} , R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00558] Also provided herein are compounds of Formula (L-5-28):

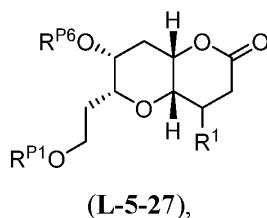


and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and

R^{P1} , R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

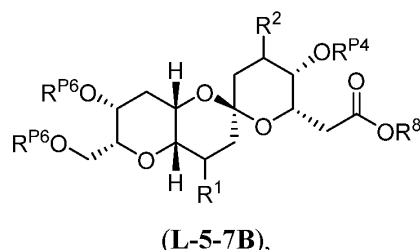
[00559] Provided herein are compounds of Formula (L-5-27):



and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group.

[00560] Provided herein are compounds of Formula (L-5-7B):

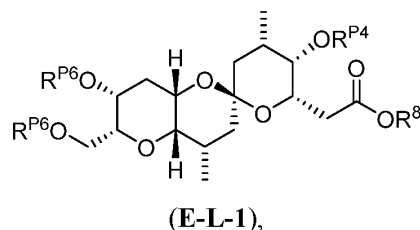


and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00561] In certain embodiments, the compound is of Formula (E-L-1):

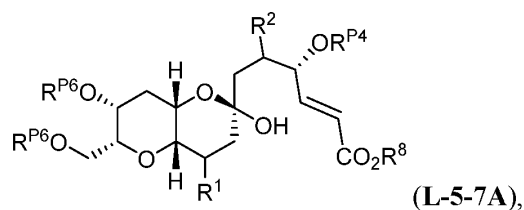


or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl; each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00562] Also provided herein are compounds of Formula (L-5-7A):



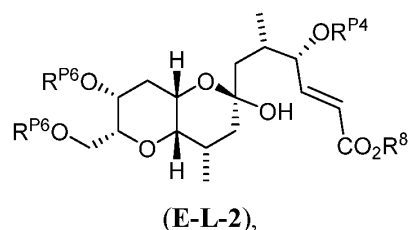
and salts thereof; wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00563] In certain embodiments, the compound is of Formula (E-L-2):

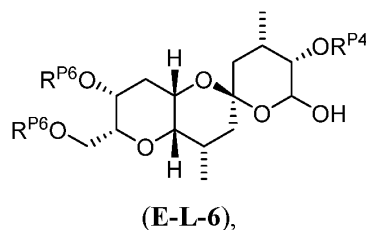


or a salt thereof; wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

[00564] Provided herein are compounds of Formula (E-L-6):

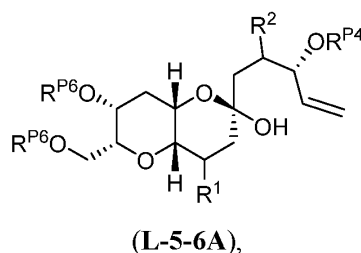


or a salt thereof; wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

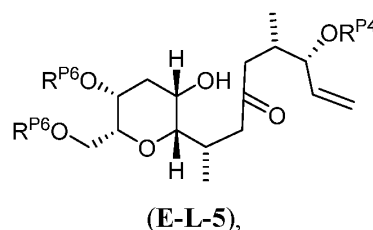
[00565] Provided herein are compounds of Formula (L-5-6A):



and salts thereof, wherein:

R^1 and R^2 are independently hydrogen, halogen, or optionally substituted alkyl; and each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

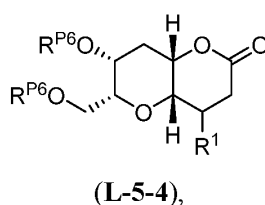
[00566] In certain embodiments, the compound is of Formula (E-L-5):



or a salt thereof, wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

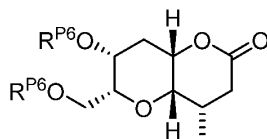
[00567] Also provided herein are compounds of Formula (L-5-4):



and salts thereof, wherein:

R^1 is independently hydrogen, halogen, or optionally substituted alkyl; and
 each R^{P6} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00568] In certain embodiments, the compound is of Formula (E-L-7):



(E-L-7),

or a salt thereof, wherein:

each instance of R^{P4} and R^{P6} are independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

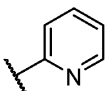
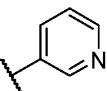
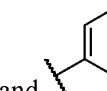
Group R^L , X^L

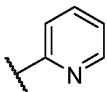
[00569] In certain embodiments, R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl. In certain embodiments, R^L is optionally substituted sulfonyl. In certain embodiments, R^L is optionally substituted sulfinyl. In certain embodiments, R^L is optionally substituted phosphoryl. In certain embodiments, R^L is optionally substituted acyl. In certain embodiments, R^L is $-\text{SO}_2$ -alkyl. In certain embodiments, R^L is mesyl ($-\text{SO}_2\text{CH}_3$; "Ms"). In certain embodiments, R^L is $-\text{SO}_2$ -aryl. In certain embodiments, R^L is $-\text{SO}_2\text{Ph}$. In certain embodiments, R^L is *p*-toluenesulfonyl ($-\text{SO}_2\text{C}_6\text{H}_4\text{p}-\text{CH}_3$; "tosyl" or "Ts"). In certain embodiments, R^L is trifluoromethanesulfonyl ($-\text{SO}_2\text{CF}_3$; "triflyl" or "Tf"). In certain embodiments, R^L is *p*-bromobenzenesulfonyl ($-\text{SO}_2\text{C}_6\text{H}_4\text{p}-\text{Br}$; "brosyl" or "Bs"). In certain embodiments, R^L is nonafluorobutanesulfonyl ($-\text{OSO}_2(\text{CF}_2)_3\text{CF}_3$; "Nf"). In certain embodiments, R^L is 2- or 4-nitrobenzenesulfonyl ($-\text{SO}_2\text{C}_6\text{H}_4\text{p}-\text{NO}_2$ or $-\text{SO}_2\text{C}_6\text{H}_4\text{o}-\text{NO}_2$; "nosyl" or "Ns"). In certain embodiments, R^L is 2,2,2-trifluoroethyl-1-sulfonyl. In certain embodiments, R^L is 5-(dimethylamino)naphthalene-1-sulfonyl ("dansyl" or "Ds").

[00570] As defined herein, X^L is halogen or a leaving group. As defined herein, in certain embodiments, X^L is halogen. In certain embodiments, X^L is $-\text{Cl}$. In certain embodiments, X^L is $-\text{Br}$. In certain embodiments, X^L is $-\text{I}$.

Group R^S

[00571] As defined herein, R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl. In certain embodiments, R^S is optionally substituted alkyl. In certain embodiments, R^S is optionally substituted C₁₋₆ alkyl. In certain embodiments, R^S is unsubstituted C₁₋₆ alkyl. In certain embodiments, R^S is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^S is optionally substituted carbocyclyl. In certain embodiments, R^S is optionally substituted aryl. In certain embodiments, R^S is optionally substituted heterocyclyl. In certain embodiments, R^S is optionally substituted heteroaryl. In certain embodiments, R^S is optionally substituted 6-membered heteroaryl. In certain embodiments, R^S is optionally substituted 6-membered heteroaryl comprising 1, 2, or 3 nitrogen atoms. In certain embodiments, R^S is optionally substituted pyridyl. In certain embodiments, R^S is unsubstituted pyridyl (Py). In certain embodiments, R^S is optionally substituted 2-pyridyl. In certain embodiments, R^S is unsubstituted 2-pyridyl (2-Py). In certain embodiments, R^S is

selected from the group consisting of: , , and . In certain

embodiments, R^S is  (abbreviated herein as “2-Py” or “Py”).

Groups X¹, X², X³, and X⁴

[00572] As defined herein, X¹ is halogen or a leaving group. In certain embodiments, X¹ is a halogen. In certain embodiments, X¹ is –Cl (*i.e.*, chloride). In certain embodiments, X¹ is –Br (*i.e.*, bromide). In certain embodiments, X¹ is –I (*i.e.*, iodide). In certain embodiments, X¹ is –F (*i.e.*, fluoride). In certain embodiments, X¹ is a leaving group.

[00573] As defined herein, X² is halogen or a leaving group. In certain embodiments, X² is a halogen. In certain embodiments, X² is –Cl. In certain embodiments, X² is –Br. In certain embodiments, X² is –I. In certain embodiments, X² is –F. In certain embodiments, X² is a leaving group.

[00574] As defined herein, X³ is halogen or a leaving group. In certain embodiments, X³ is a halogen. In certain embodiments, X³ is –Cl. In certain embodiments, X³ is –Br. In certain embodiments, X³ is –I. In certain embodiments, X³ is –F. In certain embodiments, X³ is a leaving group.

[00575] As defined herein, X^4 is halogen or a leaving group. In certain embodiments, X^4 is a halogen. In certain embodiments, X^4 is $-Cl$. In certain embodiments, X^4 is $-Br$. In certain embodiments, X^4 is $-I$. In certain embodiments, X^4 is $-F$. In certain embodiments, X^4 is a leaving group.

Groups R^1 , R^2 , R^3 , R^4 , R^5 , and R^6

[00576] As defined herein, R^1 is hydrogen, halogen, or optionally substituted alkyl. In certain embodiments, R^1 is hydrogen. In certain embodiments, R^1 is halogen. In certain embodiments, R^1 is optionally substituted alkyl. In certain embodiments, R^1 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^1 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^1 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^1 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^1 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^1 is methyl.

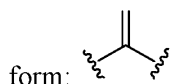
[00577] As defined herein, R^2 is hydrogen, halogen, or optionally substituted alkyl. In certain embodiments, R^2 is hydrogen. In certain embodiments, R^2 is halogen. In certain embodiments, R^2 is optionally substituted alkyl. In certain embodiments, R^2 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^2 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^2 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^2 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^2 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^2 is methyl.

[00578] As defined herein, R^3 is hydrogen, halogen, or optionally substituted alkyl. In certain embodiments, R^3 is hydrogen. In certain embodiments, R^3 is halogen. In certain embodiments, R^3 is optionally substituted alkyl. In certain embodiments, R^3 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^3 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^3 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^3 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^3 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^3 is methyl.

[00579] As defined herein, each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl; and optionally two R^4 groups are taken together to form:



. In certain embodiments, R^4 is hydrogen. In certain embodiments, R^4 is halogen. In certain embodiments, R^4 is optionally substituted alkyl. In certain embodiments, R^4 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^4 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^4 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^4 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^4 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^4 is methyl. In certain embodiments, two R^4 groups are taken together to

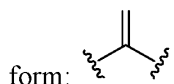


[00580] As defined herein, R^5 is hydrogen, halogen, or optionally substituted alkyl. In certain embodiments, R^5 is hydrogen. In certain embodiments, R^5 is halogen. In certain embodiments, R^5 is optionally substituted alkyl. In certain embodiments, R^5 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^5 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^5 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^5 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^5 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^5 is methyl.

[00581] As defined herein, each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl; and optionally two R^6 groups are taken together to form:



. In certain embodiments, R^6 is hydrogen. In certain embodiments, R^6 is halogen. In certain embodiments, R^6 is optionally substituted alkyl. In certain embodiments, R^6 is optionally substituted C_{1-6} alkyl. In certain embodiments, R^6 is unsubstituted C_{1-6} alkyl. In certain embodiments, R^6 is optionally substituted C_{1-3} alkyl. In certain embodiments, R^6 is unsubstituted C_{1-3} alkyl. In certain embodiments, R^6 is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^6 is methyl. In certain embodiments, two R^6 groups are taken together to



Groups R⁷ and R⁸

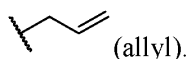
[00582] As defined herein, R⁷ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R⁷ is hydrogen. In certain embodiments, R⁷ is optionally substituted alkyl. In certain embodiments, R⁷ is optionally substituted C₁₋₆ alkyl. In certain embodiments, R⁷ is unsubstituted C₁₋₆ alkyl. In certain embodiments, R⁷ is optionally substituted C₁₋₃ alkyl. In certain embodiments, R⁷ is unsubstituted C₁₋₃ alkyl. In certain embodiments, R⁷ is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R⁷ is methyl. In certain embodiments, R⁷ is ethyl. In certain embodiments, R⁷ is optionally substituted carbocyclyl. In certain embodiments, R⁷ is optionally substituted aryl. In certain embodiments, R⁷ is optionally substituted heterocyclyl. In certain embodiments, R⁷ is optionally substituted heteroaryl. In certain embodiments, R⁷ is optionally substituted acyl. In certain embodiments, R⁷ is an oxygen protecting group. In certain embodiments, R⁷ is an optionally substituted benzyl protecting group. In certain embodiments, R⁷ is benzyl (–CH₂Ph; “Bn”).

[00583] As defined herein, R⁸ is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R⁸ is hydrogen. In certain embodiments, R⁸ is optionally substituted alkyl. In certain embodiments, R⁸ is optionally substituted C₁₋₆ alkyl. In certain embodiments, R⁸ is unsubstituted C₁₋₆ alkyl. In certain embodiments, R⁸ is optionally substituted C₁₋₃ alkyl. In certain embodiments, R⁸ is unsubstituted C₁₋₃ alkyl. In certain embodiments, R⁸ is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R⁸ is methyl. In certain embodiments, R⁸ is ethyl. In certain embodiments, R⁸ is optionally substituted carbocyclyl. In certain embodiments, R⁸ is optionally substituted aryl. In certain embodiments, R⁸ is optionally substituted heterocyclyl. In certain embodiments, R⁸ is optionally substituted heteroaryl. In certain embodiments, R⁸ is optionally substituted acyl. In certain embodiments, R⁸ is an oxygen protecting group. In certain embodiments, R⁸ is an optionally substituted benzyl protecting group. In certain embodiments, R⁸ is benzyl (–CH₂Ph; “Bn”).

Groups R^X and R^Y

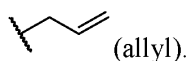
[00584] As defined herein, R^X is hydrogen or $-OR^{Xa}$. In certain embodiments, R^X is hydrogen. In certain embodiments, R^X is $-OR^{Xa}$.

[00585] As generally defined herein, R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{Xa} is hydrogen. In certain embodiments, R^{Xa} is optionally substituted alkyl. In certain embodiments, R^{Xa} is optionally substituted acyl. In certain embodiments, R^{Xa} is or an oxygen protecting group. In certain embodiments, R^{Xa} is optionally substituted allyl. In certain embodiments, R^{Xa} is

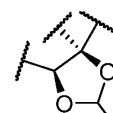


[00586] As defined herein, R^Y is hydrogen or $-OR^{Ya}$. In certain embodiments, R^Y is hydrogen. In certain embodiments, R^Y is $-OR^{Ya}$.

[00587] As generally defined herein, R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{Ya} is hydrogen. In certain embodiments, R^{Ya} is optionally substituted alkyl. In certain embodiments, R^{Ya} is optionally substituted acyl. In certain embodiments, R^{Ya} is or an oxygen protecting group. In certain embodiments, R^{Ya} is optionally substituted allyl. In certain embodiments, R^{Ya} is

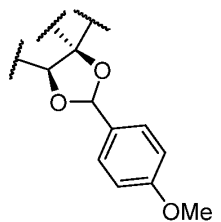


[00588] In certain embodiments, R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl. In certain embodiments, R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted 5-membered heterocyclyl. In certain embodiments, R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted 1,3-dioxolane ring. In certain embodiments, R^{Xa} and R^{Ya}



are joined together with their intervening atoms to form the following:

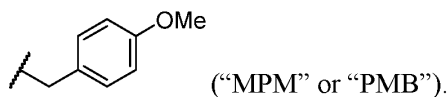
In certain embodiments, R^{Xa} and R^{Ya} are joined together with their intervening atoms to form



the following:

Groups R^{P1} , R^{P2} , R^{P3} , R^{P4} , R^{P5} , R^{P6} , R^{P7} , R^{P8} , R^{P9} , and R^{P10}

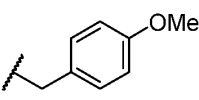
[00589] As defined herein, R^{P1} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P1} is hydrogen. In certain embodiments, R^{P1} is optionally substituted alkyl. In certain embodiments, R^{P1} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P1} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P1} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P1} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P1} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P1} is optionally substituted acyl. In certain embodiments, R^{P1} is an oxygen protecting group. In certain embodiments, R^{P1} is optionally substituted allyl. In certain embodiments, R^{P1} is allyl. In certain embodiments, R^{P1} is optionally substituted silyl. In certain embodiments, R^{P1} is trialkylsilyl. In certain embodiments, R^{P1} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P1} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P1} is *tert*-butyl dimethylsilyl ($-\text{Si}(t\text{-Bu})\text{Me}_2$; “TBS”). In certain embodiments, R^{P1} is *tert*-butyl diphenylsilyl ($-\text{Si}(t\text{-Bu})\text{Ph}_2$; “TBDPS”). In certain embodiments, R^{P1} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P1} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P1} is a methoxybenzyl protecting group. In certain embodiments, R^{P1} is *para*-methoxybenzyl:



[00590] In certain embodiments, R^{P1} and R^{P2} are joined with the intervening atoms to form optionally substituted heterocycl.

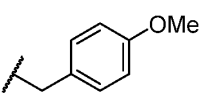
[00591] As defined herein, R^{P2} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P2} is hydrogen. In certain embodiments, R^{P2} is optionally substituted alkyl. In certain embodiments, R^{P2} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P2} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P2} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P2} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P2} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P2} is optionally substituted acyl. In certain embodiments, R^{P2} is an oxygen protecting group. In certain embodiments, R^{P2} is optionally substituted allyl. In certain embodiments, R^{P2} is allyl. In certain embodiments, R^{P2} is optionally substituted silyl. In certain embodiments, R^{P2} is trialkylsilyl. In certain embodiments, R^{P2} is triethylsilyl ($-\text{SiEt}_3$;

“TES”). In certain embodiments, R^{P2} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P2} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P2} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P2} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P2} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P2} is a methoxybenzyl protecting group. In certain

embodiments, R^{P2} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

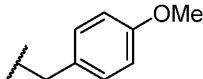
[00592] In certain embodiments, R^{P3} and R^{P3} are joined with the intervening atoms to form optionally substituted heterocyclyl.

[00593] As defined herein, R^{P3} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P3} is hydrogen. In certain embodiments, R^{P3} is optionally substituted alkyl. In certain embodiments, R^{P3} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P3} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P3} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P3} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P3} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P3} is optionally substituted acyl. In certain embodiments, R^{P3} is an oxygen protecting group. In certain embodiments, R^{P3} is optionally substituted allyl. In certain embodiments, R^{P3} is allyl. In certain embodiments, R^{P3} is optionally substituted silyl. In certain embodiments, R^{P3} is trialkylsilyl. In certain embodiments, R^{P3} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P3} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P3} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P3} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P3} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P3} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P3} is a methoxybenzyl protecting group. In certain

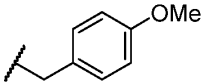
embodiments, R^{P3} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

[00594] As defined herein, R^{P4} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P4} is hydrogen. In certain embodiments, R^{P4} is optionally substituted alkyl. In certain embodiments, R^{P4} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P4} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P4} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P4} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P4} is selected from the group consisting of

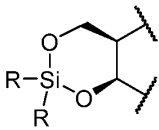
methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P4} is optionally substituted acyl. In certain embodiments, R^{P4} is an oxygen protecting group. In certain embodiments, R^{P4} is optionally substituted allyl. In certain embodiments, R^{P4} is allyl. In certain embodiments, R^{P4} is optionally substituted silyl. In certain embodiments, R^{P4} is trialkylsilyl. In certain embodiments, R^{P4} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P4} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P4} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P4} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P4} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P4} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P4} is a methoxybenzyl protecting group. In certain

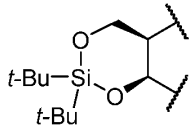
embodiments, R^{P4} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

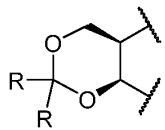
[00595] As defined herein, R^{P5} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P5} are joined with the intervening atoms to form optionally substituted heterocyclyl. In certain embodiments, R^{P5} is hydrogen. In certain embodiments, R^{P5} is optionally substituted alkyl. In certain embodiments, R^{P5} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P5} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P5} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P5} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P5} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P5} is optionally substituted acyl. In certain embodiments, R^{P5} is an oxygen protecting group. In certain embodiments, R^{P5} is optionally substituted allyl. In certain embodiments, R^{P5} is allyl. In certain embodiments, R^{P5} is optionally substituted silyl. In certain embodiments, R^{P5} is trialkylsilyl. In certain embodiments, R^{P5} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P5} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P5} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P5} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P5} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P5} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P5} is a methoxybenzyl protecting group. In certain embodiments, R^{P5} is *para*-methoxybenzyl:

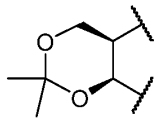
 (“MPM” or “PMB”). In certain embodiments, two R^{P5} are joined with the intervening atoms to form optionally substituted heterocyclyl. In certain embodiments, two

R^{P5} are joined with the intervening atoms to form optionally substituted six-membered heterocyclyl. In certain embodiments, two R^{P5} are joined with the intervening atoms to form a

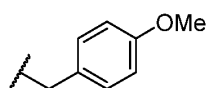
ring of the formula: . In certain embodiments, two R^{P5} are joined with the

intervening atoms to form a ring of the formula: . In certain embodiments,

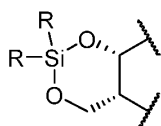
two R^{P5} are joined with the intervening atoms to form a ring of the formula: . In certain embodiments, two R^{P5} are joined with the intervening atoms to form a ring of the

formula: .

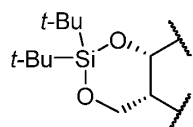
[00596] As defined herein, R^{P6} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl. In certain embodiments, R^{P6} is hydrogen. In certain embodiments, R^{P6} is optionally substituted alkyl. In certain embodiments, R^{P6} is optionally substituted C₁₋₆ alkyl. In certain embodiments, R^{P6} is unsubstituted C₁₋₆ alkyl. In certain embodiments, R^{P6} is optionally substituted C₁₋₃ alkyl. In certain embodiments, R^{P6} is unsubstituted C₁₋₃ alkyl. In certain embodiments, R^{P6} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P6} is optionally substituted acyl. In certain embodiments, R^{P6} is an oxygen protecting group. In certain embodiments, R^{P6} is optionally substituted allyl. In certain embodiments, R^{P6} is allyl. In certain embodiments, R^{P6} is optionally substituted silyl. In certain embodiments, R^{P6} is trialkylsilyl. In certain embodiments, R^{P6} is triethylsilyl (–SiEt₃; “TES”). In certain embodiments, R^{P6} is trimethylsilyl (–SiMe₃; “TMS”). In certain embodiments, R^{P6} is *tert*-butyl dimethylsilyl (–Si(*t*-Bu)Me₂; “TBS”). In certain embodiments, R^{P6} is *tert*-butyl diphenylsilyl (–Si(*t*-Bu)Ph₂; “TBDPS”). In certain embodiments, R^{P6} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P6} is benzyl (–CH₂Ph; “Bn”). In certain embodiments, R^{P6} is a methoxybenzyl protecting group. In certain embodiments, R^{P6} is *para*-methoxybenzyl:



(“MPM” or “PMB”). In certain embodiments, two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl. In certain embodiments, two R^{P6} are joined with the intervening atoms to form optionally substituted six-membered heterocyclyl. In certain embodiments, two R^{P6} are joined with the intervening atoms to form a



ring of the formula: . In certain embodiments, two R^{P6} are joined with the

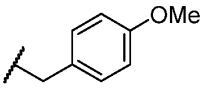


intervening atoms to form a ring of the formula:

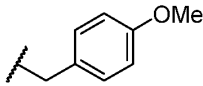
[00597] As defined herein, in certain embodiments, R^{P7} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In other embodiments, R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P7} is optionally substituted sulfonyl. In certain embodiments, R^{P7} is mesyl ($-\text{SO}_2\text{CH}_3$; “Ms”). In certain embodiments, R^{P7} is tosyl ($\text{C}-\text{SO}_2\text{C}_6\text{H}_4\text{p}-\text{CH}_3$; “Ts”). In certain embodiments, R^{P7} is triflyl ($-\text{SO}_2\text{CF}_3$; “Tf”). In certain embodiments, R^{P7} is optionally substituted sulfinyl. In certain embodiments, R^{P7} is optionally substituted phosphoryl. In certain embodiments, R^{P7} is optionally substituted acyl.

[00598] In certain embodiments, R^{P7} is hydrogen. In certain embodiments, R^{P7} is optionally substituted alkyl. In certain embodiments, R^{P7} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P7} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P7} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P7} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P7} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P7} is optionally substituted acyl. In certain embodiments, R^{P7} is an oxygen protecting group. In certain embodiments, R^{P7} is optionally substituted allyl. In certain embodiments, R^{P7} is allyl. In certain embodiments, R^{P7} is optionally substituted silyl. In certain embodiments, R^{P7} is trialkylsilyl. In certain embodiments, R^{P7} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P7} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P7} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P7} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P7} is an optionally

substituted benzyl protecting group. In certain embodiments, R^{P7} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P7} is a methoxybenzyl protecting group. In certain embodiments,

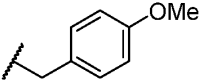
R^{P7} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

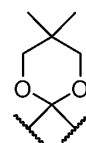
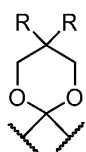
[00599] As defined herein, R^{P8} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P8} is hydrogen. In certain embodiments, R^{P8} is optionally substituted alkyl. In certain embodiments, R^{P8} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P8} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P8} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P8} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P8} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P8} is optionally substituted acyl. In certain embodiments, R^{P8} is an oxygen protecting group. In certain embodiments, R^{P8} is optionally substituted allyl. In certain embodiments, R^{P8} is allyl. In certain embodiments, R^{P8} is optionally substituted silyl. In certain embodiments, R^{P8} is trialkylsilyl. In certain embodiments, R^{P8} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P8} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P8} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P8} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P8} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P8} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P8} is a methoxybenzyl protecting group. In certain

embodiments, R^{P8} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

[00600] As defined herein, R^{P9} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P9} is hydrogen. In certain embodiments, R^{P9} is optionally substituted alkyl. In certain embodiments, R^{P9} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P9} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P9} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P9} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P9} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P9} is optionally substituted acyl. In certain embodiments, R^{P9} is an oxygen protecting group. In certain embodiments, R^{P9} is optionally substituted allyl. In certain embodiments, R^{P9} is allyl. In certain embodiments, R^{P9} is optionally substituted silyl. In certain embodiments, R^{P9} is trialkylsilyl. In certain embodiments, R^{P9} is triethylsilyl ($-\text{SiEt}_3$;

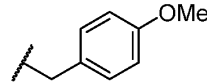
“TES”). In certain embodiments, R^{P9} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P9} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P9} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P9} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P9} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P9} is a methoxybenzyl protecting group. In certain

embodiments, R^{P9} is *para*-methoxybenzyl:  (“MPM” or “PMB”). In certain embodiments, two R^{P9} are joined together with the intervening atoms to form optionally substituted heterocycl. In certain embodiments, two R^{P9} are joined together to form



. In certain embodiments, two R^{P9} are joined together to form

[00601] As defined herein, R^{P10} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group. In certain embodiments, R^{P10} is hydrogen. In certain embodiments, R^{P10} is optionally substituted alkyl. In certain embodiments, R^{P10} is optionally substituted C_{1-6} alkyl. In certain embodiments, R^{P10} is unsubstituted C_{1-6} alkyl. In certain embodiments, R^{P10} is optionally substituted C_{1-3} alkyl. In certain embodiments, R^{P10} is unsubstituted C_{1-3} alkyl. In certain embodiments, R^{P10} is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl. In certain embodiments, R^{P10} is optionally substituted acyl. In certain embodiments, R^{P10} is an oxygen protecting group. In certain embodiments, R^{P10} is optionally substituted allyl. In certain embodiments, R^{P10} is allyl. In certain embodiments, R^{P10} is optionally substituted silyl. In certain embodiments, R^{P10} is trialkylsilyl. In certain embodiments, R^{P10} is triethylsilyl ($-\text{SiEt}_3$; “TES”). In certain embodiments, R^{P10} is trimethylsilyl ($-\text{SiMe}_3$; “TMS”). In certain embodiments, R^{P10} is *tert*-butyl dimethylsilyl ($-\text{Si}t\text{-BuMe}_2$; “TBS”). In certain embodiments, R^{P10} is *tert*-butyl diphenylsilyl ($-\text{Si}t\text{-BuPh}_2$; “TBDPS”). In certain embodiments, R^{P10} is an optionally substituted benzyl protecting group. In certain embodiments, R^{P10} is benzyl ($-\text{CH}_2\text{Ph}$; “Bn”). In certain embodiments, R^{P10} is a methoxybenzyl protecting group. In certain

embodiments, R^{P10} is *para*-methoxybenzyl:  (“MPM” or “PMB”).

Group R

[00602] As generally defined herein, R is hydrogen or optionally substituted alkyl. In certain embodiments, R is hydrogen. In certain embodiments, R is optionally substituted alkyl. In certain embodiments, R is optionally substituted C₁₋₆ alkyl. In certain embodiments, R is unsubstituted C₁₋₆ alkyl. In certain embodiments, R is optionally substituted C₁₋₃ alkyl. In certain embodiments, R is unsubstituted C₁₋₃ alkyl. In certain embodiments, R is selected from the group consisting of methyl, ethyl, *n*-propyl, *iso*-propyl, *n*-butyl, *iso*-butyl, *sec*-butyl, and *tert*-butyl.

EXAMPLES***Zr/Ni-Mediated Ketolization Reactions***

[00603] The structure (*S,S*)-**1-C** can be prepared directly via a coupling of (*S*)-**1-A** with (*S*)-**1-B** (Figure 2A). Although appealing, sequence presents challenges. For example, anion-based ketone syntheses might be problematic, because of the presence of an O-R group at the β- and β'-positions. The "umpolung" concept, represented by dithiane chemistry, is the historical solution for this type of problem (see, *e.g.*, For a review, for example see: Seebach, D. *Angew. Chem. Int. Ed.* 1979, 18, 239; Corey, E. J.; Seebach, D. *Angew. Chem. Int. Ed.*, **1965**, 4, 1077; Seebach, D.; Corey, E. J. *J. Org. Chem.* **1975**, 40, 231). Indeed, dithiane-based ketone synthesis has successfully been applied to a synthesis of complex natural products (For a review, see, *e.g.*, Yus, M.; Najera, C.; Foubelo, F. *Tetrahedron*, **2003**, 59, 6147; Smith, III. A. S.; Adams, C. M.; *Acc. Chem. Rev.* **2004**, 37, 365). Nevertheless, a direct ketone synthesis was developed which can be used in the synthesis of ketones, including complex molecules. The best chance of achieving this goal would be a radical-based, preferably one-pot, ketone synthesis. A related Zn/Pd-mediated one-pot ketone synthesis was reported (see, *e.g.*, Lee, J. H.; Kishi, Y. *J. Am. Chem. Soc.* **2016**, 138, 7178).

[00604] Recently, Weix, Gong, Reisman, and others extensively studied Ni-mediated one-pot ketone synthesis, pioneered by Mukaiyama in 1981 (see, *e.g.*, Onaka, M.; Matsuoka, Y.; Mukaiyama, T. *Chem. Lett.* **1981**, 531; Wotal, A. C.; Weix, D. J. *Org. Lett.* **2012**, 14, 1476; Wotal, A. C.; Ribson, R. D.; Weix, D. J. *Organometallics* **2014**, 33, 5874; Wu, F.; Lu, W.; Qian, Q.; Ren, Q.; Gong, H. *Org. Lett.* **2012**, 14, 3044; Zhao, C.; Jia, X.; Wang, X.; Gong, H. *J. Am. Chem. Soc.* **2014**, 136, 17645 and references cited therein; Cherney, A. H.; Kadunce, N. T.; Reisman, S. E. *J. Am. Chem. Soc.* **2013**, 135, 7442). Among a wide range of substrates reported, one specific example given by Gong and coworkers suggested a possibility that Ni-

mediated one-pot ketone synthesis might meet with our need (*Figure 2B*). The substrates shown in *Figure 2C* were arbitrarily chosen for this study. The arbitrarily chosen substrates were tested under these three conditions, thereby demonstrating the feasibility of proposed coupling, *e.g.*, via Weix and Reisman protocols. At the same time, it became evident that serious improvements were required successfully to use the Ni-mediated one-pot ketone synthesis at a late-stage coupling in a convergent synthesis of complex molecules.

[00605] More than 15 ligands were first tested to solubilize NiCl_2 , thereby showing 4,4'-di-*tert*-butyl-2,2'-bipyridine (dtbbpy) to give the best result. Noteworthy, $\text{NiBr}_2 \cdot (\text{dtbbpy})$ complex gave a better coupling efficiency than $\text{NiCl}_2 \cdot (\text{dtbbpy})$ complex (see, *e.g.*, Lu, Z.; Fu, G. C. *Angew. Chem. Int. Ed.* **2010**, 49, 6676; Serrano, E.; Martin, R. *Angew. Chem. Int. Ed.* **2016**, 55, 11207; Zhang, X.; MacMillan, D. W. C. *J. Am. Chem. Soc.* **2016**, 138, 13862).

[00606] Among the activated forms of carboxylic acid studied, 2-thiopyridine ester, originally reported by Mukaiyama, was found most effective for the coupling. 2-Thiopyridine ester was originally used for their seminal work of macrolactonization by Corey and Nicolaou (see, *e.g.*, Corey, E. J.; Nicolaou, K. C. *J. Am. Chem. Soc.* **1974**, 96, 5614) and by Nicolaou Gerlack and Thalmann (see, *e.g.*, Gerlach, H.; Thalmann, A. *Helv. Chim. Acta* **1974**, 57, 2661). Mn (powder) and Zn (powder) were found to be effective reducing-metals.

[00607] Among many solvent-systems tested, 1,3-dimethyl-2-imidazolidione (DMI) was found best. A 5:1 mixture of DMI and EtOAc was a good solvent system, when a substrate(s) exhibited a poor solubility in DMI. As expected, better coupling yields were obtained at higher concentration, typical concentration being in the range of $C = 0.1 \sim 0.5$ M. While studying additive effects, it was discovered that addition of one equivalent Cp_2ZrCl_2 dramatically enhanced the coupling rate; the coupling completed within minutes to hours with Cp_2ZrCl_2 , compared with overnight to days without Cp_2ZrCl_2 . In addition, by-product formation via a (I $\rightarrow$ SPy)-displacement was eliminated or suppressed by addition of Cp_2ZrCl_2 .

[00608] The observed, dramatic rate-acceleration indicated that Cp_2ZrCl_2 was involved in the rate-limiting step of catalytic reaction. Two different catalytic cycles had been proposed for the Ni-mediated one-pot ketone synthesis, *i.e.*, (1) the catalytic cycle involving a (L)Ni(alkyl)₂ intermediate and (2) the catalytic cycle of sequential reduction. However, in order to explain the observed results, a new mechanism is proposed, consisting of Ni-catalytic cycle, Zr-catalytic cycle, and Zr $\rightarrow$ Ni transmetallation (*Figure 3A*). The Ni-catalytic cycle starts with Ni(II) $\rightarrow$ Ni(0) reduction by Zn, followed by its oxidative addition to 2-

thiopyridine ester, *i.e.*, **2-D** → **2-E** → **2-F**. Because of the strong Zr-SR bond, it is possible that Cp_2ZrCl_2 and/or a Zr-salt could accelerate the step from **2-F** to **2-G**, thereby resulting in the dramatic rate-acceleration. On the other hand, a second catalytic cycle involving Cp_2ZrCl_2 ; Zn-reduction of Cp_2ZrCl_2 to form a low-valent Zr-species could be operative. According to the seminal work by Schwartz, such a low-valent Zr-species readily activates an alkyl iodide, *i.e.*, $\text{Cp}_2\text{ZrCl}_2 \rightarrow \text{Cp}_2\text{Zr} \rightarrow \text{Cp}_2\text{Zr-R}$ (see, *e.g.*, Williams, G. M.; Gell, K. I.; Schwartz, J. *J. Am. Chem. Soc.* **1980**, *102*, 3660; Williams, G. M.; Schwartz, J. *J. Am. Chem. Soc.* **1982**, *104*, 1122). Then, the Ni- and Zr-catalytic cycles are coupled with Zr/Ni-transmetallation, to yield **2-H** (For transmetallation from alkenyl-Zr → alkenyl-Ni, see, *e.g.*, Negishi, E., Van Horn, D. E. *J. Am. Chem. Soc.* **1977**, *99*, 3168; Loots, M. J., Schwartz, J. *J. Am. Chem. Soc.* **1977**, *99*, 8045). Overall, Cp_2ZrCl_2 plays critical dual roles in this scheme. To differentiate the previous Ni-mediated method, this transformation as Zr/Ni-mediated ketone synthesis.

[00609] Related to the mechanism proposed, several commonly known thiol-scavengers were tested, observing only insignificant effect on the acceleration of coupling rate, thereby supporting the proposed dual roles of Cp_2ZrCl_2 . As noted, the dramatic coupling-rate acceleration of coupling-rate by addition of Cp_2ZrCl_2 indicated its involvement in the rate-limiting step. Although there is no experimental support, it is possible that the rate-limiting step is likely **1-F** → **1-G**. Thus, alkyl iodide participates only after the rate-limiting step, which could explain the reason why the Zr/Ni-mediated ketone synthesis is uniquely different from the Zn/Pd- and Fe/Cu-mediated ketone syntheses. As noted, the coupling-rate acceleration by addition of Cp_2ZrCl_2 indicates its involvement in the rate-limiting step. Therefore, the rate-limiting step is likely **1-E** → **1-F**. Thus, alkyl iodide participates only after the rate-limiting step, which could explain the reason why the Zr/Ni-mediated one-pot ketone synthesis is uniquely different from other Zn/Pd- and Fe/Cu-mediated ketone syntheses.

[00610] The behavior of common radical probes were tested (*Figure 3B*). The observation on **4e** showed the radical nature of coupling reaction. On the other hand, **4a~d** gave the normal ketones, thereby suggesting that a radical intermediate was involved only in a very short time-scale (for the reactivity and stability-instability of β -alkoxyalkyl-Zr(IV)-species, see, *e.g.*, Buchwald, S. L.; Nielsen, R. B.; Dewan, J. C. *Organometallics* **1988**, *7*, 2324; Wipf, P.; Smitrovich, J. H. *J. Org. Chem.* **1991**, *56*, 6494).

[00611] In order to establish exemplary optimum conditions, the effect of molar ratio of **1-1** ($X = I$) and **1-2** ($Y = SPy$) on the coupling efficiency were studied under the condition of $NiBr_2(dtbbpy)$ (5 mol%), Cp_2ZrCl_2 (1.0 equiv.), Zn (excess) in DMI (C : 0.5 M) at rt, to give the following results: 89% isolated yield for **1-1:1-2** = 1.5:1.0, 89% for **1-1:1-2** = 1.2:1.0, 85% for **1-1:1-2** = 1.1:1.0, 83% for **1-1:1-2** = 1.0:1.0, 75% for **1-1:1-2** = 0.8:1.0. Considering all of these observations, the conditions were chosen as: $NiBr_2(dtbbpy)$ (5 mol%), Cp_2ZrCl_2 (1 equiv.), Mn or Zn (excess) in DMI or 5:1 DMI-EtOAc (C : 0.5~0.1 M) at ~20 °C, with (1.2:1.0)-molar ratio of nucleophile and electrophile for further studies. However, based on the molecular size and complexity of coupling partners, the molar ratio could accordingly be adjusted without any noticeable drawback. *Figure 4* summarizes the substrates bearing an OR or relevant group at the α -position. The new method gave the expected products in excellent yields.

[00612] *Figure 5A* summarizes further examples. Common protecting groups were tolerated well (**a**, *Figure 5A*). The coupling was effective for mono- and di-methylated substrates at the β -position, as well as mono-methylated substrate at the α -position, but not effective for dimethylated substrate at the α -position or adamantyl substrate (**b**, *Figure 5A*). This method allows one to selectively to activate, and couple, an alkyl iodide over an alkyl bromide or chloride, as well as an aryl bromide (**c**, *Figure 5A*). As mentioned, this reaction exhibited a radical nature, thereby suggesting the possibility that it might be effective for substrates bearing a free hydroxyl and/or acidic group. Indeed, the coupling with these substrates gave the desired products, but further improvements were obviously required for practical uses (**d**, *Figure 5A*).

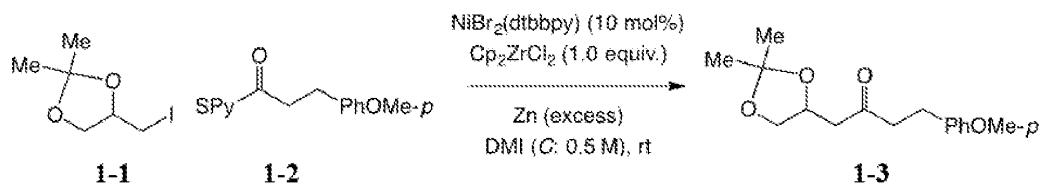
[00613] Finally, in order to demonstrate the applicability of the Zr/Ni-mediated one-pot ketone synthesis to the structure motif given in *Figure 2*, we studied the coupling of (*S*)-**1-11** with (*S*)-**1-12** and (*R*)-**1-12** and obtained expected products (*S,S*)-**1-13** and (*S,R*)-**1-13**, respectively (*Figure 5B*). During the coupling, the stereochemical purity of products, as well as starting materials, could be lost, for example, *via* retro-oxy-Michael/oxy-Michael process. Experimentally, it was found that (*S,S*)-**1-13** and (*S,R*)-**1-13** gave virtually identical ^1H NMR spectra, but exhibited a very similar but distinctly different ^{13}C NMR spectra. With use of ^{13}C NMR spectra, the stereochemical purity of (*S,S*)-**1-13** and (*S,R*)-**1-13** was studied, thereby demonstrating that no stereochemistry scrambling took place in the ketone coupling.

[00614] A new Zr/Ni-mediated one-pot ketone synthesis was reported, where Cp_2ZrCl_2 dramatically accelerated the coupling rate and, at the same time, suppressed by-product

formation via a (I→SPy)-displacement. Unlike Zn/Pd- and Fe/Cu-mediated one-pot ketone syntheses, the new method was found effective for the nucleophiles bearing an OR or relevant group at α -position. A mechanism, consisting of Ni-catalytic cycle, Zr-catalytic cycle, and Zr→Ni transmetallation, was proposed, where Cp_2ZrCl_2 was suggested to play critical dual roles. The newly developed Zr/Ni-mediated method gives a realistic hope of incorporating one-pot ketone at the late-stage in a convergent synthesis of complex molecules.

Experimental Procedures for Ni/Zr-Mediated Ketolization Reactions

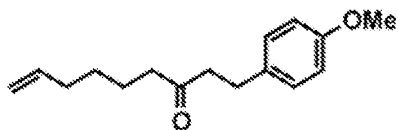
[00615] Solvents and reagents are commercial grade and were used as supplied, unless otherwise noted. Reactions involving air or moisture sensitive reagents or intermediates were performed under an inert atmosphere of nitrogen or argon in glassware that was oven dried. Analytical thin layer chromatography (TLC) was performed with E. Merck pre-coated TLC plates, silica gel 60F-254, layer thickness 0.25 mm. TLC plates were visualized by staining with AMCAN (ammonium molybdate/cerium ammonium nitrate), potassium permanganate, or *p*-anisaldehyde. Flash chromatography separations were performed on E. Merck Silica Gel 60 (40-63 μm), Kanto Chemical Silica Gel 60N (spherical, neutral, 40-50 μm), or Wako Pure Chemical Industry Wakogel 50NH₂ (38-63 μm). Medium pressure column chromatography was performed with YAMAZEN Smart Flash. NMR spectra were recorded on a Varian Inova 600 MHz or Varian Inova 500 MHz. Chemical shifts were reported in parts per million (ppm). The residual solvent peak was used as an internal reference (for ¹H NMR spectra: 7.26 ppm in CDCl_3 , 7.16 ppm in C_6D_6 , 3.31 ppm in CD_3OD , and 5.33 in CD_2Cl_2 ; for ¹³C NMR: 77.0 ppm in CDCl_3 , 128.0 ppm in C_6D_6 , 49.0 ppm in CD_3OD , and 53.8 ppm in CD_2Cl_2). Coupling constants (*J*) are reported in Hz and the splitting abbreviations used are: s for singlet, d for doublet, t for triplet, q for quartet, m for multiplet, and br for broad. Optical rotations were measured at 20 °C using Perkin-Elmer 241 polarimeter. IR spectra were recorded on Bruker Alpha FT-IR spectrometer. Electrospray ionization experiments were performed on Micromass Inc., Platform II Atmospheric Pressure Ionization Mass Spectrometer.

A General Procedure for Ni/Zr-Mediated Coupling Reactions

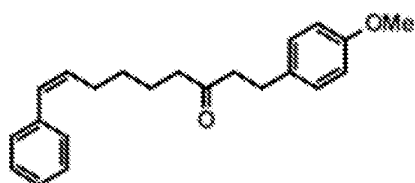
[00616] In a glove box, to a solution of iodide **1-1** (29.1 mg, 0.12 mmol, 1.2 eq.) and thioester **1-2** (27.3 mg, 0.10 mmol, 1.0 eq.) in DMI (0.2 mL, Sigma-aldrich, 99.5%) were added Cp_2ZrCl_2 (29.3 mg, 0.10 mmol, 1.0 eq. Sigma-aldrich, 98%), Zn powder (19.6 mg, 0.3 mmol, 3.0 eq. Sigma-aldrich, used without any activation), and $\text{NiBr}_2 \cdot \text{dtbbpy}$ (4.8 mg, 0.01 mmol, 10 mol%, preparation see page 8) at room temperature. After being stirred at the same temperature for mins to hrs (monitored by TLC), the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel to give **1-3** as colorless oil.

Experimental Procedures for the Reactions Outlined in Figure 3B

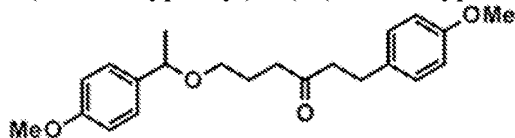
[00617] In a glove box, to a solution of iodide **4a-e** (0.24 mmol, 1.2 eq.) and thioester **1-5** (54.6 mg, 0.20 mmol, 1.0 eq.) in either DMI (0.4 mL, sigma aldrich, 99.5%) or DMI/EtOAc (0.334 mL/0.066mL) were added Cp_2ZrCl_2 (58.5 mg, 0.20 mmol, 1.0 eq. Sigma-aldrich, 98%), Zn powder (39.2 mg, 0.6 mmol, 3.0 eq. Sigma-aldrich, used without any activation), and $\text{NiBr}_2 \cdot \text{dtbbpy}$ (9.7 mg, 0.02 mmol, 10 mol%, preparation see page 8) at room temperature. After being stirred at the same temperature for 10 min to 1 h (monitored by TLC), the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel to give **4a-d**, **S1** as colorless oils. Note: DMI or DMI/EtOAc depending on the solubilities of substrates was used as solvent.

1-(4-methoxyphenyl)non-8-en-3-one (6a)

[00618] 42.1 mg (0.171 mmol, 86%); IR (film) 2930, 2856, 1712, 1612, 1513, 1463, 1300, 1246, 1178, 1109, 1037, 910, 831 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.09 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 5.81-5.74 (m, 1H), 4.99 (dd, J = 17.4, 1.7 Hz, 1H), 4.94 (dd, J = 10.2, 1.7 Hz, 1H), 3.78 (s, 3H), 2.83 (t, J = 7.8 Hz, 2H), 2.69 (t, J = 7.8 Hz, 2H), 2.38 (t, J = 7.2 Hz, 2H), 2.03 (q, J = 7.2 Hz, 2H), 1.60-1.54 (m, 2H), 1.38-1.32 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 210.5, 158.1, 138.6, 133.3, 129.4, 114.8, 114.0, 55.4, 44.7, 43.0, 33.6, 29.1, 28.6, 23.4; HRMS (ESI) m/z calc. for $\text{C}_{16}\text{H}_{23}\text{O}_2$ $[\text{M}+\text{H}]^+$ 247.1708; found 247.1693.

(Z)-1-(4-methoxyphenyl)-9-phenylnon-8-en-3-one (6b)

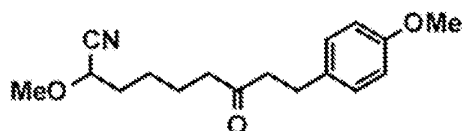
[00619] 59.8 mg, (0.186 mmol, 93%); IR (film) 2931, 2859, 1712, 1612, 1513, 1594, 1463, 1447, 1408, 1373, 1300, 1246, 1178, 1101, 1036, 826, 771, 700 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.32 (t, J = 7.8 Hz, 2H), 7.25 (d, J = 7.8 Hz, 2H), 7.22 (t, J = 7.8 Hz, 1H), 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 6.42 (d, J = 11.4 Hz, 1H), 5.65-5.60 (m, 1H), 3.78 (s, 3H), 2.82 (t, J = 7.2 Hz, 2H), 2.66 (t, J = 8.4 Hz, 2H), 2.35 (t, J = 7.8 Hz, 2H), 2.32 (qd, J = 7.2 Hz, 2.0 Hz, 2H), 1.62-1.56 (m, 2H), 1.44-1.38 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 210.4, 158.1, 137.8, 133.3, 132.6, 129.4, 129.3, 128.9, 128.3, 126.6, 114.0, 55.4, 44.7, 42.9, 29.5, 29.1, 28.4, 23.5; HRMS (ESI) m/z calc. for $\text{C}_{22}\text{H}_{26}\text{NaO}_2$ $[\text{M}+\text{Na}]^+$ 345.1825; found 345.1830.

1-(4-methoxyphenyl)-6-(1-(4-methoxyphenyl)ethoxy)hexan-3-one (6c)

[00620] 61.6 mg, (0.173 mmol, 87%); IR (film) 2953, 2932, 2836, 1712, 1612, 1512, 1464, 1442, 1369, 1301, 1287, 1245, 1177, 1099, 1035, 832 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.20 (d, J = 8.4 Hz, 2H), 7.08 (d, J = 8.4 Hz, 2H), 6.87 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4

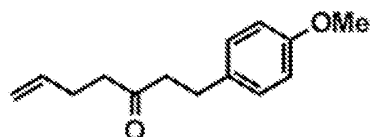
Hz, 2H), 4.29 (q, $J = 6.6$ Hz, 1H), 3.80 (s, 3H), 3.78 (s, 3H), 3.28-3.20 (m, 2H), 2.83-2.78 (m, 2H), 2.70-2.66 (m, 2H), 2.50-2.39 (m, 2H), 1.84-1.77 (m, 2H), 1.39 (d, $J = 6.6$ Hz, 3H); ^{13}C NMR (126 MHz, CDCl_3) $\delta = 210.1, 159.0, 158.0, 136.1, 133.3, 129.3, 127.5, 114.0, 113.9, 77.5, 67.4, 55.4, 44.7, 39.9, 29.0, 24.2, 24.1$; HRMS (ESI) m/z calc. for $\text{C}_{22}\text{H}_{28}\text{NaO}_4$ $[\text{M}+\text{Na}]^+$ 379.1880; found 379.1885.

2-methoxy-9-(4-methoxyphenyl)-7-oxononanenitrile (6d)



[00621] 52.3 mg (0.180 mmol, 90%); IR (film) 2937, 2868, 2834, 1711, 1612, 1513, 1463, 1410, 1372, 1300, 1246, 1179, 1113, 1073, 1035, 829 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) $\delta = 7.09$ (d, $J = 8.4$ Hz, 2H), 6.82 (d, $J = 8.4$ Hz, 2H), 4.02 (t, $J = 7.2$ Hz, 1H), 3.78 (s, 3H), 3.47 (s, 3H), 2.83 (t, $J = 7.8$ Hz, 2H), 2.69 (t, $J = 7.8$ Hz, 2H), 2.39 (t, $J = 7.8$ Hz, 2H), 1.81 (q, $J = 7.2$ Hz, 2H), 1.62-1.56 (m, 2H), 1.47-1.41 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) $\delta = 209.8, 158.1, 133.2, 129.4, 118.1, 114.0, 70.5, 58.1, 55.4, 44.7, 42.7, 33.3, 29.1, 24.4, 23.0$; HRMS (ESI) m/z calc. for $\text{C}_{17}\text{H}_{24}\text{NO}_3$ $[\text{M}+\text{H}]^+$ 290.1751; found 290.1760.

1-(4-methoxyphenyl)hept-6-en-3-one (S1, the product from 4e)



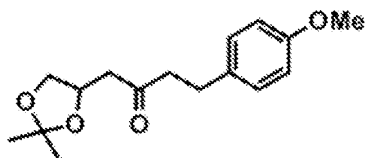
[00622] 33.8 mg (0.155 mmol, 77%); IR (film) 2926, 1753, 1612, 1513, 1442, 1365, 1301, 1246, 1178, 1109, 1036, 911, 829 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) $\delta = 7.09$ (d, $J = 8.5$ Hz, 2H), 6.82 (d, $J = 8.5$ Hz, 2H), 5.83-5.73 (m, 1H), 5.01 (dd, $J = 17.5$ Hz, 1.4 Hz, 1H), 4.97 (dd, $J = 10.0$ Hz, 1.4 Hz, 1H), 3.78 (s, 3H), 2.84 (t, $J = 7.5$ Hz, 2H), 2.70 (t, $J = 7.5$ Hz, 2H), 2.48 (t, $J = 7.5$ Hz, 2H), 2.31 (q, $J = 7.5$ Hz, 2H); ^{13}C NMR (126 MHz, C_6D_6) $\delta = 207.1, 158.6, 137.6, 133.6, 129.6, 115.0, 114.2, 54.7, 44.5, 41.8, 29.1, 28.0$; HRMS (ESI) m/z calc. for $\text{C}_{14}\text{H}_{19}\text{O}_2$ $[\text{M}+\text{H}]^+$ 219.1380; found 219.1374.

Experimental Details for the Reactions Outlined in Figure 4

[00623] In a glove box, to a solution of iodide **7a-m** (0.24 mmol, 1.2 eq.) and thioester **1-5** (54.6 mg, 0.20 mmol, 1.0 eq.) in either DMI (0.4 mL, sigma aldrich, 99.5%) or DMI/EtOAc (0.334 mL/0.066mL) were added Cp_2ZrCl_2 (58.5 mg, 0.20 mmol, 1.0 eq. Sigma-aldrich,

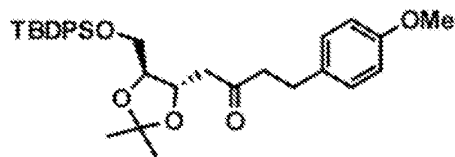
98%), Zn powder (39.2 mg, 0.6 mmol, 3.0 eq. Sigma-aldrich, used without any activation), and $\text{NiBr}_2 \cdot \text{dtbbpy}$ (9.7 mg, 0.02 mmol, 10 mol%, preparation see page 8) at room temperature. After being stirred at the same temperature for 10 min to 2 hr (monitored by TLC), the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel to give **8a-m** as colorless oils or white amorphous solids. Note 1: DMI or DMI/EtOAc depending on the solubilities of substrates was used as the solvent. Note 2: 2.0 eq. of lutidine was added before addition of Cp_2ZrCl_2 for the syntheses of **8d** and **8e**.

1-(2,2-dimethyl-1,3-dioxolan-4-yl)-4-(4-methoxyphenyl)butan-2-one (8a)



[00624] 51.7 mg (0.186 mmol, 93%); IR (film) 3035, 2988, 2935, 1711, 1612, 1513, 1478, 1370, 1246, 1178, 1058, 1036, 829, 669 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ = 7.09 (d, J = 8.5 Hz, 2H), 6.81 (d, J = 8.5 Hz, 2H), 4.44 (quin, J = 6.0 Hz, 1H), 4.15 (dd, J = 8.5 Hz, 8.0 Hz, 1H), 3.77 (s, 3H), 3.50 (dd, J = 8.5 Hz, 8.0 Hz, 1H), 2.88-2.80 (m, 3H), 2.76-2.71 (m, 2H), 2.52 (dd, J = 16.5 Hz, 7.0 Hz, 1H), 1.38 (s, 3H), 1.33 (s, 3H); ^{13}C NMR (126 MHz, CDCl_3) δ = 207.8, 158.0, 132.8, 129.2, 113.9, 108.8, 71.7, 69.4, 55.2, 47.2, 45.2, 28.7, 26.9, 25.5; HRMS (ESI) m/z calc. for $\text{C}_{16}\text{H}_{22}\text{NaO}_4$ $[\text{M}+\text{Na}]^+$ 301.1410; found 301.1425.

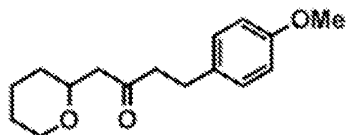
1-(((4*S*,5*S*)-5-(((*tert*-butyldiphenylsilyl)oxy)methyl)-2,2-dimethyl-1,3-dioxolan-4-yl)-4-(4-methoxyphenyl)butan-2-one (8b)



[00625] 102.3 mg (0.187 mg, 94%); $[\alpha]_{\text{D}}^{22}$ = -8.9 (c 1.8, CHCl_3); IR (film) 2985, 2955, 2932, 2898, 2858, 1716, 1612, 1513, 1472, 1463, 1428, 1379, 1370, 1301, 1247, 1177, 1112, 1981, 1037, 998, 823, 787, 742, 704, 603, 505, 490 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) δ = 7.69-7.64 (m, 4H), 7.45-7.35 (m, 6H), 7.10 (d, J = 8.5 Hz, 2H), 6.82 (d, J = 8.5 Hz, 2H), 4.42-4.37

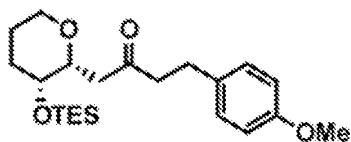
(m, 1H), 3.84-3.71 (m, 3H), 3.78 (s, 3H), 2.85 (t, $J = 8.0$ Hz, 2H), 2.77-2.73 (m, 2H), 2.69-2.65 (m, 2H), 1.38 (s, 3H), 1.37 (s, 3H), 1.06 (s, 9H); ^{13}C NMR (126 MHz, CDCl_3) $\delta = 207.4$, 158.2, 135.8, 133.3, 133.2, 130.0, 129.5, 128.0, 114.1, 109.4, 80.6, 74.8, 64.3, 55.5, 46.7, 45.5, 28.8, 27.4, 27.1, 27.0, 19.4; HRMS (ESI) m/z calc. for $\text{C}_{33}\text{H}_{43}\text{O}_5\text{Si}$ $[\text{M}+\text{H}]^+$ 547.2874; found 547.2869.

4-(4-methoxyphenyl)-1-(tetrahydro-2H-pyran-2-yl)butan-2-one (8c)



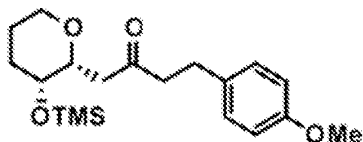
[00626] 49.1 mg (0.188 mg, 94%); IR (film) 2934, 2849, 1712, 1612, 1513, 1441, 1300, 1246, 1178, 1087, 1043, 828 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) $\delta = 7.09$ (d, $J = 8.4$ Hz, 2H), 6.81 (d, $J = 8.4$ Hz, 2H), 3.91 (d, $J = 11.4$ Hz, 1H), 3.78 (s, 3H), 3.77-3.72 (m, 1H), 3.41 (t, $J = 10.8$ Hz, 1H), 2.83 (t, $J = 7.8$ Hz, 2H), 2.74 (q, $J = 5.4$ Hz, 2H), 2.64 (dd, $J = 15.6$ Hz, 7.8 Hz, 1H), 2.36 (dd, $J = 15.6$ Hz, 5.2 Hz, 1H), 1.80 (d, $J = 5.2$ Hz, 1H), 1.58 (d, $J = 12.6$ Hz, 1H), 1.53-1.46 (m, 3H), 1.29-1.21 (m, 1H); ^{13}C NMR (126 MHz, CDCl_3) $\delta = 210.2$, 158.1, 135.7, 133.9, 133.3, 129.8, 129.3, 127.8, 114.0, 63.1, 55.4, 44.7, 39.5, 29.1, 27.0, 26.7, 19.3; HRMS (ESI) m/z calc. for $\text{C}_{16}\text{H}_{23}\text{O}_3$ $[\text{M}+\text{H}]^+$ 263.1642; found 263.1649.

4-(4-methoxyphenyl)-1-((2R,3R)-3-((triethylsilyl)oxy)tetrahydro-2H-pyran-2-yl)butan-2-one (8d)



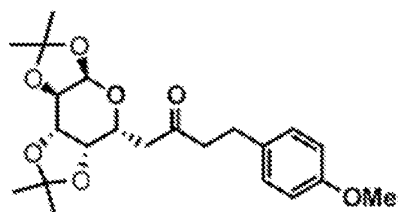
[00627] 73.4 mg (0.187 mg, 94%); $[\alpha]_{\text{D}}^{22} = -11.8$ (c 1.0, CHCl_3); IR (film) 2953, 2915, 2875, 1714, 1612, 1513, 1463, 1300, 1246, 1178, 1098, 1071, 1023, 828, 743 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) $\delta = 7.09$ (d, $J = 8.4$ Hz, 2H), 6.81 (d, $J = 8.4$ Hz, 2H), 3.89 (d, $J = 14.3$ Hz, 1H), 3.81 (ddd, $J = 7.8$, 7.5, 2.5 Hz, 1H), 3.78 (s, 3H), 3.71 (s, 1H), 3.43 (td, $J = 14.4$, 2.5 Hz, 1H), 2.85-2.67 (m, 5H), 2.44 (dd, $J = 19.8$, 6.6 Hz, 1H), 1.96 (m, 1H), 1.81 (m, 1H), 1.71-1.63 (m, 1H), 1.32 (d, $J = 15.5$ Hz, 1H), 0.95 (t, $J = 9.6$ Hz, 9H), 0.59 (q, $J = 9.6$ Hz, 6H); ^{13}C NMR (126 MHz, CDCl_3) $\delta = 208.9$, 158.0, 133.3, 129.4, 114.1, 76.2, 67.9, 67.5, 55.4, 45.8, 45.2, 31.3, 28.8, 20.6, 7.1, 5.1; HRMS (ESI) m/z calc. for $\text{C}_{22}\text{H}_{37}\text{O}_4\text{Si}$ $[\text{M}+\text{H}]^+$ 393.2461; found 393.2449.

4-(4-methoxyphenyl)-1-((2*R*,3*R*)-3-((trimethylsilyl)oxy)tetrahydro-2*H*-pyran-2-yl)butan-2-one (8e)



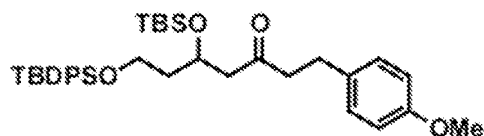
[00628] 61.9 mg (0.177 mg, 89%); $[\alpha]_D^{22} = -15.8$ (*c* 1.0, CHCl₃); IR (film) 2952, 2852, 2839, 1713, 1612, 1513, 1441, 1409, 1300, 1247, 1178, 1137, 1098, 1071, 1023, 840, 763 cm⁻¹; ¹H NMR (500 MHz, C₆D₆) δ = 6.93 (d, *J* = 8.4 Hz, 2H), 6.81 (d, *J* = 8.4 Hz, 2H), 3.77 (ddd, *J* = 6.0, 5.8, 1.5 Hz, 1H), 3.72 (d, *J* = 10.2 Hz, 1H), 3.44 (s, 1H), 3.27 (s, 3H), 3.16 (ddd, *J* = 11.2, 10.8, 1.5 Hz, 1H), 2.81 (m, 2H), 2.71 (dd, *J* = 15.0, 9.0 Hz, 1H), 2.52-2.46 (m, 1H), 2.44-2.38 (m, 1H), 2.26 (dd, *J* = 15.0, 5.1 Hz, 1H), 1.90 (m, 1H), 1.53 (d, *J* = 12.6 Hz, 1H), 1.26 (m, 1H), 0.87 (d, *J* = 12.6 Hz, 1H), 0.00 (s, 9H); ¹³C NMR (126 MHz, C₆D₆) δ = 207.3, 158.6, 133.5, 129.7, 114.3, 76.3, 67.7, 67.5, 54.7, 45.8, 45.4, 31.3, 29.0, 20.6, 0.2; HRMS (ESI) *m/z* calc. for C₁₉H₃₁O₄Si [M+H]⁺ 351.1992; found 351.1978.

4-(4-methoxyphenyl)-1-((3*aR*,5*R*,5*aS*,8*aS*,8*bR*)-2,2,7,7-tetramethyltetrahydro-5*H*-bis([1,3]dioxolo)[4,5-*b*:4',5'-*d*]pyran-5-yl)butan-2-one (8f)



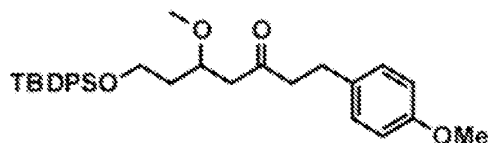
[00629] 75.7 mg (0.186 mmol, 93%); $[\alpha]_D^{22} = -10.7$ (*c* 1.0, CHCl₃); IR (film) 2987, 2935, 1713, 1612, 1513, 1465, 1382, 1456, 1382, 1372, 1246, 1211, 1178, 1099, 1066, 1037, 1000, 861, 547 cm⁻¹; ¹H NMR (600 MHz, CDCl₃) δ = 7.09 (d, *J* = 7.8 Hz, 2H), 6.81 (d, *J* = 7.8 Hz, 2H), 5.46 (d, *J* = 4.7 Hz, 1H), 4.60 (dd, *J* = 7.2, 2.5 Hz, 1H), 4.33-4.28 (m, 2H), 4.18 (d, *J* = 7.2 Hz, 1H), 3.77 (s, 3H), 2.85 (dd, *J* = 6.4, 6.0 Hz, 2H), 2.82-2.73 (m, 3H), 2.65 (dd, *J* = 17.2, 5.0 Hz, 1H), 1.58 (s, 3H), 1.44 (s, 3H), 1.33 (s, 6H); ¹³C NMR (151 MHz, CDCl₃) δ = 207.8, 158.1, 133.2, 129.4, 114.1, 109.3, 108.9, 96.5, 72.6, 70.9, 70.5, 64.2, 55.4, 45.4, 43.6, 28.7, 26.2, 25.2, 24.6; HRMS (ESI) *m/z* calc. for C₂₂H₃₁O₇ [M+H]⁺ 407.2064; found 407.2050.

5-((*tert*-butyldimethylsilyl)oxy)-7-((*tert*-butyldiphenylsilyl)oxy)-1-(4-methoxyphenyl)heptan-3-one (8g)

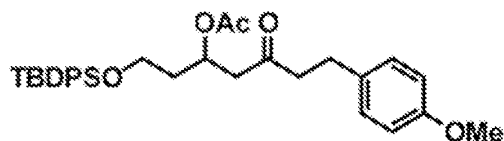


[00630] 103.4 mg, (0.171 mmol, 86%); IR (film) 2955, 2930, 2893, 2856, 1716, 1513, 1472, 1428, 1361, 1248, 1178, 1111, 1084, 1038, 836, 776, 739, 702, 615, 505 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.68-7.65 (m, 4H), 7.45-7.35 (m, 6H), 7.09 (d, J = 7.8 Hz, 2H), 6.82 (d, J = 7.8 Hz, 2H), 4.42-4.37 (m, 1H), 3.78 (s, 3H), 3.71 (t, J = 6.0 Hz, 2H), 2.85-2.79 (m, 1H), 2.72-2.68 (m, 1H), 2.60 (dd, J = 15.6 Hz, 6.6 Hz, 1H), 2.49 (dd, J = 15.6 Hz, 4.8 Hz, 1H), 1.78-1.65 (m, 2H), 1.06 (s, 9H), 0.83 (s, 9H), 0.03 (s, 3H), 0.01 (s, 3H); ^{13}C NMR (151 MHz, CDCl_3) δ = 209.0, 158.0, 135.72, 135.69, 133.87, 133.84, 133.3, 129.75, 129.73, 129.3, 127.8, 114.0, 66.7, 60.5, 55.4, 50.5, 46.5, 40.3, 28.7, 27.0, 26.0, 19.3, 19.1, -4.5, -4.6; HRMS (ESI) m/z calc. for $\text{C}_{36}\text{H}_{53}\text{O}_4\text{Si}_2$ $[\text{M}+\text{H}]^+$ 605.3477; found 605.3464.

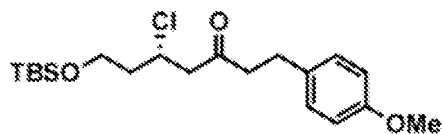
7-((*tert*-butyldiphenylsilyl)oxy)-5-methoxy-1-(4-methoxyphenyl)heptan-3-one (8h)



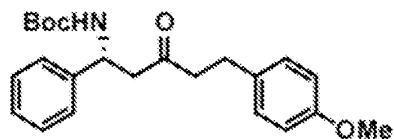
[00631] 94.8 mg (0.188 mmol, 94%); IR (film) 2931, 2896, 2835, 1715, 1612, 1513, 1471, 1464, 1428, 1362, 1300, 1247, 1178, 1111, 1087, 1037, 823, 738, 703, 688, 622, 615, 505, 490, 429 cm^{-1} ; ^1H NMR (600 MHz, C_6D_6) δ = 7.81-7.77 (m, 4H), 7.26-7.22 (m, 6H), 6.98 (d, J = 8.5 Hz, 2H), 6.77 (d, J = 8.5 Hz, 2H), 3.99-3.93 (m, 1H), 3.86-3.80 (m, 1H), 3.76-3.70 (m, 1H), 3.33 (s, 3H), 3.10 (s, 3H), 2.82 (t, J = 8.0 Hz, 2H), 2.44-2.36 (m, 3H), 2.13 (dd, J = 15.5 Hz, 4.8 Hz, 1H), 1.73-1.67 (m, 2H), 1.18 (s, 9H); ^{13}C NMR (126 MHz, C_6D_6) δ = 207.3, 159.0, 136.4, 134.6, 134.0, 130.4, 130.0, 128.5, 114.0, 75.0, 61.1, 57.2, 55.1, 48.1, 46.1, 37.5, 29.4, 27.5, 17.8; HRMS (ESI) m/z calc. for $\text{C}_{31}\text{H}_{40}\text{NaO}_4\text{Si}$ $[\text{M}+\text{Na}]^+$ 527.2588; found 527.2593.

1-((*tert*-butyldiphenylsilyl)oxy)-7-(4-methoxyphenyl)-5-oxoheptan-3-yl acetate (8i)

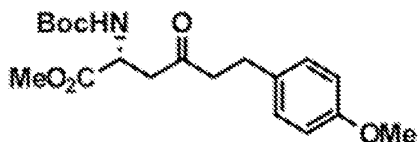
[00632] 90.0 mg (0.169 mmol, 85%); IR (film) 2956, 2931, 2857, 1738, 1716, 1513, 1428, 1363, 1244, 1179, 1111, 1036, 824, 739, 704, 614, 505 cm^{-1} ; ^1H NMR (600 MHz, C_6D_6) δ = 7.79-7.74 (m, 4H), 7.26-7.21 (m, 6H), 6.98 (d, J = 7.8 Hz, 2H), 6.77 (d, J = 7.8 Hz, 2H), 5.67-5.62 (m, 1H), 3.73-3.64 (m, 2H), 3.32 (s, 3H), 2.79 (t, J = 7.2 Hz, 2H), 2.46 (dd, J = 16.2 Hz, 6.6 Hz, 1H), 2.42-2.35 (m, 1H), 2.33-2.26 (m, 1H), 2.22 (dd, J = 16.2 Hz, 6.6 Hz, 2H), 1.86-1.79 (m, 1H), 1.78-1.72 (m, 1H), 1.62 (s, 3H), 1.18 (s, 9H); ^{13}C NMR (126 MHz, C_6D_6) δ = 205.8, 170.0, 159.0, 136.4, 134.4, 133.8, 130.4, 130.0, 128.5, 114.6, 68.4, 60.8, 55.2, 47.7, 45.4, 37.4, 29.4, 27.4, 21.0, 19.8; HRMS (ESI) m/z calc. for $\text{C}_{32}\text{H}_{40}\text{NaO}_5\text{Si}$ $[\text{M}+\text{Na}]^+$ 555.2537; found 555.2533.

((*S*)-7-((*tert*-butyldimethylsilyl)oxy)-5-chloro-1-(4-methoxyphenyl)heptan-3-one (8j)

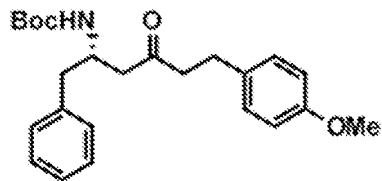
[00633] 28.4 mg (0.074 mmol, 37%); $[\alpha]_{\text{D}}^{22}$ = -11.6 (c 0.5, CHCl_3) IR (film) 2954, 2928, 2856, 1738, 1716, 1612, 1513, 1463, 1300, 1247, 1178, 1123, 1038, 838, 779 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 6.10 (d, J = 7.8 Hz, 2H), 6.82 (d, J = 7.8 Hz, 2H), 3.88 (m, 1H), 3.78 (s, 3H), 3.77 (dd, J = 10.2, 5.1 Hz, 1H), 3.67 (dd, J = 10.2, 6.6 Hz, 1H), 2.85 (t, J = 7.2 Hz, 2H), 2.72 (t, J = 7.2 Hz, 2H), 2.68-2.55 (m, 2H), 2.25-2.19 (m, 1H), 1.83-1.77 (m, 1H), 0.89 (s, 9H), 0.07 (s, 6H); ^{13}C NMR (126 MHz, C_6D_6) δ = 209.3, 158.2, 133.1, 129.4, 114.1, 67.4, 62.2, 55.4, 44.8, 39.5, 29.1, 28.3, 26.0, 18.5, -5.9, -5.3; HRMS (ESI) m/z calc. for $\text{C}_{20}\text{H}_{34}\text{ClO}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 385.1960; found 385.1943.

***R*-(5-(4-methoxyphenyl)-3-oxo-1-phenylpentyl)carbamate (8k)**

[00634] 69.6 mg (0.182 mmol, 91%); $[\alpha]_D^{22} = +14.7$ (*c* 0.3, CHCl₃); IR (film) 3376, 2979, 2932, 1707, 1612, 1513, 1455, 1366, 1247, 1175, 1037, 819, 701 cm⁻¹; ¹H NMR (600 MHz, CDCl₃) δ = 7.33-7.29 (m, 2H), 7.27-7.22 (m, 3H), 7.00 (d, *J* = 8.4 Hz, 2H), 6.78 (d, *J* = 8.4 Hz, 2H), 5.46 (brs, 1H), 5.07 (brs, 1H), 3.78 (s, 3H), 3.00 (brs, 1H), 2.85 (dd, *J* = 17.4 Hz, 4.3 Hz, 1H), 2.73 (t, *J* = 7.8 Hz, 2H), 2.69-2.62 (m, 1H), 2.59-2.52 (m, 1H), 1.41 (s, 9H); ¹³C NMR (126 MHz, CDCl₃) δ = 208.5, 158.1, 155.3, 141.7, 132.9, 129.3, 128.8, 127.5, 126.4, 114.0, 79.9, 55.4, 51.3, 48.8, 45.4, 28.7, 28.5; HRMS (ESI) *m/z* calc. for C₂₃H₂₉NNaO₄ [M+Na]⁺ 406.1989; found 406.1980.

methyl (*R*)-2-((*tert*-butoxycarbonyl)amino)-6-(4-methoxyphenyl)-4-oxohexanoate (8l)

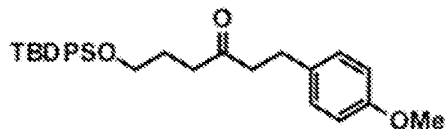
[00635] 62.8 mg (0.172 mmol, 86%); $[\alpha]_D^{22} = +19.0$ (*c* 0.8, CHCl₃); IR (film) 3383, 2974, 2953, 2932, 1749, 1713, 1612, 1513, 1454, 1439, 1367, 1342, 1299, 1247, 1165, 1110, 1088, 1034, 830 cm⁻¹; ¹H NMR (600 MHz, C₆D₆) δ = 6.83 (d, *J* = 8.4 Hz, 2H), 6.74 (d, *J* = 8.4 Hz, 2H), 5.63 (brs, 1H), 4.62 (brs, 1H), 3.32 (s, 3H), 3.27 (s, 3H), 2.70 (d, *J* = 19.2 Hz, 1H), 2.61 (q, *J* = 6.6 Hz, 2H), 2.50 (d, *J* = 19.2 Hz, 1H), 2.16-2.06 (m, 2H), 1.42 (s, 9H); ¹³C NMR (126 MHz, C₆D₆) δ = 207.7, 172.2, 159.0, 156.0, 133.4, 129.9, 114.6, 79.9, 55.1, 52.4, 50.4, 44.9, 44.7, 29.2, 28.7; HRMS (ESI) *m/z* calc. for C₁₉H₂₇NNaO₆ [M+Na]⁺ 388.1731; found 388.1740.

***tert*-butyl (*S*)-(6-(4-methoxyphenyl)-4-oxo-1-phenylhexan-2-yl)carbamate (8m)**

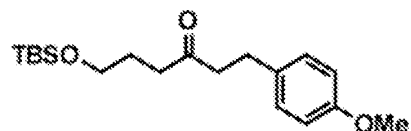
[00636] 71.9 mg (0.181 mmol, 91%); $[\alpha]_D^{22} = -5.7$ (*c* 1.1, CHCl₃); IR (film) 3360, 2977, 2931, 1708, 1612, 1513, 1455, 1391, 1366, 1301, 1247, 1174, 1109, 1077, 1037, 824, 778, 702 cm⁻¹; ¹H NMR (600 MHz, CDCl₃) δ = 7.29-7.25 (m, 2H), 7.21 (t, *J* = 7.2 Hz, 1H), 7.10 (d, *J* = 7.2 Hz, 2H), 7.08 (d, *J* = 9.0 Hz, 2H), 6.82 (d, *J* = 9.0 Hz, 2H), 5.04 (brs, 1H), 4.11 (brs, 1H), 3.78 (s, 3H), 2.91 (brs, 1H), 2.84-2.75 (m, 3H), 2.71-2.58 (m, 2H), 2.54 (d, *J* = 4.9 Hz, 2H), 1.40 (s, 9H); ¹³C NMR (126 MHz, CDCl₃) δ = 209.5, 158.1, 155.4, 138.2, 132.9, 129.4, 129.3, 128.7, 126.7, 114.1, 79.5, 55.4, 48.9, 45.6, 45.1, 40.4, 28.8, 28.5; HRMS (ESI) *m/z* calc. for C₂₄H₃₂NO₄ [M+H]⁺ 398.2331; found 398.2326.

Experimental Procedures for the Reactions Outlined in Figure 5A

[00637] In a glove box, to a solution of iodide **9a-u** (0.24 mmol, 1.2 eq.) and thioester **1-5** (54.6 mg, 0.20 mmol, 1.0 eq.) in either DMI (0.4 mL, sigma aldrich, 99.5%) or DMI/EtOAc (0.334 mL/0.066mL) were added Cp₂ZrCl₂ (58.5 mg, 0.20 mmol, 1.0 eq. Sigma-aldrich, 98%), Zn powder (39.2 mg, 0.6 mmol, 3.0 eq. Sigma-aldrich, used without any activation), and NiBr₂·dtbbpy (9.7 mg, 0.02 mmol, 10 mol%, preparation see page 8) at room temperature. After being stirred at the same temperature for 10 min to 3 hr (monitored by TLC), the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel to give **10a-u** as colorless oils or white amorphous solids. Note 1: DMI or DMI/EtOAc depending on the solubilities of substrates was used as the solvent. Note 2: 2.0 eq. of lutidine was added before addition of Cp₂ZrCl₂ for the syntheses of **10c**. Note 3: 1.5 eq. of iodide **9p-u**, and 1.5 eq. of Cp₂ZrCl₂ were used during syntheses of **10p-u**.

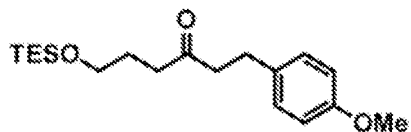
6-((*tert*-butyldiphenylsilyl)oxy)-1-(4-methoxyphenyl)hexan-3-one (10a)

[00638] 87.9 mg (0.191 mmol, 96%); IR (film) 2952, 2931, 2834, 1714, 1513, 1247, 1036, 975, 823, 688, 613, 487 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.66-7.64 (m, 4H), 7.44-7.36 (m, 6H), 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 3.78 (s, 3H), 3.66 (t, J = 5.9 Hz, 2H), 2.83 (t, J = 7.2 Hz, 2H), 2.69 (t, J = 7.8 Hz, 2H), 2.51 (t, J = 7.2 Hz, 2H), 1.85-1.80 (m, 2H), 1.05 (s, 9H); ^{13}C NMR (151 MHz, CDCl_3) δ = 210.2, 158.1, 135.7, 133.9, 133.3, 129.8, 129.3, 127.8, 114.0, 63.1, 55.4, 44.7, 39.5, 29.1, 27.0, 26.7, 19.3; HRMS (ESI) m/z calc. for $\text{C}_{29}\text{H}_{37}\text{O}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 461.2506; found 461.2508.

6-((*tert*-butyldimethylsilyl)oxy)-1-(4-methoxyphenyl)hexan-3-one (10b)

[00639] 63.8 mg (0.190 mmol, 95%); IR (film) 2954, 2929, 2857, 1715, 1513, 1247, 1097, 1038, 835, 776 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 3.78 (s, 3H), 3.59 (t, J = 6.0 Hz, 2H), 2.84 (t, J = 7.2 Hz, 2H), 2.71 (t, J = 7.8 Hz, 2H), 2.46 (t, J = 7.2 Hz, 2H), 1.79-1.74 (m, 2H), 0.88 (s, 9H), 0.03 (s, 6H); ^{13}C NMR (151 MHz, CDCl_3) δ = 210.3, 158.1, 133.3, 129.4, 114.0, 62.3, 55.4, 44.8, 39.5, 29.1, 26.9, 26.1, 18.4, -5.2; HRMS (ESI) m/z calc. for $\text{C}_{19}\text{H}_{33}\text{O}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 337.2193; found 337.2186.

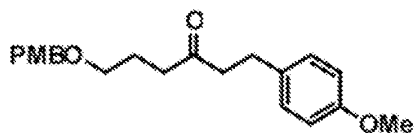
1-(4-methoxyphenyl)-6-((triethylsilyl)oxy)hexan-3-one (10c)



[00640] 61.5 mg (0.183 mmol, 92%); IR (film) 2953, 2876, 2835, 1715, 1612, 1513, 1464, 1247, 1178, 1095, 1038, 1005, 826, 808, 743 cm^{-1} ; ^1H NMR (600 MHz, C_6D_6) δ = 6.97 (d, J = 8.4 Hz, 2H), 6.77 (d, J = 8.4 Hz, 2H), 3.47 (t, J = 5.9 Hz, 2H), 3.33 (s, 3H), 2.80 (t, J = 7.8 Hz, 2H), 2.31 (t, J = 7.8 Hz, 2H), 2.15 (t, J = 7.2 Hz, 2H), 1.80-1.75 (m, 2H), 0.99 (t, J = 7.2 Hz, 9H), 0.58 (q, J = 7.2 Hz, 6H); ^{13}C NMR (151 MHz, C_6D_6) δ = 208.3, 159.0, 134.0, 130.0,

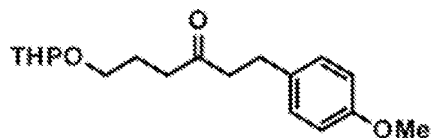
114.6, 62.4, 55.1, 44.9, 39.5, 29.6, 27.6, 7.5, 5.2; HRMS (ESI) m/z calc. for $C_{19}H_{33}O_3Si$ $[M+H]^+$ 337.2193; found 337.2186.

6-((4-methoxybenzyl)oxy)-1-(4-methoxyphenyl)hexan-3-one (10d)



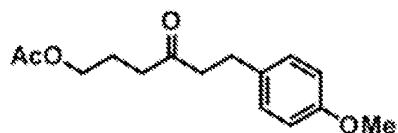
[00641] 64.9 mg (0.190 mmol, 95%); IR (film) 2932, 2855, 2835, 1711, 1612, 1585, 1512, 1464, 1441, 1363, 1301, 1245, 1177, 1095, 1034, 819 cm^{-1} ; ^1H NMR 7.23 (d, J = 9.0 Hz, 2H), 7.08 (d, J = 9.0 Hz, 2H), 6.87 (d, J = 9.0 Hz, 2H), 6.81 (d, J = 9.0 Hz, 2H), 4.39 (s, 2H), 3.80 (s, 3H), 3.78 (s, 3H), 3.43 (t, J = 6.0 Hz, 2H), 2.82 (t, J = 7.2 Hz, 2H), 2.68 (t, J = 7.2 Hz, 2H), 2.48 (t, J = 7.2 Hz, 2H), 1.89-1.84 (m, 2H); ^{13}C NMR (126 MHz, $CDCl_3$) δ = 210.1, 159.3, 158.1, 133.3, 130.6, 129.39, 129.36, 114.0, 113.9, 72.6, 69.1, 55.40, 55.39, 44.7, 39.8, 29.0, 24.0; HRMS (ESI) m/z calc. for $C_{21}H_{26}NaO_4$ $[M+Na]^+$ 365.1723; found 365.1724.

1-(4-methoxyphenyl)-6-((tetrahydro-2H-pyran-2-yl)oxy)hexan-3-one (10e)



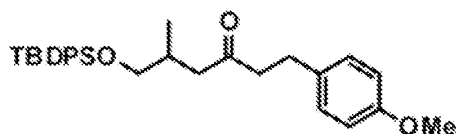
[00642] 55.4 mg (0.181 mmol, 91%); IR (film) 2940, 2870, 1712, 1612, 1513, 1442, 1331, 1246, 1179, 1076, 1034, 991, 815 cm^{-1} ; ^1H NMR (600 MHz, $CDCl_3$) δ = 7.09 (d, J = 9.0 Hz, 2H), 6.81 (d, J = 9.0 Hz, 2H), 4.53 (s, 1H), 3.82 (t, J = 9.6 Hz, 1H), 3.78 (s, 3H), 3.71 (q, J = 6.0 Hz, 1H), 3.48 (t, J = 5.0 Hz, 1H), 3.38 (q, J = 6.0 Hz, 1H), 2.84 (t, J = 7.8 Hz, 2H), 2.72 (t, J = 7.8 Hz, 2H), 2.54-2.44 (m, 2H), 1.89-1.83 (m, 2H), 1.80 (d, J = 8.4 Hz, 1H), 1.71-1.65 (m, 1H), 1.58-1.48 (m, 4H); ^{13}C NMR (151 MHz, $CDCl_3$) δ = 209.9, 157.9, 133.2, 129.2, 113.9, 76.8, 66.5, 62.4, 55.2, 44.6, 39.8, 30.7, 28.9, 25.4, 23.9, 19.7; HRMS (ESI) m/z calc. for $C_{18}H_{26}NaO_4$ $[M+Na]^+$ 329.1723; found 329.1722.

6-(4-methoxyphenyl)-4-oxohexyl acetate (10f)



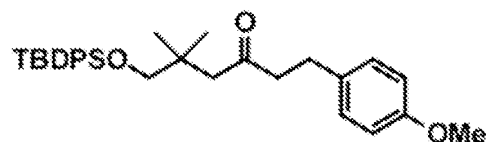
[00643] 49.9 mg (0.189 mmol, 95%); IR (film) 2959, 2935, 1734, 1711, 1512, 1364, 1238, 1177, 1109, 1034, 761 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.09 (d, J = 9.0 Hz, 2H), 6.81 (d, J = 9.0 Hz, 2H), 4.03 (t, J = 6.6 Hz, 2H), 3.77 (s, 3H), 2.84 (t, J = 7.2 Hz, 2H), 2.70 (t, J = 7.2 Hz, 2H), 2.45 (t, J = 7.2 Hz, 2H), 2.02 (s, 3H), 1.92-1.86 (m, 2H); ^{13}C NMR (151 MHz, CDCl_3) δ = 209.2, 171.2, 158.1, 133.1, 129.4, 114.0, 63.7, 55.4, 44.7, 39.3, 29.0, 22.7, 21.0; HRMS (ESI) m/z calc. for $\text{C}_{15}\text{H}_{21}\text{O}_4$ $[\text{M}+\text{H}]^+$ 265.1434; found 265.1433.

6-((*tert*-butyldiphenylsilyl)oxy)-1-(4-methoxyphenyl)-5-methylhexan-3-one (10g)

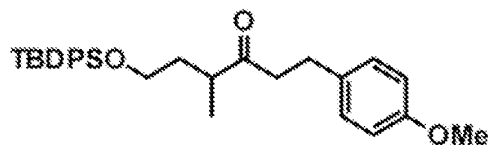


[00644] 85.4 mg (0.180 mmol, 90%); IR (film) 2959, 2931, 2857, 1713, 1513, 1463, 1442, 1247, 1178, 1111, 1037, 824, 741, 702, 614, 506 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.66-7.63 (m, 4H), 7.44-7.41 (m, 2H), 7.40-7.36 (m, 4H), 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 3.78 (s, 3H), 3.52 (dd, J = 9.6 Hz, 5.2 Hz, 1H), 3.43 (dd, J = 10.2 Hz, 6.6 Hz, 1H), 2.82 (t, J = 8.4 Hz, 2H), 2.68 (td, J = 7.8 Hz, 2.0 Hz, 2H), 2.63 (dd, J = 16.2 Hz, 5.2 Hz, 1H), 2.28-2.22 (m, 1H), 2.18 (dd, J = 16.2 Hz, 16.0 Hz, 1H), 1.05 (s, 9H), 0.88 (d, J = 6.6 Hz, 3H); ^{13}C NMR (126 MHz, CDCl_3) δ = 209.9, 157.9, 135.6, 133.7, 133.2, 129.6, 129.2, 127.7, 113.9, 68.3, 55.3, 46.8, 45.1, 32.0, 28.9, 26.9, 19.3, 16.8; HRMS (ESI) m/z calc. for $\text{C}_{30}\text{H}_{39}\text{O}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 475.2663; found 475.2654.

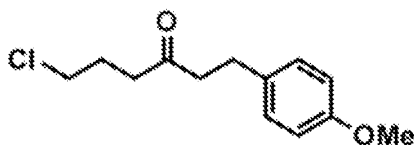
6-((*tert*-butyldiphenylsilyl)oxy)-1-(4-methoxyphenyl)-5,5-dimethylhexan-3-one (10h)



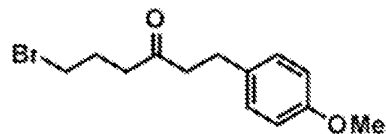
[00645] 88.4 mg (0.181 mmol, 91%); IR (film) 2958, 2858, 1711, 1512, 1264, 1178, 907, 825, 731, 703, 650, 436 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.64-7.62 (m, 4H), 7.44-7.41 (m, 2H), 7.39-7.36 (m, 4H), 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 3.78 (s, 3H), 3.39 (s, 2H), 2.79 (t, J = 7.8 Hz, 2H), 2.68 (t, J = 7.8 Hz, 2H), 2.42 (s, 2H), 1.06 (s, 9H), 0.97 (s, 6H); ^{13}C NMR (126 MHz, CDCl_3) δ = 210.2, 158.0, 135.8, 133.8, 133.4, 129.8, 129.4, 127.8, 114.0, 72.2, 55.4, 50.2, 46.9, 36.3, 29.0, 27.1, 24.5, 19.6; HRMS (ESI) m/z calc. for $\text{C}_{31}\text{H}_{41}\text{O}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 489.2819; found 489.2832.

6-((*tert*-butyldiphenylsilyl)oxy)-1-(4-methoxyphenyl)-4-methylhexan-3-one (10i)

[00646] 90.1 mg (0.190 mmol, 95%); IR (film) 2959, 2931, 2857, 1710, 1612, 1513, 1463, 1247, 1178, 1111, 1038, 823, 740, 703, 614, 519 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.66-7.64 (m, 4H), 7.45-7.41 (m, 2H), 7.40=7.37 (m, 4H), 7.09 (d, J = 9.0 Hz, 2H), 6.82 (d, J = 9.0 Hz, 2H), 3.79 (s, 3H), 3.65 (t, J = 6.6 Hz, 2H), 2.84-2.70 (m, 5H), 1.97-1.90 (m, 1H), 1.53-1.47 (m, 1H), 1.06 (s, 9H), 1.02 (d, J = 7.2 Hz, 3H); ^{13}C NMR (126 MHz, CDCl_3) δ = 213.6, 157.9, 135.6, 133.7, 133.4, 129.7, 129.3, 127.7, 113.9, 61.6, 55.3, 43.1, 42.9, 35.3, 28.9, 26.9, 19.2, 16.2; HRMS (ESI) m/z calc. for $\text{C}_{30}\text{H}_{39}\text{O}_3\text{Si}$ $[\text{M}+\text{H}]^+$ 475.2663; found 475.2657.

6-chloro-1-(4-methoxyphenyl)hexan-3-one (10l)

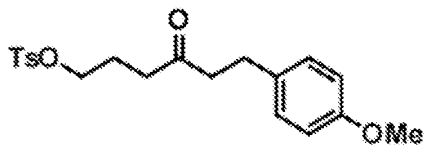
[00647] 46.0 mg (0.192 mmol, 96%); IR (film) 2932, 2836, 1712, 1612, 1513, 1442, 1374, 1300, 1245, 1178, 1091, 1034, 829, 546 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.10 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 3.55 (t, J = 6.6 Hz, 2H), 2.85 (t, J = 7.8 Hz, 2H), 2.72 (t, J = 7.8 Hz, 2H), 2.58 (t, J = 7.8 Hz, 2H), 2.05-2.00 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 209.1, 158.2, 133.0, 129.4, 114.1, 55.4, 44.8, 44.6, 29.7, 29.1, 26.4; HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{18}\text{ClO}_2$ $[\text{M}+\text{H}]^+$ 241.0990; found 241.0998.

6-bromo-1-(4-methoxyphenyl)hexan-3-one (10m)

[00648] 54.3 mg (0.191 mmol, 96%); IR (film) 2933, 2835, 1712, 1611, 1512, 1441, 1409, 1372, 1300, 1245, 1178, 1035, 828, 555 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.09 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 3.42 (t, J = 6.6 Hz, 2H), 2.85 (t, J = 7.8 Hz, 2H), 2.72 (t, J = 7.8 Hz, 2H), 2.58 (t, J = 7.8 Hz, 2H), 2.13-2.08 (m, 2H); ^{13}C NMR (126

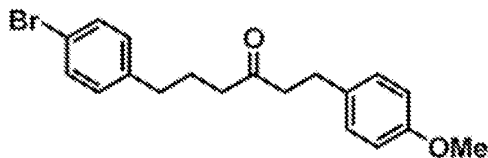
MHz, CDCl₃) δ = 209.0, 158.1, 133.0, 129.4, 114.1, 55.4, 44.8, 40.9, 33.4, 29.1, 26.4; HRMS (ESI) m/z calc. for C₁₃H₁₈BrO₂ [M+H]⁺ 285.0485; found 285.0476.

6-(4-methoxyphenyl)-4-oxohexyl 4-methylbenzenesulfonate (10n)

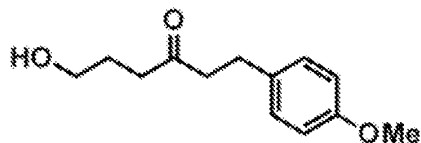


[00649] 61.7 mg (0.164 mmol, 82%); IR (film) 2960, 2936, 1714, 1612, 1513, 1465, 1455, 1443, 1416, 1359, 1302, 1246, 1189, 1175, 1098, 1037, 1037, 1019, 963, 931, 921, 903, 830, 814, 795, 664, 543 cm⁻¹; ¹H NMR (600 MHz, C₆D₆) δ = 7.72 (d, J = 8.4 Hz, 2H), 6.94 (d, J = 8.4 Hz, 2H), 6.79 (d, J = 8.4 Hz, 2H), 6.69 (d, J = 8.4 Hz, 2H), 3.79 (t, J = 6.6 Hz, 2H), 3.34 (s, 3H), 2.70 (t, J = 7.2 Hz, 2H), 2.14 (t, J = 7.2 Hz, 2H), 1.85 (t, J = 6.6 Hz, 2H), 1.82 (s, 3H), 1.59-1.54 (m, 2H); ¹³C NMR (126 MHz, C₆D₆) δ = 207.4, 159.0, 144.6, 134.7, 133.8, 130.2, 129.9, 114.6, 70.0, 55.2, 55.1, 44.7, 38.3, 29.5, 23.5, 21.5; HRMS (ESI) m/z calc. for C₂₀H₂₄NaO₅S [M+Na]⁺ 399.1237; found 399.1221.

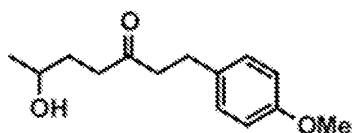
6-(4-bromophenyl)-1-(4-methoxyphenyl)hexan-3-one (10o)



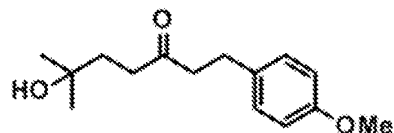
[00650] 57.6 mg (0.160 mmol, 82%); IR (film) 2934, 1712, 1612, 1512, 1488, 1454, 1404, 1370, 1300, 1246, 1178, 1109, 1035, 1011, 824, 518 cm⁻¹; ¹H NMR (600 MHz, CDCl₃) δ = 7.38 (d, J = 8.4 Hz, 2H), 7.08 (d, J = 8.4 Hz, 2H), 7.00 (d, J = 8.4 Hz, 2H), 6.81 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 2.82 (t, J = 8.4 Hz, 2H), 2.66 (t, J = 8.4 Hz, 2H), 2.53 (t, J = 7.2 Hz, 2H), 2.36 (t, J = 7.2 Hz, 2H), 1.88-1.82 (m, 2H); ¹³C NMR (151 MHz, CDCl₃) δ = 209.9, 158.1, 140.7, 133.2, 131.5, 130.3, 129.4, 119.8, 114.0, 55.4, 44.7, 42.1, 34.5, 29.0, 25.0; HRMS (ESI) m/z calc. for C₁₉H₂₁BrNaO₂ [M+Na]⁺ 383.0617; found 383.0608.

6-hydroxy-1-(4-methoxyphenyl)hexan-3-one (10p)

[00651] 27.3 mg (0.123 mmol, 62%); IR (film) 3523-3306 (br), 2918, 1708, 1612, 1513, 1299, 1246, 1179, 1107, 1066, 848 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.10 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 3.63 (t, J = 6.6 Hz, 2H), 2.85 (t, J = 6.6 Hz, 2H), 2.73 (t, J = 6.6 Hz, 2H), 2.53 (t, J = 6.6 Hz, 2H), 1.85-1.80 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 210.9, 158.1, 133.1, 129.3, 114.1, 62.5, 55.4, 44.8, 39.9, 29.1, 26.5; HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{17}\text{O}_2$ $[\text{M}+\text{H}-\text{H}_2\text{O}]^+$ 205.1223; found 205.1223. Note: Exists as a mixture of ketone and hemiacetal (30:1).

6-hydroxy-1-(4-methoxyphenyl)heptan-3-one (10q)

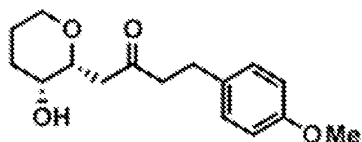
[00652] 30.9 mg (0.131 mmol, 66%); IR (film) 3513-3300 (br), 2916, 1705, 1610, 1513, 1299, 1246, 1179, 1107, 1100, 1087, 845 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 7.10 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 3.78-3.76 (m, 1H), 2.87-2.80 (m, 2H), 2.77-2.68 (m, 2H), 2.58-2.49 (m, 2H), 1.65-1.55 (m, 2H), 1.18 (d, J = 5.2 Hz, 3H); ^{13}C NMR (126 MHz, CDCl_3) δ = 211.1, 158.0, 133.4, 129.3, 114.1, 67.6, 55.4, 44.8, 39.5, 32.7, 29.1, 23.9; HRMS (ESI) m/z calc. for $\text{C}_{14}\text{H}_{20}\text{O}_3$ $[\text{M}+\text{H}]^+$ 237.1491; found 237.1485. Note: Exists as a mixture of ketone and hemiacetal (20:1).

6-hydroxy-1-(4-methoxyphenyl)-6-methylheptan-3-one (10r)

[00653] 24.8 mg (0.099 mmol, 50%); IR (film) 3550-3450 (br), 2966, 2928, 1708, 1611, 1512, 1464, 1366, 1300, 1244, 1177, 1138, 1035, 822 cm^{-1} ; ^1H NMR (500 MHz, CDCl_3) (only peaks of the ketone in the mixture are shown) δ = 7.09 (d, J = 8.4 Hz, 2H), 6.82 (d, J = 8.4 Hz, 2H), 3.78 (s, 3H), 2.84 (t, J = 9.6 Hz, 2H), 2.74 (t, J = 9.6 Hz, 2H), 2.52 (t, J = 9.6 Hz,

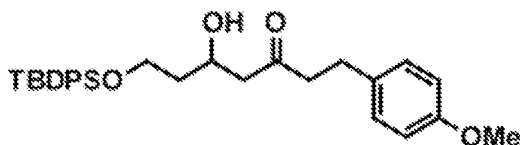
2H), 1.74 (t J = 9.6 Hz, 2H), 1.19 (s, 6H); ^{13}C NMR (126 MHz, CDCl_3) (all peaks of the mixture are shown) δ = 210.9, 133.0, 129.2, 129.1, 113.9, 113.8, 70.1, 55.3, 44.8, 43.2, 38.0, 37.2, 36.6, 36.2, 30.4, 30.0, 29.4, 29.0 HRMS (ESI) m/z calc. for $\text{C}_{15}\text{H}_{23}\text{O}_3$ $[\text{M}+\text{H}]^+$ 251.1647; found 251.1639. Note: Exists as a mixture of ketone and hemiacetal (2.5:1).

1-((2*R*,3*R*)-3-hydroxytetrahydro-2*H*-pyran-2-yl)-4-(4-methoxyphenyl)butan-2-one (10s)

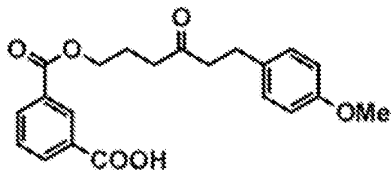


[00654] 25.3 mg (0.091 mmol); ^1H NMR shows complex mixtures which are considered as a mixture of the ketone and two hemiacetal isomers. ^1H NMR of **10s** is shown in Part 8 of this supporting information. In order to confirm the structure, **10s** was subject to TESOTf (1.2 eq.) and 2,6-lutidine (1.5 eq.) in dichloromethane. The expected **8e** was isolated as the major product for 87% yield.

7-((*tert*-butyldiphenylsilyl)oxy)-5-hydroxy-1-(4-methoxyphenyl)heptan-3-one (10t)



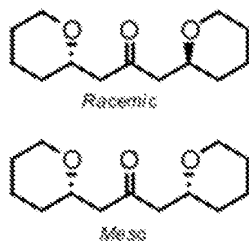
[00655] 40.7 mg (0.083 mmol, 42%); IR (film) 3489 (br), 2930, 2857, 1711, 1612, 1513, 1471, 1428, 1301, 1247, 1178, 1111, 1038, 823, 739, 703, 689, 617, 504 cm^{-1} ; ^1H NMR (600 MHz, C_6D_6) δ = 7.80-7.75 (m, 4H), 7.25-7.20 (m, 6H), 6.95 (d, J = 9.0 Hz, 2H), 6.78 (d, J = 9.0 Hz, 2H), 4.33-4.27 (m, 1H), 3.88-3.83 (m, 1H), 3.80-3.76 (m, 1H), 3.32 (s, 3H), 3.28 (d, J = 2.9 Hz, 1H), 2.77-2.73 (m, 2H), 2.27 (t, J = 7.8 Hz, 2H), 2.19 (dd, J = 16.8 Hz, 9.0 Hz, 1H), 2.02 (dd, J = 16.8 Hz, 3.4 Hz, 2H), 1.68-1.61 (m, 1H), 1.52-1.46 (m, 1H), 1.16 (s, 9H); ^{13}C NMR (126 MHz, C_6D_6) δ = 209.7, 159.0, 136.4, 134.3, 133.8, 130.4, 130.0, 114.6, 110.8, 110.7, 66.6, 62.5, 55.2, 50.1, 45.7, 39.6, 29.3, 27.5, 19.7; HRMS (ESI) m/z calc. for $\text{C}_{30}\text{H}_{39}\text{O}_4\text{Si}$ $[\text{M}+\text{H}]^+$ 491.2612; found 491.2604.

3-(((6-(4-methoxyphenyl)-4-oxohexyl)oxy)carbonyl)benzoic acid (10u)

[00656] 42.9 mg (0.116 mmol, 58%); IR (film) 2951, 2905, 2834, 1721, 1610, 1508, 1483, 14691, 1280, 1170, 1105, 1087, 845, 721 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 8.73 (s, 1H), 8.30 (d, J = 5.6 Hz, 1H), 8.26 (d, J = 5.6 Hz, 1H), 7.59 (t, J = 5.6 Hz, 1H), 7.08 (d, J = 8.4 Hz, 2H), 6.80 (d, J = 8.4 Hz, 2H), 4.35 (t, J = 6.6 Hz, 2H), 3.75 (s, 3H), 2.86 (t, J = 7.8 Hz, 2H), 2.73 (t, J = 7.8 Hz, 2H), 2.56 (t, J = 6.6 Hz, 2H), 2.07 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 209.2, 170.8, 165.8, 158.4, 134.8, 134.5, 133.0, 131.4, 131.0, 129.8, 129.4, 128.9, 114.0, 64.8, 55.4, 44.8, 39.4, 29.3, 22.9; HRMS (ESI) m/z calc. for $\text{C}_{21}\text{H}_{22}\text{NaO}_6$ $[\text{M}+\text{Na}]^+$ 393.1314; found 393.1303.

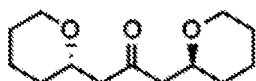
Experimental Procedures for the Reactions Outlined in Figure 5B

[00657] In a glove box, to a solution of iodide **1-12** (27.1 mg, 0.12 mmol, 1.2 eq.) and thioester **1-11** (23.7 mg, 0.10 mmol, 1.0 eq.) in DMI (0.2 mL, Sigma-aldrich, 99.5%) were added Cp_2ZrCl_2 (29.3 mg, 0.10 mmol, 1.0 eq. Sigma-aldrich, 98%), Zn powder (19.6 mg, 0.3 mmol, 3.0 eq. Sigma-aldrich, used without any activation), and $\text{NiBr}_2 \cdot \text{dtbbpy}$ (4.8 mg, 0.01 mmol, 10 mol%, preparation see page 8) at room temperature. After being stirred at the same temperature for 40 mins (monitored by TLC), the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with ethyl acetate. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel to give **1-13** as a colorless oil.

1,3-bis(tetrahydro-2H-pyran-2-yl)propan-2-one (1:1 mixture-1-13)

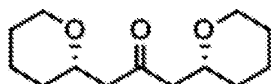
[00658] 19.4 mg (0.086 mmol, 86%); IR (film) 2933, 2487, 1713, 1440, 1378, 1356, 1203, 1175, 1088 cm^{-1} ; ^1H NMR (600 MHz, CDCl_3) δ = 3.91 (d, J = 9.2 Hz, 2H), 3.79-3.73 (m, 2H), 3.43 (dd, J = 11.2, 10.8 Hz, 2H), 2.67 (dd, J = 14.8, 5.8 Hz, 2H), 2.44 (dd, J = 14.8, 5.8 Hz, 2H), 1.80 (d, J = 7.2 Hz, 2H), 1.62-1.58 (m, 3H), 1.52-1.46 (m, 5H), 1.30-1.21 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 207.7, 74.1, 68.7, 50.6, 50.4, 31.9, 25.9, 23.5 HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{22}\text{NaO}_3$ $[\text{M}+\text{Na}]^+$ 249.1467; found 249.1460.

1,3-bis((*S*)-tetrahydro-2*H*-pyran-2-yl)propan-2-one [(*S*)-1-13]



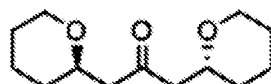
[00659] 20.2 mg (0.089 mmol, 89%); $[\alpha]_D^{22}$ = -7.3 (c 0.74, CHCl_3); ^1H NMR (600 MHz, CDCl_3) δ = 3.91 (d, J = 9.2 Hz, 2H), 3.79-3.73 (m, 2H), 3.43 (dd, J = 11.2, 10.8 Hz, 2H), 2.67 (dd, J = 14.8, 5.8 Hz, 2H), 2.44 (dd, J = 14.8, 5.8 Hz, 2H), 1.80 (d, J = 7.2 Hz, 2H), 1.62-1.58 (m, 3H), 1.52-1.46 (m, 5H), 1.30-1.21 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 207.7, 74.1, 68.7, 50.6, 31.9, 25.9, 23.5 ppm; HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{22}\text{NaO}_3$ $[\text{M}+\text{Na}]^+$ 249.1467; found 249.1463.

1-((*R*)-tetrahydro-2*H*-pyran-2-yl)-3-((*S*)-tetrahydro-2*H*-pyran-2-yl)propan-2-one[(*S*,*R*)-1-13]



[00660] 19.2 mg (0.085 mmol, 85%) from (*S*)-1-11; 19.4 mg (0.086 mmol, 86%) from (*R*)-1-11. ^1H NMR (600 MHz, CDCl_3) δ = 3.91 (d, J = 9.2 Hz, 2H), 3.79-3.73 (m, 2H), 3.43 (dd, J = 11.2, 10.8 Hz, 2H), 2.67 (dd, J = 14.8, 5.8 Hz, 2H), 2.44 (dd, J = 14.8, 5.8 Hz, 2H), 1.80 (d, J = 7.2 Hz, 2H), 1.62-1.58 (m, 3H), 1.52-1.46 (m, 5H), 1.30-1.21 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 207.7, 74.1, 68.7, 50.4, 31.9, 25.9, 23.5 ppm; HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{22}\text{NaO}_3$ $[\text{M}+\text{Na}]^+$ 249.1467; found 249.1463.

1,3-bis((*R*)-tetrahydro-2*H*-pyran-2-yl)propan-2-one [(*R*)-1-13]



[00661] 19.0 mg (0.084 mmol, 84%); $[\alpha]_D^{25} = +7.6$ (c 0.77, CHCl_3); ^1H NMR (600 MHz, CDCl_3) δ = 3.91 (d, J = 9.2 Hz, 2H), 3.79-3.73 (m, 2H), 3.43 (dd, J = 11.2, 10.8 Hz, 2H), 2.67 (dd, J = 14.8, 5.8 Hz, 2H), 2.44 (dd, J = 14.8, 5.8 Hz, 2H), 1.80 (d, J = 7.2 Hz, 2H), 1.62-1.58 (m, 3H), 1.52-1.46 (m, 5H), 1.30-1.21 (m, 2H); ^{13}C NMR (126 MHz, CDCl_3) δ = 207.7, 74.1, 68.7, 50.6, 31.9, 25.9, 23.5 ppm; HRMS (ESI) m/z calc. for $\text{C}_{13}\text{H}_{22}\text{NaO}_3$ $[\text{M}+\text{Na}]^+$ 249.1467; found 249.1455.

Synthesis of Halichondrins and Analogs

[00662] A unified, efficient, and scalable synthesis of halichondrins, with use of Zr/Ni-mediated one-pot ketone synthesis as the final coupling reaction has been developed. In a previous synthesis, the key intermediate for construction of the [6,6] and [5,5] spiroketals was enone **2-3**, which was synthesized via a Ni/Cr-mediated coupling of **2-1** with **2-2** in an excellent overall yield (Figure 7). The best combination of protecting groups at C35, C41, and C48 was recently identified to be TES, TBS, and TES, respectively. During this transformation, three chiral centers were introduced at C38, C40, and C44, cf., **2-3** $\rightarrow$ **2-A** $\rightarrow$ **2-B** $\rightarrow$ **2-4**. Based on the synthetic work of calcimycin, the desired stereochemistry should be preferentially formed under a basic condition (see, e.g., Negri, D. P.; Kishi, Y. *Tetrahedron Lett.*, **1987**, 28, 1063). Indeed, this approach worked nicely for a synthesis of halichondrins A-Cs. However, an alternative route for the final transformation was desired.

[00663] Ketone **2-B** is available via an alternative, well-defined route. The Zr/Ni-mediated one-pot ketone synthesis showed a potential to meet these needs; specifically, this method was proved effective for coupling of (*S*)-**2-C** + (*S*)-**2-D** $\rightarrow$ (Σ,Σ)-**2-E**. The requisite ketone **2-B** could be synthesized from iodide **2-5** and 2-thiopyridine ester **2-6**. Ketone **2-B** could also be obtained via coupling at the C38-C39 bond, but we focused on the former route because of the overall synthetic efficiency of **2-5**. The feasibility of this disconnection was demonstrated with use of the combination of CH_2I at C40 with C(=O)SPy at C38. Py = 2-pyridyl.

[00664] Being encouraged with the successful ((*S*)-**2-C** + (*S*)-**2-D** $\rightarrow$ (Σ,Σ)-**2-E**)-coupling, the feasibility study for the proposed synthesis began. For this study, the right half **2-5** of halichondrin Bs was chosen. The C35-protecting group was selected for two reasons, i.e., (1) the rate of ketone coupling with **2-5** was significantly faster than that with the corresponding C35-TBS substrate and (2) deprotection of the C35-TES group in the following step was noticeably faster than that of the corresponding C35-TBS substrate. On the other hand, the left half **2-6** was chosen, because of its availability in a larger quantity at the time of

preliminary study. The C41-protecting group was chosen primarily for the ease of deprotection.

[00665] The desired product **2-7** was obtained in the first attempt under the conditions used for ((*S*)-**2-C** + (*S*)-**2-D** → (Σ,Σ)-**2-E**)-coupling. Conditions were then optimized for this case. First, Cp₂ZrCl₂ was important to accelerate the ketone coupling and, at the same time, suppress by-product formation via a (I→SPy)-displacement at C37. Second, a 5:1 mixture of DMI and EtOAc was found to be the best solvent. Third, the coupling proceeded well at 0.1 M concentration, although a higher concentration, for example 0.4 M, was better. Fourth, both Zn and Mn metals were effective. Fifth, 2,6-di-*tert*-butyl-4-methylpyridine was used to avoid partial deprotection of the TES groups during the reaction and/or workup. Lastly, as expected, the coupling efficiency depended on the molar ratio of **2-5** and **2-6**, for example 84% yield with **2-5:2-6** = 1.0:1.3; 62% with 1.0:1.0; 71% with 1.0:1.2.

[00666] Considering all these factors, the coupling condition specified in *Figure 7* is shown as an example procedure. For all the couplings, the molar ratio of **2-5:2-6** = 1.0:1.3 was used, considering the molecular size and complexity of **2-5** vs. **2-6**. Under this condition, the ketone coupling was carried out in 0.5-1.0 g scales, to furnish the desired product **2-7** in 80-90% yields.

[00667] In this coupling, three by-products were isolated in very small amounts (~3% yields). Spectroscopic analysis (¹H NMR, MS) suggested these by-products to be **2-8**, **2-9**, and **2-10**, respectively. The first two by-products were derived from **2-5**, formation of which was not surprising in light of the results discussed in the method-development work. The third by-product **2-10** was obviously derived from **2-6**, which was, as speculated, formed via a Ni-mediated decarbonylation, the transformation depicted in *Figure 7*.

[00668] Ketone **2-7** also served for a model study on the second stage of synthesis, *i.e.*, deprotection of the silyl groups, followed by acid-catalyzed [5,5]-spiroketal formation. As expected, the C50/C52-dioxasilinane group in **2-7** was readily removed on a treatment with HF•Py, to give the corresponding diol. A treatment of the resultant C49/C52-diol with TBAF (4 equiv.) buffered with pivalic acid (2 equiv.) gave the completely deprotected product within 6 hours, thereby confirming the ease of deprotection of the two TES group at C35 and C41. This transformation was also done in one step, *i.e.*, treatment directly with TBAF, buffered with pivalic acid.

[00669] The completely desilylated product was treated with an acid, to furnish **2-11**; namely, PPTS in CH₂Cl₂ at room temperature gave a ~5:1 mixture of **2-11** and its C38-*epi*-

11, which were separated by reverse phase, medium-pressure column chromatography, to furnish **2-11** (67% overall yield from **2-5**) and C38-*epi*-**2-11** (13% overall yield from **2-5**). With the method previously reported, C38-*epi*-**2-11** was isomerized to give additional **2-11** (9% isolated yield), thereby making the overall yield of **2-11** from **5** 76%. The structure of **2-11** was concluded from spectroscopic analysis; ^1H and ^{13}C NMR spectra were found beautifully to correspond to those of norhalichondrin B.

[00670] The results given in the previous section made a convincing case that the Zr/Ni-mediated one-pot ketone synthesis should lead to the development of a unified synthesis of the halichondrin class of natural products, and analogs thereof. To demonstrate experimentally, three types of right- and left-halves were prepared, respectively (*Figure 8A*). Combinations of these right- and left-halves should give all the nine halichondrins (*Figure 8B*).

[00671] The first stage in this approach was to apply Zr/Ni-mediated one-pot ketone synthesis for each combination. The ketone coupling was conducted under the previously defined condition, to furnish the expected products in 80-90% isolated yields. All the ketones were isolated by medium-pressured column chromatography (neutral silica gel) and fully characterized. The results were virtually identical with those found for **2-5** + **2-6** $\rightarrow$ **2-7**, including coupling rates, isolated yields, and detected by-products. For example, the (**2-5** + **2-14**)-coupling was carried out in a 200 mg scale of **2-5**, to give the expected, desired ketone in 88% isolated yield, along with three by-products **2-8**, **2-9**, and one corresponding to **2-11** in small amounts (~3%). Noteworthy, the C12 allyl group of halichondrins-C was found intact in the time-scale of ketone synthesis.

[00672] The second stage was deprotection of the silyl protecting groups, followed by [5,5]-spiroketal formation under acidic conditions. Halichondrin-B synthesis was first studied, where deprotection of the silyl groups and formation of the [5,5]-spiroketal were effected with TBAF buffered with pivalic acid in DMF and then PPTS in CH_2Cl_2 , to give a ~5:1 mixture of halichondrin B and its C38-epimer. Reverse-phase medium-pressure column chromatography was adopted for separation/isolation, to furnish halichondrin B and C38-epimer in an excellent overall yield; for example, 200 mg of **2-5** gave 133 mg (68%) and 25 mg (13%) of halichondrin B and C38-*epi*-halichondrin B, respectively. With the method previously reported, C38-*epi*-halichondrin B was isomerized to give additional 17 mg halichondrin B (9% isolated yield). Thus, the overall yield of halichondrin B was 77% from **2-5**. Spectroscopic comparison (HR-MS, ^1H and ^{13}C NMR) confirmed that halichondrin B

was identical to the authentic sample. The reproducibility of overall transformation was excellent and no potential issue was noticed for scaling.

[00673] Similarly, the synthesis of halichondrin A (**2-12** + **2-14** → **2-20**) was carried out. In this series, an additional step was required to remove the C12/C13 anisylidene, *i.e.*, PPTS treatment in a mixture of isopropanol and 2,2-dimethyl-1,3-propanediol. During the acid-treatment, the ratio of halichondrin A and its C38-epimer changed from ~5:1 down to ~3:1. As before, C38-epimer was isomerized, furnishing halichondrin A in 61% total yield from **2-12**. Spectroscopic comparison confirmed that halichondrin A was identical to the authentic sample.

[00674] The synthesis of halichondrin C (**2-13** + **2-14** → **2-23**) was also carried out. In this series, an additional step was required to remove the allyl group at C12, which was uneventfully achieved with the method used in the previous synthesis. Synthetic halichondrin C and C38-epimer were isolated in 55% and 11% yields, respectively. Spectroscopic comparison confirmed that halichondrin C was identical to the authentic sample. Noteworthy, attempted TMSOTf-induced isomerization in CH₂Cl₂ did not give halichondrin C. This phenomenon was observed for all the members in the halichondrin-C sub-group, but not for any member of other sub-groups, thereby indicating that the reason for the unsuccessful isomerization was due to the chemical property of halichondrin-C polycycle. Spectroscopic analysis of a product formed during the attempted reaction suggested a rearrangement of the halichondrin-C polycycle to a C12 ketal.

[00675] Synthesis in the norhalichondrin series proceeded equally well, although an extra step was required to hydrolyze the methyl ester at C53, which was achieved under the condition used in the previous work. It should be noted that, for synthesis of norhalichondrin C, base-induced hydrolysis of the methyl ester was done before deprotection of the allyl group, because of the base-instability of halichondrin-C polycycle. Spectroscopic comparison established that norhalichondrins A-C thus obtained were identical to the authentic samples.

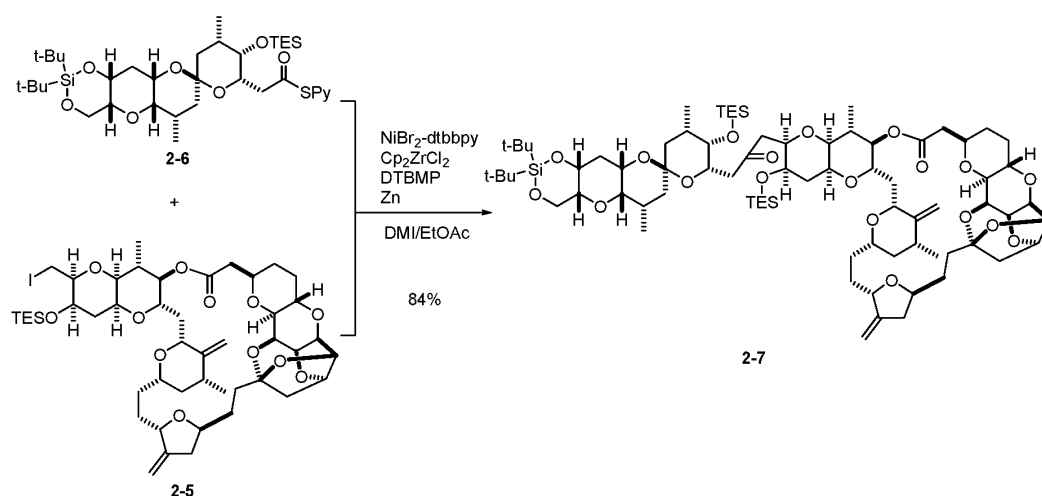
[00676] Lastly, the ketone route was applied to the homohalichondrin series. It is noteworthy that the previous enone route was not effective for a synthesis of homohalichondrins; it was successful only for homohalichondrin A, but with a very low efficiency (5% isolated yield). To our delight, the new synthetic route was found effective for a total synthesis of all the homohalichondrins; the overall efficiency in the homohalichondrin series was comparable to that in the halichondrin and norhalichondrin series. For instance, 100 mg **2-5** furnished 72 mg homohalichondrin B (75% overall yield). Spectroscopic comparison (HR-MS, ¹H and ¹³C

NMR) confirmed that homohalichondrins A-C were identical to the authentic samples. The reproducibility of overall transformation was excellent and no potential issue was noticed for scaling.

[00677] In summary, a unified, efficient, and scalable synthesis of the halichondrin class of natural products was completed. Newly developed Zr/Ni-mediated one-pot ketone synthesis was used for coupling of right halves with left halves, where Cp_2ZrCl_2 was found crucial to accelerate the coupling rate and, at the same time, suppress by-product formation.

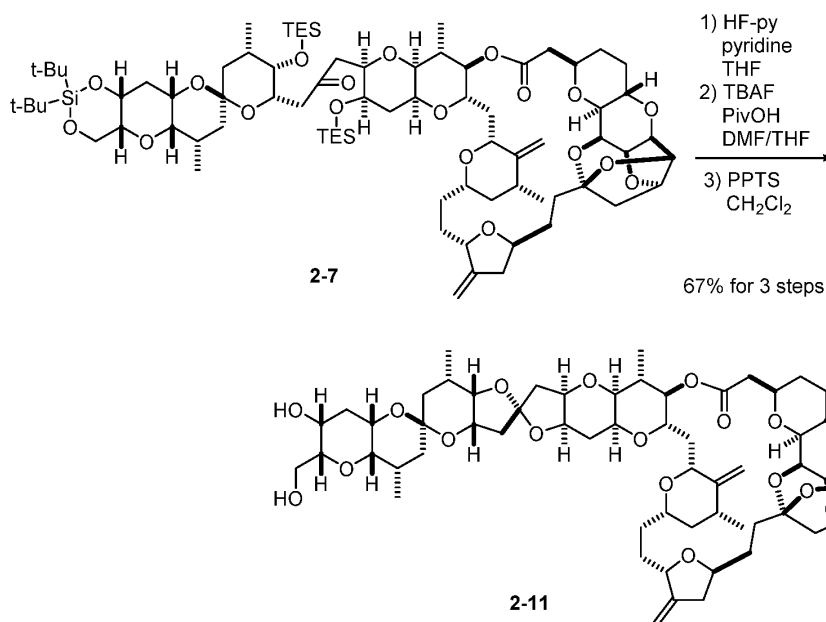
Halichondrins were obtained from these ketones basically in two operations, *i.e.*, desilylation and then [5,5]-spiroketal formation. Notably, the new synthetic route was successfully applied for a total synthesis of all the homohalichondrins. All the halichondrins thus synthesized were isolated as crystalline solids. We succeeded in growing a single crystal for an X-ray analysis for some of them; thus far, the analysis completed for halichondrin C, which was the first successful X-ray analysis of intact halichondrin. To demonstrate the scalability, halichondrin B was chosen, where 150 mg of halichondrin B (77% yield) was obtained from 200 mg of the right half **2-5**.

Experimental Procedures for the Synthesis of Halichondrins and Analogs



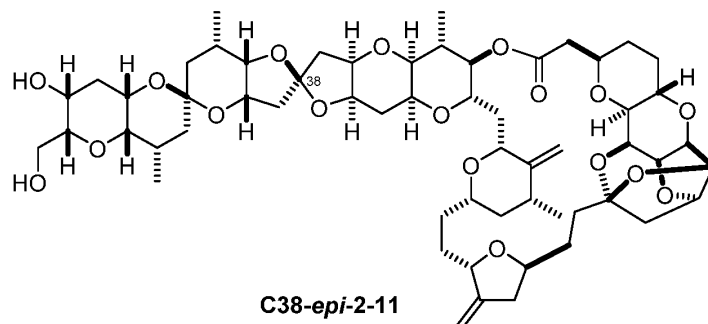
[00678] In a glove box, to a mixture of **2-5** (41.6 mg, 0.0424 mmol, 1 eq.), **2-6** (39 mg, 0.0551 mol, 1.3 eq.), DTBMP (21.8 mg, 0.106 mmol, 2.5 eq.), Zn (16.6 mg, 0.254 mmol, 6 eq.), and Cp_2ZrCl_2 (24.8 mg, 0.0848 mmol, 2 eq.) were added 5:1 mixture of DMI-EtOAc (0.2 mL) and $\text{NiBr}_2 \cdot \text{dtbbpy}$ (7.2 mg, 0.0148 mmol, 35 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction was removed from the glove box and quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer

was extracted with Et₂O. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by YAMAZEN purification system with neutral silica gel to give **2-7** (51.7 mg, 0.0356 mmol, 84%) as a colorless amorphous solid. (**2-7**): $[\alpha]_D^{20}$ -59.7 (*c* 1.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ: 5.20 (1H, s), 5.10 (1H, s), 4.92 (1H, s), 4.84-4.74 (3H, m), 4.68 (1H, d, *J* = 10.6 Hz), 4.52 (1H, ddd, *J* = 10.0, 10.0, 4.1 Hz), 4.35 (1H, m), 4.27 (1H, m), 4.21 (1H, d, *J* = 12.3 Hz), 4.17-4.06 (4H, m), 4.03-3.94 (4H, m), 3.89 (1H, dd, *J* = 6.5, 4.7 Hz), 3.84-3.70 (3H, m), 3.64 (1H, dd, *J* = 6.5, 4.1 Hz), 3.45 (1H, ddd, *J* = 4.7, 4.7, 4.7 Hz), 3.33 (1H, s), 3.19 (1H, dd, *J* = 16.4, 10.0 Hz), 3.14 (1H, dd, *J* = 5.3, 4.1 Hz), 3.07-2.95 (3H, m), 2.84-2.72 (3H, m), 2.61 (1H, dd, *J* = 9.4, 1.8 Hz), 2.45-2.02 (15H, m), 2.02-1.90 (2H, m), 1.83 (1H, m), 1.79-1.66 (6H, m), 1.59 (1H, ddd, *J* = 14.1, 4.7, 4.7 Hz), 1.56-1.37 (6H, m), 1.37-1.27 (10H, m), 1.17 (3H, d, *J* = 7.0 Hz), 1.13 (9H, s), 1.10-1.02 (22H, m), 1.00 (3H, d, *J* = 6.5 Hz), 0.96 (3H, d, *J* = 6.5 Hz), 0.72-0.62 (12H, m) ppm. ¹³C NMR (125 MHz, C₆D₆) δ: 206.9, 171.3, 153.0, 152.7, 110.0, 15.0, 103.7, 97.2, 82.4, 81.0, 78.3, 78.0, 77.8, 77.7, 77.6, 76.9, 76.2, 75.5, 74.8, 74.7 (x2), 74.2, 74.0, 73.8, 73.2, 70.4, 69.3, 68.6, 68.5, 67.3, 66.0, 64.7, 63.8, 48.6, 46.7, 46.3, 43.9, 41.3, 39.5, 39.2, 38.5, 37.7, 36.8, 36.6, 36.3, 35.5, 35.3, 32.5, 31.1, 30.7, 30.6, 30.4, 29.5, 29.1, 27.9, 27.7, 23.4, 21.0, 18.6, 18.1, 17.4, 16.4, 7.5, 7.3, 6.0, 5.3 ppm. IR (film): 2955, 2933, 2875, 1723, 1371, 1133, 1097, 1084, 1017 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₇₈H₁₂₈NaO₁₉Si₃, 1475.8250; found, 1475.8251.



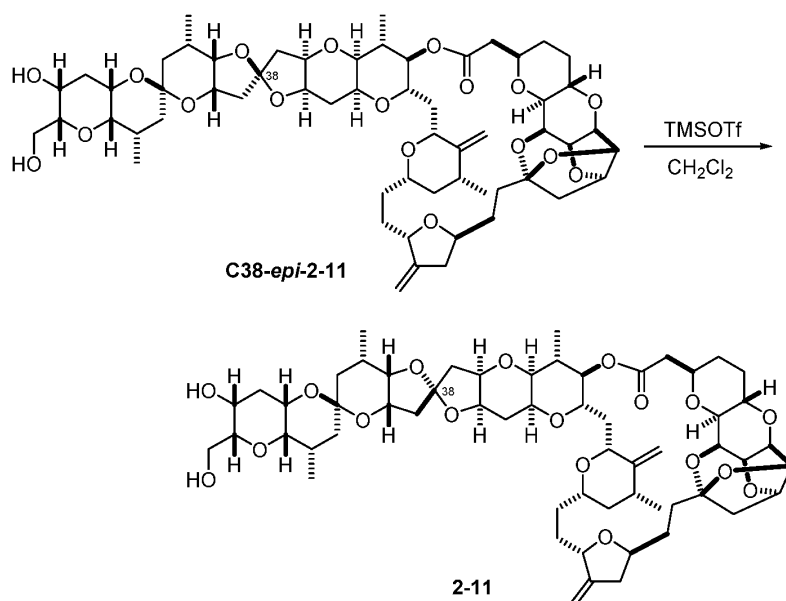
[00679] To a stirred solution of **2-7** (108 mg, 0.0743 mmol, 1 equiv.) in dry THF (7.5 mL, 0.01M) in a plastic tube was added pyridine-buffered pyridinium hydrofluoride solution (0.16 mL, 20 equiv.; freshly prepared from 0.20 mL of pyridinium hydrofluoride available from Aldrich, 0.60 mL of pyridine) at 0 °C. After being stirred for 2 hours at the same temperature, the reaction was quenched with sat. aq. NaHCO₃ until gas evolution stopped. The aqueous layer was extracted with EtOAc. The combined organic phase was washed with brine, dried over Na₂SO₄, and concentrated under reduced pressure. The crude material was used for the next step without further purification. To a stirred solution of crude diol (calculated as 0.0743 mmol, 1 equiv.) in DMF (3.7 mL, 0.02M) was added the buffered TBAF solution (0.37 mL, 5 equiv., freshly prepared by 0.74 mL TBAF solution (1 M in THF) and 38 mg PivOH) at room temperature. After being stirred for 4 h at the same temperature, CaCO₃ (2.0 g) and DOWEX 50WX8-400 (2.0 g) were added. After being stirred for 1 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of the crude tetraol (calculated as 0.0743 mmol, 1 eq.) in CH₂Cl₂ (3.7 mL, 0.02M) was added PPTS (93.3 mg, 0.371mmol, 5 equiv.) at room temperature. After being stirred for 2.5 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (100% EtOAc, then 9% MeOH in EtOAc) to give a crude **2-11** with its C38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give **2-11** (53.1 mg,

0.0498 mmol, 67% for 3 steps) as a white solid and **C38-*epi*-2-11** (10.2 mg, 0.0096 mmol, 13% for 3 steps) as a white solid. (**2-11**): $[\alpha]_D^{20}$ -62.0 (*c* 0.30, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.06 (1H, s), 5.02 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.70 (1H, dd, $J = 4.5, 4.5$ Hz), 4.63 (1H, dd, $J = 7.8, 4.8$ Hz), 4.60 (1H, dd, $J = 4.2, 4.2$ Hz), 4.45 (1H, d, $J = 12.6$ Hz), 4.33 (1H, ddd, $J = 9.6, 9.6, 4.2$ Hz), 4.30 (1H, m), 4.25-4.23 (1H, m), 4.18 (1H, dd, $J = 6.6, 4.8$ Hz), 4.13-4.06 (4H, m), 3.99 (1H, d, $J = 2.4$ Hz), 3.90-3.86 (2H, m), 3.81 (1H, s), 3.72-3.69 (3H, m), 3.61 (1H, d, $J = 10.8$ Hz), 3.41 (1H, dd, $J = 6.0, 6.6$ Hz), 3.22 (1H, ddd, $J = 6.6, 4.8, 4.8$ Hz), 2.98 (1H, dd, $J = 10.4, 1.5$ Hz), 2.82-2.79 (1H, m), 2.56 (1H, dd, $J = 17.4, 3.6$ Hz), 2.45 (1H, dd, $J = 17.4, 1.8$ Hz), 2.40 (1H, dd, $J = 13.2, 6.0$ Hz), 2.38-2.25 (6H, m), 2.22-2.16 (3H, m), 2.11-1.97 (9H, m), 1.94-1.90 (3H, m), 1.86-1.80 (3H, m), 1.74-1.67 (3H, m), 1.60 (1H, ddd, $J = 12.0, 12.0, 6.0$ Hz), 1.51-1.29 (9H, m), 1.11 (3H, d, $J = 7.8$ Hz), 1.06 (3H, d, $J = 7.8$ Hz), 1.05-0.99 (1H, m), 0.95 (3H, d, $J = 7.2$ Hz), 0.94 (3H, d, $J = 7.2$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.8, 153.3, 153.2, 114.8, 111.2, 105.7, 104.7, 98.6, 83.8, 82.4, 81.4, 80.6, 79.1, 78.0, 78.0, 77.9, 77.3, 77.3, 77.2, 76.3, 76.1, 75.8, 75.3, 75.0, 75.0, 74.9, 73.8, 72.7, 69.6, 68.5, 66.3, 65.7, 63.2, 49.4, 45.5, 44.9, 44.8, 41.2, 39.7, 38.2, 38.1, 37.8, 37.4, 37.2, 35.8, 35.4, 33.0, 31.8, 31.2, 31.0, 30.8, 30.1, 29.4, 27.3, 18.4, 18.1, 17.4, 15.8 ppm. FTIR (film): 3476, 2956, 2918, 2850, 1733, 1668, 1589, 1433, 1207, 1134, 1097, 1021 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{58}\text{H}_{82}\text{O}_{18}\text{Na}$, 1089.5393; found, 1089.5378.



[00680] **C38-*epi*-2-11**: $[\alpha]_D^{20}$ -68.3 (*c* 0.20, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.04 (1H, s), 5.00 (1H, s), 4.87 (1H, s), 4.80 (1H, s), 4.72 (1H, dd, $J = 12.0, 7.2$ Hz), 4.70 (1H, dd, $J = 6.0, 5.4$ Hz), 4.60 (1H, dd, $J = 5.4, 5.4$ Hz), 4.43 (1H, d, $J = 12.0$ Hz), 4.36 (1H, ddd, $J = 12.0, 12.0, 4.8$ Hz), 4.27 (1H, m), 4.18-4.05 (6H, m), 4.10 (1H, dd, $J = 5.4, 1.8$ Hz), 3.91-3.84 (3H, m), 3.78 (1H, s), 3.70-3.60 (4H, m), 3.57 (1H, d, $J = 13.8$ Hz), 3.42 (1H, dd, $J = 7.8, 6.6$ Hz), 3.33 (1H, d, $J = 2.4$ Hz), 3.32-3.31 (2H, m), 3.16 (1H, dd, $J = 10.6, 7.6$ Hz), 2.99 (1H, d, $J = 11.4$ Hz), 2.84-2.79 (1H, m), 2.55 (1H, dd, $J = 20.7, 10.5$ Hz), 2.45 (1H, dd, $J = 20.7, 2.4$ Hz), 2.35-1.90 (20H, m), 1.86-1.70 (3H, m), 1.74-1.51 (5H, m), 1.51-1.29 (9H, m), 1.10 (3H, d, $J = 7.8$ Hz), 1.03 (3H, d, $J = 8.4$ Hz), 1.05-0.99 (1H, m), 1.01 (3H, d, $J = 7.8$ Hz), 1.00

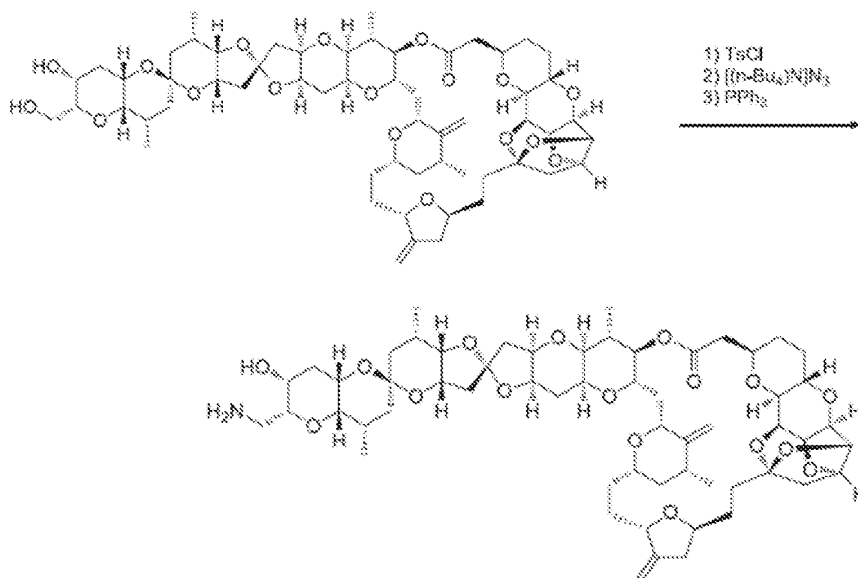
(3H, d, $J = 7.8$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.8, 153.3, 152.8, 115.6, 111.3, 105.1, 104.7, 98.4, 83.8, 82.4, 81.5, 79.8, 79.2, 79.0, 78.9, 78.4, 77.9, 77.9, 77.0, 76.5, 76.1, 76.1, 76.0, 75.2, 75.2, 75.0, 74.7, 73.2, 73.2, 69.5, 68.5, 68.3, 66.3, 63.2, 49.5, 45.5, 45.0, 44.8, 41.2, 39.6, 38.7, 38.2, 38.2, 37.5, 37.4, 37.2, 35.4, 35.3, 34.6, 33.3, 31.8, 31.3, 31.0, 30.7, 30.1, 29.2, 27.0, 18.4, 18.3, 17.4, 15.2 ppm. FTIR (film): 3465, 2960, 2918, 2850, 1735, 1668, 1590, 1433, 1210, 1134, 1097, 1022 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{58}\text{H}_{82}\text{O}_{18}\text{Na}$, 1089.5393; found, 1089.5367. **C38-*epi*-2-11** was epimerized to **2-11** by the following procedure:



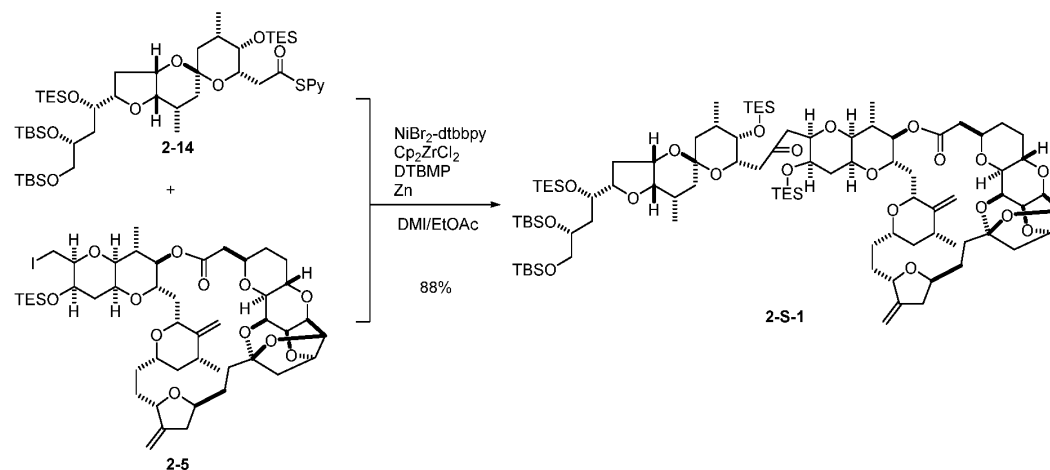
[00681] To a solution of **C38-*epi*-2-11** (10.1 mg, 0.0095 mmol, 1 eq.) in CH_2Cl_2 (4.7 mL) was added TMSOTf (95 μL , 0.525 mmol, excess) at -78°C . After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. After being stirred for 1 h at 0°C , the organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H_2O to 100% MeCN) to give **2-11** (6.9 mg, 0.0065 mmol, 68%) as a white solid.

[00682] An exemplary reaction sequence converting Compound (2) to Compound (1) is shown below in *Scheme 3*. Exemplary experimental procedures are provided below.

Scheme 3

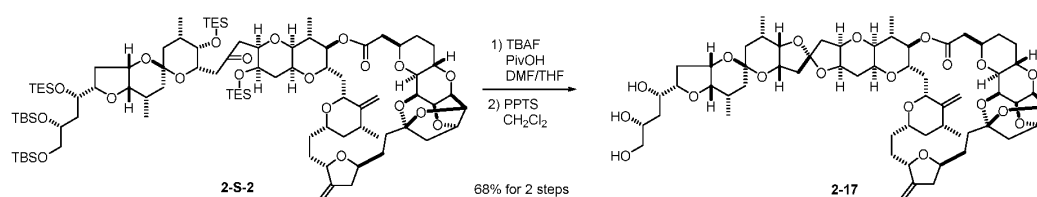


Halichondrin B (17)



[00683] In a glove box, to a solution of iodide **2-5** (200 mg, 0.203 mmol, 1 eq.) and thioester **2-14** (252.5 mg, 0.264 mmol, 1.3 equiv.) in DMI (1.7 mL) and EtOAc (0.34 mL) were added DTBMP (167 mg, 0.816 mmol, 4 eq.), Zn powder (80.0 mg, 1.22 mmol, 6 eq.), Cp_2ZrCl_2 (178.4 mg, 0.612 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (29.7 mg, 0.062 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was

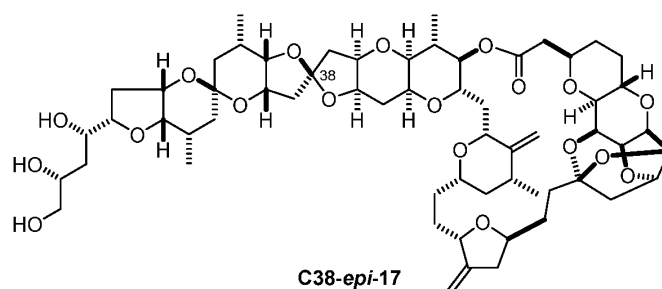
separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by YAMAZEN purification system on neutral silica gel to give ketone **2-S-1** (303 mg, 0.178 mmol, 88%) as a white amorphous solid. (**2-S-1**): [α]_D²⁰ -58.3 (*c* 1.20, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 5.21 (1H, s), 5.11 (1H, s), 4.94 (1H, s), 4.85 (1H, d, *J* = 7.2 Hz), 4.81-4.78 (2H, m), 4.69 (1H, d, *J* = 10.2 Hz), 4.54-4.51 (2H, m), 4.36 (1H, d, *J* = 7.8 Hz), 4.27 (1H, s), 4.24 (1H, m), 4.18-4.13 (2H, m), 4.10-4.07 (2H, m), 3.93-3.88 (2H, m), 3.83-3.81 (3H, m), 3.78-3.75 (2H, m), 3.63 (1H, dd, *J* = 6.0, 4.2 Hz), 3.44 (2H, m), 3.33 (1H, s), 3.19 (1H, dd, *J* = 16.2, 10.2 Hz), 3.16 (1H, d, *J* = 5.4 Hz), 3.11-3.02 (2H, m), 2.78 (1H, dd, *J* = 16.8, 7.2 Hz), 2.60 (1H, d, *J* = 9.6 Hz), 2.49-2.43 (1H, m), 2.41-2.31 (5H, m), 2.28-2.24 (3H, m), 2.19-1.96 (10H, m), 1.93 (1H, d, *J* = 13.2 Hz), 1.87-1.64 (7H, m), 1.61 (1H, ddd, *J* = 15.0, 4.8, 4.8 Hz), 1.56-1.43 (7H, m), 1.40 (1H, dd, *J* = 13.2, 4.8 Hz), 1.33 (1H, dd, *J* = 9.6, 9.6 Hz), 1.18 (3H, d, *J* = 6.6 Hz), 1.15 (3H, d, *J* = 7.2 Hz), 1.12-1.04 (27H, m), 1.10 (9H, s), 1.04 (9H, s), 1.00 (3H, d, *J* = 6.6 Hz), 0.96 (3H, d, *J* = 6.0 Hz), 0.78 (6H, q, *J* = 8.0 Hz), 0.69-0.65 (12H, m), 0.28 (6H, s), 0.150 (3H, s), 0.148 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 206.8, 171.3, 153.0, 152.7, 110.0, 104.9, 103.8, 97.0, 82.4, 81.5, 81.0, 80.4, 78.4, 78.1, 77.6, 76.9, 75.5, 74.9, 74.7, 74.1, 74.0, 73.8, 72.9, 72.0, 71.8, 71.5, 70.5, 69.9, 68.4, 68.3, 65.9, 64.6, 48.6, 46.8, 46.3, 43.9, 41.3, 39.5, 39.3, 38.5, 38.2, 37.8, 36.4, 35.5, 35.4, 35.3, 32.5, 31.3, 30.7, 30.6, 29.0, 26.6, 26.3($\times 6$), 26.3($\times 6$), 18.7, 18.6, 18.5, 18.4, 18.1, 16.4, 7.4($\times 6$), 7.4($\times 6$), 7.3($\times 6$), 6.0($\times 3$), 5.7($\times 3$), 5.3($\times 3$), -4.0, -4.2, -5.1, -5.2 ppm. FTIR (film): 3450, 2936, 2864, 1734, 1642, 1547, 1147, 1112, 1055, 1021, 997 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₉₀H₁₅₈O₂₀Si₅Na, 1722.0085; found, 1722.0061.



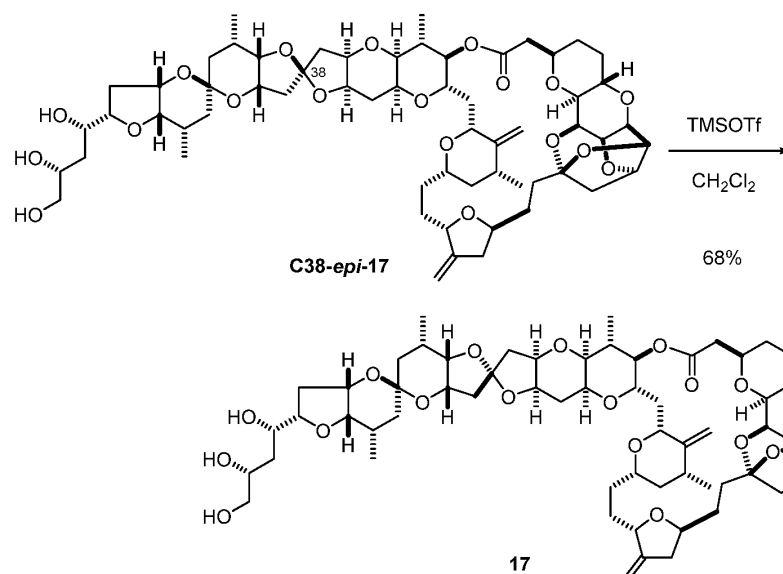
[00684] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 3.52 mL of 1 M in THF, 3.52 mmol, 10 eq.) and PivOH (180 mg, 1.76 mmol, 5 eq.). To a stirred solution of **2-S-1** (303 mg, 0.178 mmol, 1 equiv.) in DMF (8.8 mL) was added the buffered TBAF solution at room temperature. After being stirred for 4 h at the same temperature, CaCO₃ (6.0 g) and DOWEX 50WX8-400 (6.0 g) were added. After being stirred for 2 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through

a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude material, which was used in the next step without further purification. To a stirred solution of the crude material (calculated as 0.178 mmol, 1 eq.) in CH₂Cl₂ (17.6 mL) was added PPTS (221.8 mg, 0.882 mmol, 5 eq.) at room temperature. After being stirred for 4 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (100% EtOAc, then 9% MeOH in EtOAc) to give a crude **2-17** with its C38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give halichondrin B **17** (133.0 mg, 0.120 mmol, 68% for 2 steps) as a white crystalline solid and **C38-epi-17** (25.0 mg, 0.0225 mmol, 13% for 2 steps) as a white solid.

Halichondrin B (17): $[\alpha]_D^{20}$ -62.3 (*c* 1.00, MeOH). MP: 164-166 °C (recrystallized from Hexanes-CH₂Cl₂). ¹H NMR (600 MHz, CD₃OD) δ : 5.07 (1H, d, *J* = 1.8 Hz), 5.02 (1H, d, *J* = 1.8 Hz), 4.89 (1H, s), 4.81 (1H, s), 4.70 (1H, dd, *J* = 4.8, 3.6 Hz), 4.63 (1H, dd, *J* = 7.2, 4.8 Hz), 4.60 (1H, dd, *J* = 4.2, 4.2 Hz), 4.45 (1H, d, *J* = 10.8 Hz), 4.33 (1H, ddd, *J* = 9.6, 9.6, 4.2 Hz), 4.30 (1H, m), 4.25-4.23 (1H, m), 4.18 (1H, dd, *J* = 6.6, 4.2 Hz), 4.13-4.05 (6H, m), 3.99 (1H, ddd, *J* = 9.6, 4.8, 4.8 Hz), 3.90-3.85 (3H, m), 3.71 (1H, dd, *J* = 10.2, 10.2 Hz), 3.70 (1H, m), 3.61 (1H, d, *J* = 7.6 Hz), 3.56 (1H, s), 3.53 (1H, dd, *J* = 10.4, 4.2 Hz), 3.47 (1H, dd, *J* = 10.8, 6.0 Hz), 3.22 (1H, dd, *J* = 6.6, 4.8 Hz), 2.98 (1H, dd, *J* = 9.6, 2.4 Hz), 2.82-2.78 (1H, m), 2.56 (1H, dd, *J* = 17.4, 9.6 Hz), 2.45 (1H, dd, *J* = 17.4, 2.4 Hz), 2.39 (1H, dd, *J* = 13.8, 5.7 Hz), 2.38-2.22 (7H, m), 2.22-2.16 (2H, m), 2.09-1.97 (7H, m), 1.86-1.81 (3H, m), 1.77-1.67 (4H, m), 1.62-1.58 (2H, m), 1.57-1.29 (9H, m), 1.10 (3H, d, *J* = 6.6 Hz), 1.06 (3H, d, *J* = 6.6 Hz), 1.05-0.99 (1H, m), 1.02 (3H, d, *J* = 6.6 Hz), 0.97 (3H, d, *J* = 6.6 Hz) ppm. ¹³C NMR (125 MHz, CD₃OD) δ : 172.8, 153.3, 153.2, 114.8, 111.3, 105.7, 104.8, 98.4, 83.8, 82.4, 81.3, 81.3, 80.7, 79.1, 78.1, 77.9, 77.9, 77.4, 77.2, 76.3, 76.1, 75.8, 75.4, 75.0, 75.0, 74.9, 73.7, 73.3, 73.1, 73.0, 71.6, 69.6, 67.2, 65.7, 49.4, 45.5, 44.9, 44.9, 41.2, 39.7, 37.9, 37.9, 37.8, 37.5, 37.5, 37.2, 36.3, 35.8, 33.0, 31.8, 31.3, 31.0, 30.8, 29.4, 27.1, 27.1, 18.4, 18.3, 18.1, 15.8 ppm. FTIR (film): 3460, 2936, 2864, 1736, 1642, 1557, 1167, 1122, 1105, 1054, 1041, 1021, 997 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₆₀H₈₆O₁₉Na, 1133.5656; found, 1133.5651.

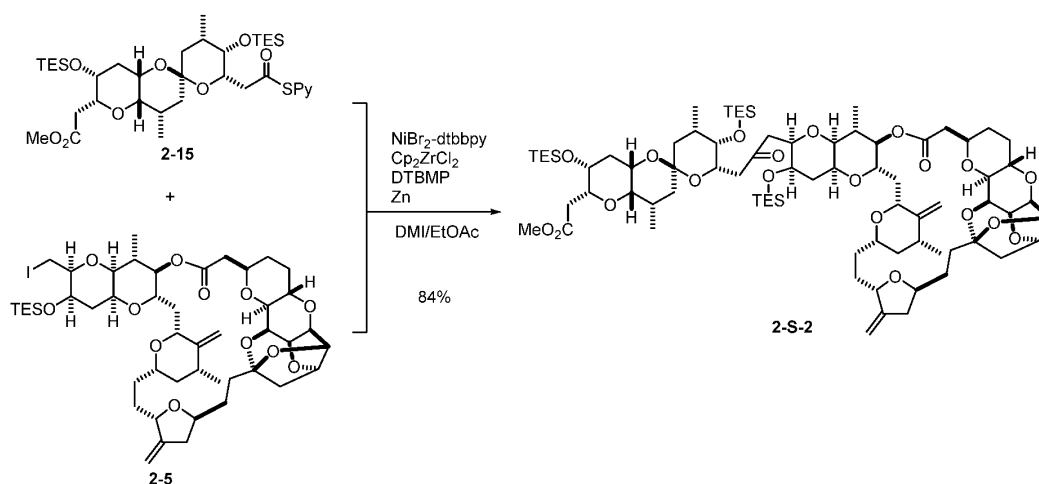


[00685] C38-*epi*-Halichondrin B: $[\alpha]_D^{20}$ -66.0 (*c* 1.00, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.04 (1H, s), 5.00 (1H, s), 4.87 (1H, s), 4.81 (1H, s), 4.72 (1H, dd, $J = 10.2, 6.6$ Hz), 4.70 (1H, dd, $J = 4.2, 4.2$ Hz), 4.60 (1H, dd, $J = 4.8, 4.8$ Hz), 4.43 (1H, d, $J = 10.8$ Hz), 4.37 (1H, ddd, $J = 12.0, 12.0, 4.8$ Hz), 4.27 (1H, m), 4.19-4.06 (8H, m), 3.99 (1H, ddd, $J = 9.6, 5.4, 4.2$ Hz), 3.91-3.82 (4H, m), 3.78 (1H, ddd, $J = 14.4, 4.8, 4.2$ Hz), 3.64-3.56 (3H, m), 3.53 (1H, dd, $J = 11.4, 4.5$ Hz), 3.46 (1H, dd, $J = 11.4, 6.0$ Hz), 3.34 (2H, m), 3.17 (1H, dd, $J = 8.7, 6.3$ Hz), 2.99 (1H, dd, $J = 9.6, 1.8$ Hz), 2.84-2.79 (1H, m), 2.55 (1H, dd, $J = 16.8, 8.4$ Hz), 2.47 (1H, dd, $J = 16.8, 2.4$ Hz), 2.35-1.93 (20H, m), 1.86-1.82 (2H, m), 1.79-1.70 (5H, m), 1.67-1.33 (12H, m), 1.10 (3H, d, $J = 6.6$ Hz), 1.04 (3H, d, $J = 8.4$ Hz), 1.05-0.99 (1H, m), 1.02 (3H, d, $J = 7.8$ Hz), 1.00 (3H, d, $J = 6.6$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.8, 153.3, 152.8, 115.5, 111.3, 105.1, 104.7, 98.2, 83.8, 82.4, 81.3, 81.1, 79.9, 79.2, 78.9, 78.9, 78.4, 77.9, 77.9, 76.5, 76.1, 76.1, 76.0, 75.2, 75.2, 74.7, 73.5, 73.3, 73.1, 73.0, 71.7, 69.5, 68.2, 67.1, 49.9, 45.6, 45.0, 44.7, 41.2, 39.6, 38.3, 38.2, 38.1, 37.5, 37.5, 37.2, 36.2, 35.4, 33.3, 31.8, 31.3, 30.9, 30.5, 30.2, 29.3, 27.1, 26.8, 18.4, 18.3, 15.2 ppm. FTIR (film): 3460, 2936, 2864, 1736, 1642, 1557, 1167, 1122, 1105, 1054, 1041, 1021, 997 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{60}\text{H}_{86}\text{O}_{19}\text{Na}$, 1133.5656; found, 1133.5651. **C38-*epi*-17** was epimerized to halichondrin B (**17**) by the following procedure:



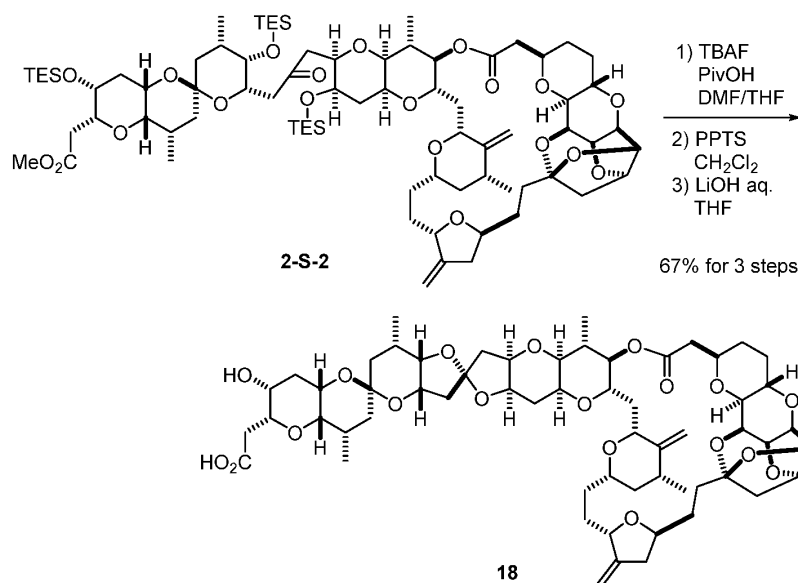
[00686] To a solution of **C38-*epi*-17** (25.0 mg, 0.0225 mmol, 1 eq.) in CH₂Cl₂ (11.2 mL) was added TMSOTf (0.225 mL, 0.719 mmol, excess) at -78 °C. After being stirred for 15 min, the reaction was quenched with sat. NaHCO₃ aq. After being stirred for 1 h at 0 °C, the organic layer was separated and the aqueous layer was extracted with CH₂Cl₂. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give halichondrin B (**17**) (17.1 mg, 0.0154 mmol, 68%) as a colorless solid.

Norhalichondrin B (18)



[00687] In a glove box, to a solution of iodide **2-5** (100 mg, 0.102 mmol, 1 eq.) and thioester **2-16** (95.5 mg, 0.132 mmol, 1.3 eq.) in DMI (0.85 mL) and EtOAc (0.17 mL) were added

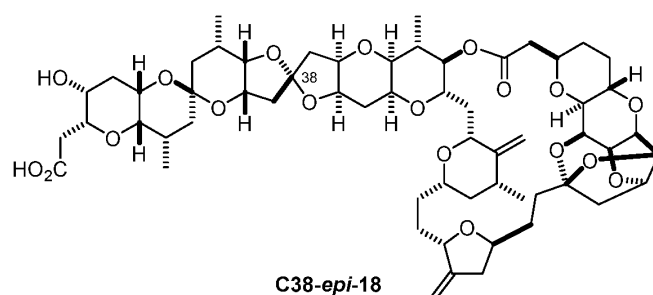
DTBMP (83.8 mg, 0.408 mmol, 4 eq.), Zn powder (40.0 mg, 0.612 mmol, 6 eq.), Cp_2ZrCl_2 (89.4 mg, 0.306 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (14.9 mg, 0.0306 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et_2O and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with Et_2O . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 15%, 25% EtOAc in Hexanes) to give ketone **2-S-2** (125.7 mg, 0.0856 mmol, 84%) as a colorless amorphous solid. (**2-S-2**): $[\alpha]_D^{20}$ -68.4 (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 5.21 (1H, s), 5.11 (1H, s), 4.94 (1H, s), 4.85 (1H, t, $J = 6.6$ Hz), 4.81-4.77 (2H, m), 4.69 (1H, d, $J = 10.2$ Hz), 4.52 (1H, ddd, $J = 9.8, 9.8, 4.2$ Hz), 4.36 (1H, d, $J = 9.6$ Hz), 4.27 (1H, s), 4.14 (1H, dd, $J = 4.2, 4.2$ Hz), 4.11-4.06 (2H, m), 4.03-3.97 (3H, m), 3.89 (1H, dd, $J = 5.7, 5.7$ Hz), 3.84-3.72 (4H, m), 3.78-3.68 (5H, m), 3.64 (1H, dd, $J = 6.3, 3.9$ Hz), 3.59 (1H, brs), 3.45 (1H, q, $J = 4.0$ Hz), 3.38 (3H, s), 3.37 (1H, s), 3.20-3.14 (2H, m), 3.13 (1H, s), 3.07 (1H, dd, $J = 17.8, 6.0$ Hz), 2.99 (1H, dd, $J = 17.8, 6.0$ Hz), 2.84 (1H, dd, $J = 14.4, 7.8$ Hz), 2.81-2.75 (2H, m), 2.61 (1H, d, $J = 10.2$ Hz), 2.58 (1H, dd, $J = 14.8, 5.4$ Hz), 2.42-2.21 (7H, m), 2.21-2.06 (5H, m), 1.99 (1H, dd, $J = 12.6, 12.6$ Hz), 1.93 (1H, d, $J = 13.2$ Hz), 1.89-1.82 (1H, m), 1.79-1.64 (3H, m), 1.62-1.30 (9H, m), 1.18 (3H, d, $J = 6.6$ Hz), 1.12-1.04 (31H, m), 1.01 (3H, d, $J = 6.6$ Hz), 0.98 (3H, d, $J = 6.6$ Hz), 0.70-0.61 (18H, m) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.9, 171.7, 171.3, 153.0, 152.6, 110.0, 104.9, 103.7, 96.9, 82.4, 81.0, 78.3, 78.0, 77.7, 77.2, 76.9, 76.5, 76.1, 75.5, 74.8, 74.70, 74.67, 74.1, 74.0, 73.8, 73.0, 70.3, 69.6, 68.4, 65.9, 65.7, 64.6, 64.5, 50.9, 48.6, 46.8, 46.3, 43.9, 41.3, 39.5, 39.3, 38.6, 37.5, 36.3, 35.5, 35.4, 32.5, 31.1, 31.0, 30.7, 30.6, 29.2, 29.0, 26.2, 18.6, 18.1, 17.2, 16.4, 7.4, 7.28, 7.25, 6.0, 5.4, 5.3 ppm; FTIR (film): 2954, 2921, 2876, 1737, 1458, 1436, 1372, 1287, 1262, 1239, 1207, 1187, 1154, 1073, 740, 728 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{78}\text{H}_{128}\text{O}_{20}\text{Si}_3\text{Na}$, 1491.8204; found, 1491.8181.



[00688] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.86 mL of 1 M in THF, 0.86 mmol, 10 eq.) and PivOH (43.9 mg, 0.430 mmol, 5 eq.). To a stirred solution of ketone **2-S-2** (125.7 mg, 0.0856 mmol, 1 eq.) in DMF (4.3 mL) was added the buffered TBAF solution at room temperature. After being stirred for 6 h at the same temperature, CaCO₃ (2.4 g) and DOWEX 50WX8-400 (2.4 g) were added after diluting with 10 mL EtOAc. After being stirred for 1 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of the crude tetraol (calculated as 0.0856 mmol, 1 eq.) in CH₂Cl₂ (8.5 mL) was added PPTS (86.4 mg, 0.344 mmol, 4 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (CH₂Cl₂ then 25%, 50%, 75%, then 100% EtOAc in Hexanes then 2% MeOH in EtOAc) to give a crude Norhalichondrin B methyl ester with its C38 epimer. The compound was used in the next step after concentration without further purification.

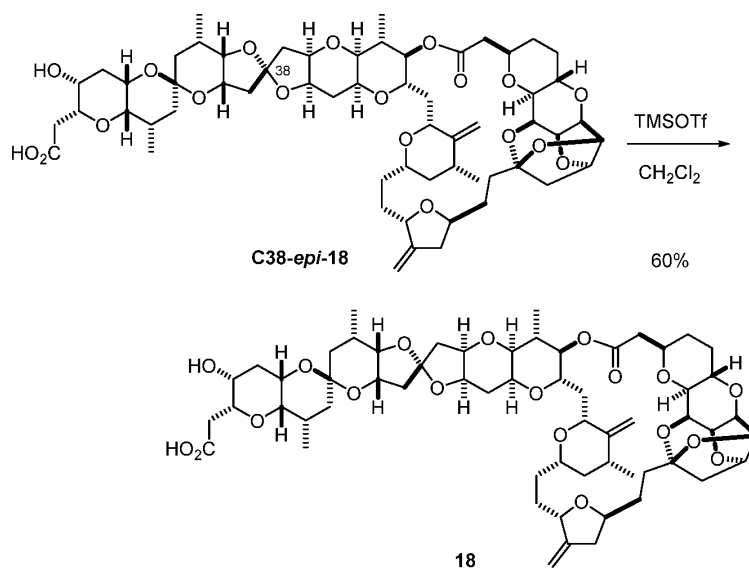
To a stirred solution of the crude methyl ester (calculated as 0.0856 mmol, 1 eq.) in THF (10 mL) was added 1M LiOH aq. (3.3 mL) at room temperature.³ After being stirred for 2 h at the same temperature, the reaction mixture was diluted with 6.6 mL of water. The THF was then removed from the mixture by evaporator. After the reaction was cooled down to 0 °C, 1 M HCl aq. (3.3 mL) was added and the reaction mixture was allowed for further 2 min stirring. The resulting mixture was extracted by EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The resulting mixture was

purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give Norhalichondrin B (**18**) (62.4 mg, 0.0570 mmol, 67% for 3 steps) as a colorless solid and 38-*epi*-Norhalichondrin B (**C38-*epi*-18**) (8.4 mg, 0.0077 mmol, 9% for 3 steps) as a colorless solid. **Norhalichondrin B (18)**: $[\alpha]_D^{20}$ -54.6 (*c* 1.00, MeOH).. ¹H NMR (600 MHz, CD₃OD) δ : 5.06 (1H, d, *J* = 1.5 Hz), 5.01 (1H, d, *J* = 1.5 Hz), 4.88 (1H, s), 4.81 (1H, d, *J* = 1.5 Hz), 4.70 (1H, t, *J* = 4.0 Hz), 4.63 (1H, dd, *J* = 7.8, 4.7 Hz), 4.60 (1H, t, *J* = 4.0 Hz), 4.45 (1H, d, *J* = 9.6 Hz), 4.32 (1H, td, *J* = 10.2, 4.6 Hz), 4.31–4.29 (1H, m), 4.24 (1H, ddd, *J* = 11.2, 4.2, 1.8 Hz), 4.18 (1H, dd, *J* = 6.6, 4.8 Hz), 4.14–4.09 (3H, m), 4.07 (1H, dd, *J* = 9.6, 9.3 Hz), 3.99 (1H, dd, *J* = 5.8, 2.4 Hz), 3.91–3.85 (2H, m), 3.82–3.78 (2H, m), 3.74–3.69 (2H, m), 3.61 (1H, d, *J* = 10.4 Hz), 3.59–3.56 (1H, m), 3.30 (1H, m), 3.22 (1H, dd, *J* = 6.6, 5.1 Hz), 2.98 (1H, dd, *J* = 9.6, 1.8 Hz), 2.81 (1H, ddd, *J* = 16.0, 8.0, 2.1 Hz), 2.59 (1H, dd, *J* = 15.0, 7.8 Hz), 2.57–2.52 (2H, m), 2.45 (1H, dd, *J* = 17.6, 1.8 Hz), 2.40 (1H, dd, *J* = 13.2, 6.2 Hz), 2.34–2.32 (2H, m), 2.32–2.24 (4H, m), 2.21–2.15 (3H, m), 2.13–1.93 (8H, m), 1.87–1.79 (2H, m), 1.76–1.71 (3H, m), 1.70–1.66 (1H, m), 1.64–1.57 (1H, m), 1.56–1.47 (4H, m), 1.46–1.29 (5H, m), 1.10 (3H, d, *J* = 6.6 Hz), 1.06 (3H, d, *J* = 7.0 Hz), 1.02 (1H, d, *J* = 12.0 Hz), 0.98 (3H, d, *J* = 7.2 Hz), 0.95 (3H, d, *J* = 7.2 Hz) ppm. ¹³C NMR (150 MHz, CD₃OD) δ : 172.8 (2C), 153.3, 153.2, 114.7, 111.2, 105.6, 104.8, 98.5, 83.8, 82.4, 80.6, 79.0, 78.1, 77.90, 77.85, 77.76, 77.4, 77.23, 77.18, 76.3, 76.1, 75.8, 75.4, 75.02, 74.98, 74.9, 73.7, 72.7, 69.6, 68.0, 67.8, 65.8, 49.4, 45.4, 44.9, 44.7, 41.2, 39.8, 38.22, 38.20, 38.0, 37.8, 37.5, 37.2, 35.7, 35.5, 33.0, 31.8, 31.3, 31.0, 30.8, 30.0, 29.4, 27.3, 18.4, 18.1, 17.3, 15.8 ppm. FTIR (film): 3480, 2926, 2873, 2853, 1736, 1676, 1565, 1395, 1334, 1265, 1207, 1188, 1152, 1134, 1118, 1086, 1072, 1045, 1020, 996 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₅₉H₈₂O₁₉Na, 1117.5348; found 1117.5292.



[00689] 38-*epi*-Norhalichondrin B (C38-*epi*-18): $[\alpha]_D^{20}$ -69.7 (*c* 0.400, MeOH). ¹H NMR (600 MHz, CD₃OD) δ : 5.04 (1H, d, *J* = 1.5 Hz), 4.96 (1H, d, *J* = 1.5 Hz), 4.87 (1H, d, *J* = 1.5 Hz), 4.80 (1H, s), 4.74–4.68 (2H, m), 4.60 (1H, t, *J* = 4.5 Hz), 4.43 (1H, d, *J* = 9.6 Hz), 4.37

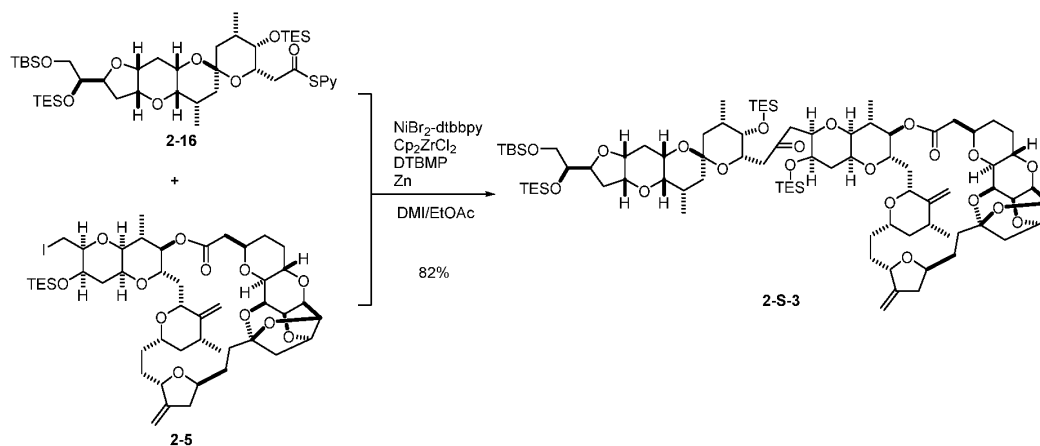
(1H, td, $J = 10.2, 4.6$ Hz), 4.30–4.25 (1H, m), 4.20–4.04 (4H, m), 4.01 (1H, dd, $J = 5.8, 2.4$ Hz), 3.91–3.83 (2H, m), 3.80 (1H, t, $J = 7.8$ Hz), 3.75 (1H, brs), 3.65–3.55 (2H, m), 3.34 (1H, m), 3.17 (1H, dd, $J = 6.6, 5.0$ Hz), 2.99 (1H, dd, $J = 9.6, 1.8$ Hz), 2.82 (1H, ddd, $J = 16.0, 8.0, 2.1$ Hz), 2.63–2.51 (2H, m), 2.48 (1H, dd, $J = 15.0, 7.8$ Hz), 2.36–2.24 (4H, m), 2.23–2.13 (4H, m), 2.13–1.92 (5H, m), 2.47 (1H, dd, $J = 17.6, 1.8$ Hz), 2.40 (1H, dd, $J = 13.2, 6.2$ Hz), 2.34–2.32 (2H, m), 2.32–2.24 (4H, m), 2.21–2.15 (2H, m), 2.13–1.93 (6H, m), 1.82 (1H, td, $J = 12.0, 2.0$ Hz), 1.77 (1H, d, $J = 12.0$ Hz), 1.72 (1H, d, $J = 12.0$ Hz), 1.69–1.60 (2H, m), 1.59–1.47 (3H, m), 1.47–1.34 (4H, m), 1.10 (3H, d, $J = 6.6$ Hz), 1.06 (1H, d, $J = 12.0$ Hz), 1.04 (3H, d, $J = 7.0$ Hz), 1.00 (3H, d, $J = 7.2$ Hz), 0.97 (3H, d, $J = 7.2$ Hz) ppm. ^{13}C NMR (150 MHz, CD_3OD) δ : 172.9 (2C), 153.3, 152.9, 115.6, 111.4, 105.1, 104.7, 98.4, 83.8, 82.4, 79.9, 79.2, 79.0, 78.4, 78.0, 77.9, 77.1, 76.5, 76.1, 76.0, 75.2, 74.8, 73.3, 73.2, 69.5, 68.3, 68.0, 67.9, 45.6, 45.0, 44.7, 41.2, 38.9, 38.7, 38.3, 38.2, 37.5, 37.2, 35.5, 33.3, 31.8, 31.3, 31.0, 30.2, 30.0, 29.3, 27.0, 18.4, 18.3, 17.4, 15.2 ppm. FTIR (film): HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{59}\text{H}_{82}\text{O}_{19}\text{Na}$, 117.5348; found 117.5292. **C38-*epi*-18** was epimerized to Norhalichondrin B (**18**) by the following procedure:



[00690] To a solution of **C38-*epi*-18** (8.4 mg, 0.0077 mmol, 1 eq.) in CH_2Cl_2 (3 mL) was added TMSOTf (0.07 mL, 0.385 mmol, excess) at -78°C . After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. After being stirred for 1 h at 0°C , the organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with

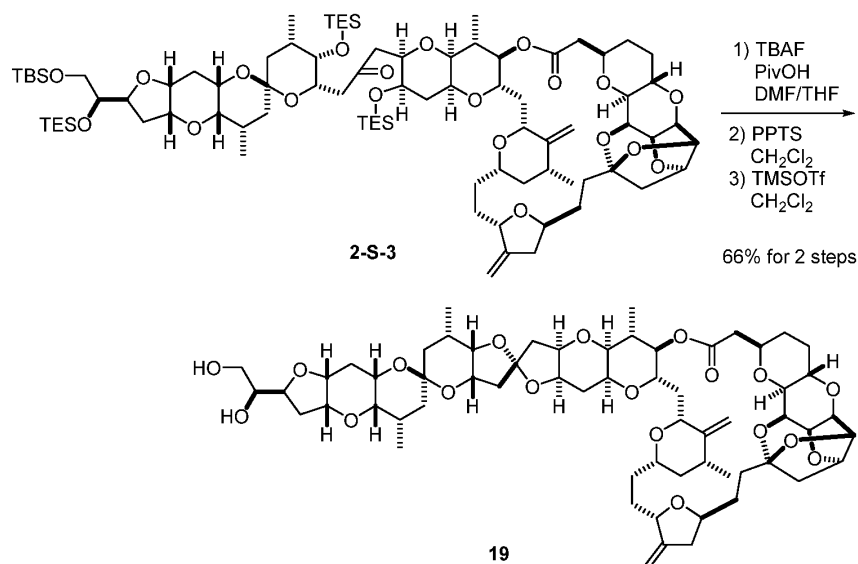
ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give Norhalichondrin B (**18**) (5.0 mg, 0.0046 mmol, 60%) as a colorless solid.

Homohalichondrin B (19)



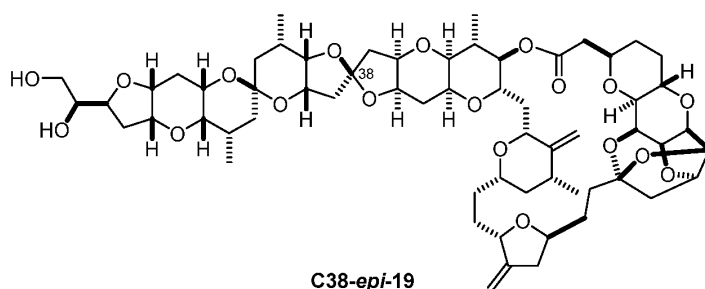
[00691] In a glove box, to a solution of iodide **2-5** (103 mg, 0.105 mmol, 1 eq.) and thioester **2-16** (113 mg, 0.132 mmol, 1.3 eq.) in DMI (0.85 mL) and EtOAc (0.17 mL) were added DTBMP (83.8 mg, 0.408 mmol, 4 eq.), Zn powder (40.0 mg, 0.612 mmol, 6 eq.), Cp₂ZrCl₂ (89.4 mg, 0.306 mmol, 3 eq.), and NiBr₂-dtbbpy (14.9 mg, 0.0306 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et₂O and sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with Et₂O. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 9%, 17% EtOAc in Hexanes) to give ketone **2-S-3** (137 mg, 0.0857 mmol, 82%) as a colorless amorphous solid. In a preliminary study, the coupling reaction of iodide **5** (25.0 mg, 0.0254 mmol) and thioester **16** (30.0 mg, 0.0352 mmol) gave desired ketone (35.7 mg 0.0223 mmol) in 88% yield. (**2-S-3**): [α]_D²⁰ -50.8 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 5.21 (1H, s), 5.11 (1H, s), 4.93 (1H, s), 4.85-4.78 (3H, m), 4.69 (1H, d, *J* = 10.2 Hz), 4.52 (1H, ddd, *J* = 9.8, 9.8, 4.2 Hz), 4.47 (1H, ddd, *J* = 10.4, 5.3, 5.3 Hz), 4.34 (1H, d, *J* = 9.6 Hz), 4.27 (1H, s), 4.14 (1H, dd, *J* = 4.2, 4.2 Hz), 4.11-4.07 (2H, m), 4.02-4.00 (3H, m), 3.89 (1H, dd, *J* = 5.7, 5.7 Hz), 3.84-3.80 (3H, m), 3.78-3.68 (5H, m), 3.64 (1H, dd, *J* = 6.3, 3.9 Hz), 3.45 (1H, d, *J* = 3.6 Hz), 3.29 (1H, s), 3.20-3.16 (2H, m), 3.06 (1H, dd, *J* = 17.6, 5.7 Hz), 3.01 (1H, dd, *J* = 17.6, 6.9 Hz), 2.93 (1H, s), 2.80 (1H, d, *J* = 7.8 Hz), 2.77 (1H, d, *J* = 7.2 Hz), 2.61 (1H, d, *J* = 10.2 Hz), 2.46 (1H, d, *J* = 15.0 Hz), 2.39-2.22 (9H, m), 2.20-2.05 (6H, m), 1.98 (1H, dd, *J* =

12.6, 12.6 Hz), 1.93 (1H, d, $J = 13.2$ Hz), 1.87-1.84 (2H, m), 1.75-1.65 (4H, m), 1.62-1.30 (11H, m), 1.18 (3H, d, $J = 6.6$ Hz), 1.12-1.04 (30H, m), 1.04-1.03 (12H, m), 0.96 (3H, d, $J = 6.6$ Hz), 0.77 (6H, q, $J = 8.0$ Hz), 0.69-0.64 (12H, m), 0.14 (3H, s), 0.13 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.7, 171.3, 153.0, 152.6, 110.0, 104.9, 103.7, 96.9, 82.4, 81.0, 79.1, 78.3, 78.0, 77.9, 77.7, 76.9, 76.8, 76.1, 75.5, 74.8, 74.7, 74.2, 74.1, 74.0, 73.8, 73.6, 73.0, 70.4, 69.3, 68.4, 66.1, 65.9, 64.7, 63.7, 48.6, 46.8, 46.3, 43.9, 41.3, 39.5, 39.3, 38.6, 37.7, 37.6, 36.5, 36.3, 35.5, 35.3, 32.5, 31.6, 31.1, 30.7, 30.65, 30.60, 29.5, 29.0, 26.2, 18.7, 18.6, 18.1, 17.6, 16.4, 7.5, 7.34, 7.28, 6.0, 5.7, 5.3, -5.1, -5.3 ppm. FTIR (film): 2953, 2927, 2875, 1720, 1459, 1086, 1015, 834, 725 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{85}\text{H}_{144}\text{O}_{20}\text{Si}_4\text{Na}$, 1619.9220; found, 1619.9298.



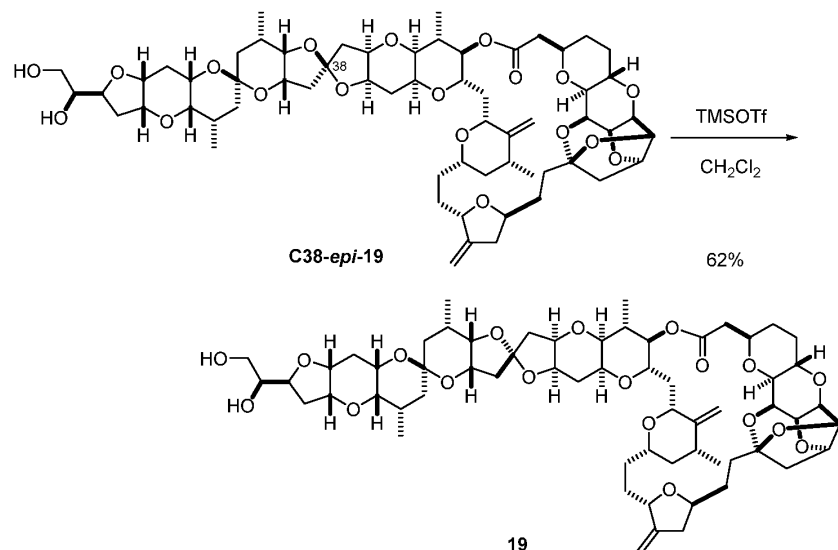
[00692] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.86 mL of 1 M in THF, 0.86 mmol, 10 eq.) and PivOH (43.9 mg, 0.430 mmol, 5 eq.). To a stirred solution of ketone **2-S-3** (137 mg, 0.0857 mmol, 1 eq.) in DMF (4.3 mL) was added the buffered TBAF solution at room temperature. After being stirred for 7 h at the same temperature, CaCO_3 (2.4 g) and DOWEX 50WX8-400 (2.4 g) were added. After being stirred for 1 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of the crude tetraol (calculated as 0.0857 mmol, 1 eq.) in CH_2Cl_2 (8.6 mL) was added PPTS (108 mg, 0.430 mmol, 5 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was

directly subjected to column chromatography on amino silica gel (CH_2Cl_2 then 25%, 50%, 75%, then 100% EtOAc in Hexanes then 2% MeOH in EtOAc) to give a crude Homohalichondrin B with its C38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H_2O to 100% MeCN) to give Homohalichondrin B (**19**) (63.8 mg, 0.0568 mmol, 66% for 2 steps) as a colorless solid and 38-*epi*-Homohalichondrin B (**C38-*epi*-19**) (14.4 mg, 0.0128 mmol, 15% for 2 steps) as a colorless solid. **Homohalichondrin B (19)**: $[\alpha]_D^{20}$ -43.7 (c 1.02, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.06 (1H, s), 5.01 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.70 (1H, dd, J = 4.5, 4.5 Hz), 4.63 (1H, dd, J = 7.8, 4.8 Hz), 4.60 (1H, dd, J = 4.5, 4.5 Hz), 4.45 (1H, d, J = 10.8 Hz), 4.33 (1H, ddd, J = 9.6, 9.6, 4.2 Hz), 4.30 (1H, s), 4.25-4.21 (2H, m), 4.18 (1H, dd, J = 5.7, 5.7 Hz), 4.06-4.13 (4H, m), 4.02 (1H, s), 3.95 (1H, s), 3.87-3.88 (3H, m), 3.71 (1H, dd, J = 10.2, 10.2 Hz), 3.66 (1H, s), 3.61 (1H, d, J = 10.8 Hz), 3.55-3.60 (3H, m), 3.50 (1H, ddd, J = 5.4, 5.4, 5.4 Hz), 3.21 (1H, dd, J = 5.7, 5.7 Hz), 3.12 (1H, s), 2.98 (1H, d, J = 10.2 Hz), 2.80 (1H, dd, J = 9.0, 6.6 Hz), 2.56 (1H, dd, J = 17.4, 9.6 Hz), 2.45 (1H, d, J = 17.4 Hz), 2.39 (1H, dd, J = 13.2, 5.4 Hz), 2.38-2.24 (6H, m), 2.22-2.13 (4H, m), 2.09-1.97 (9H, m), 1.90 (1H, ddd, J = 15.6, 4.2, 4.2 Hz), 1.80-1.84 (2H, m), 1.74-1.67 (3H, m), 1.60 (1H, ddd, J = 12.0, 12.0, 6.0 Hz), 1.51-1.29 (9H, m), 1.10 (3H, d, J = 6.6 Hz), 1.05 (3H, d, J = 7.2 Hz), 1.05-0.99 (1H, m), 0.95 (3H, d, J = 6.6 Hz), 0.94 (3H, d, J = 5.4 Hz) ppm. ^{13}C NMR (125 MHz, CDCl_3) δ : 171.2, 151.8, 151.6, 112.4, 110.1, 104.4, 104.1, 96.6, 82.2, 81.1, 79.8, 78.4, 77.7, 77.6, 76.6, 76.29, 76.25, 75.4, 75.3, 75.1, 74.9, 74.8, 74.4, 73.9, 73.7, 73.5, 72.8, 71.9, 71.2, 70.8, 68.2, 66.7, 65.7, 63.7, 48.4, 43.4, 42.5, 40.4, 38.7, 37.3, 37.0, 36.92, 36.87, 36.6, 36.0, 34.4, 32.1, 31.4, 30.7, 30.1, 29.4, 29.0, 28.9, 28.2, 25.8, 18.0, 17.8, 17.1, 15.0 ppm. FTIR (film): 3460, 2926, 2874, 1736, 1652, 1567, 1187, 1132, 1105, 1074, 1041, 1021, 997 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{61}\text{H}_{86}\text{O}_{19}\text{Na}$, 1145.5656; found, 1145.5770.



[00693] 38-*epi*-Homohalichondrin B (C38-*epi*-19): $[\alpha]_D^{20}$ -86.6 (c 0.860, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.04 (1H, d, J = 1.8 Hz), 4.96 (1H, d, J = 1.8 Hz), 4.87 (1H, s), 4.80 (1H, s), 4.72 (1H, dd, J = 10.2, 6.6 Hz), 4.70 (1H, dd, J = 4.2, 4.2 Hz), 4.60 (1H, dd, J = 4.5,

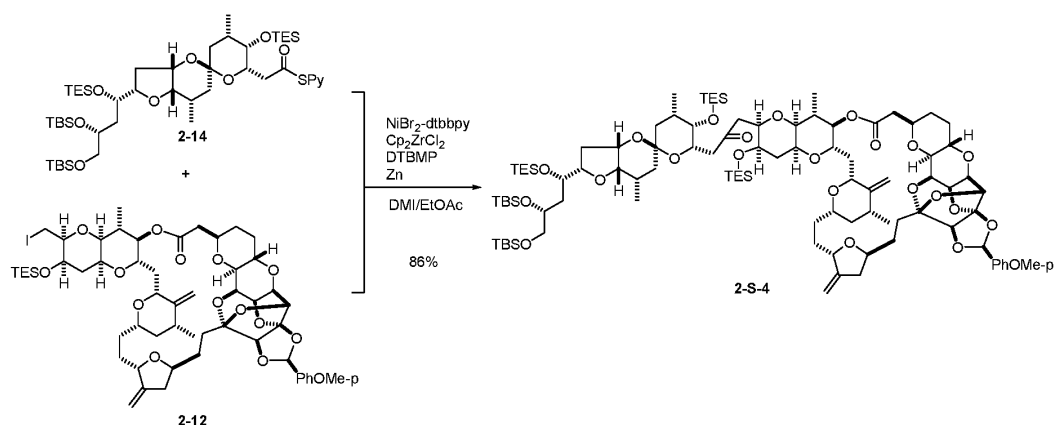
4.5 Hz), 4.44 (1H, d, $J = 10.8$ Hz), 4.37 (1H, ddd, $J = 10.2, 10.2, 4.2$ Hz), 4.28 (1H, d, $J = 2.4$ Hz), 4.23 (1H, ddd, $J = 9.6, 4.8, 4.8$ Hz), 4.19-4.05 (6H, m), 4.02 (1H, s), 3.98 (1H, dd, $J = 4.8, 1.8$ Hz), 3.91-3.85 (3H, m), 3.81 (1H, s), 3.63 (1H, dd, $J = 10.5, 10.5$ Hz), 3.59-3.56 (3H, m), 3.54 (1H, d, $J = 2.4$ Hz), 3.51-3.47 (1H, m), 3.17 (1H, dd, $J = 8.4, 6.0$ Hz), 3.15 (1H, s), 2.99 (1H, dd, $J = 9.6, 1.2$ Hz), 2.83-2.79 (1H, m), 2.56 (1H, dd, $J = 17.0, 8.7$ Hz), 2.47 (1H, dd, $J = 17.0, 2.4$ Hz), 2.35-1.93 (21H, m), 1.91 (1H, ddd, $J = 15.6, 4.5, 4.5$ Hz), 1.83 (1H, ddd, $J = 11.1, 11.1, 2.4$ Hz), 1.77 (1H, d, $J = 13.2$ Hz), 1.71 (1H, dd, $J = 13.2, 2.4$ Hz), 1.68-1.60 (2H, m), 1.58-1.54 (1H, m), 1.50-1.33 (7H, m), 1.29 (1H, dd, $J = 12.6, 4.2$ Hz), 1.10 (3H, d, $J = 7.2$ Hz), 1.05-0.98 (1H, m), 1.004 (3H, d, $J = 7.2$ Hz), 0.995 (3H, d, $J = 6.6$ Hz), 0.96 (3H, d, $J = 7.2$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.9, 153.3, 152.8, 115.6, 111.3, 105.1, 104.7, 98.0, 83.8, 82.4, 80.2, 79.8, 79.2, 78.9, 78.5, 78.4, 77.9, 77.8, 76.5, 76.12, 76.09, 76.0, 75.7, 75.2, 75.1, 74.8, 74.4, 73.2, 72.9, 69.5, 68.3, 65.3, 65.1, 45.6, 45.0, 44.7, 41.2, 39.6, 38.6, 38.2, 38.1, 37.5, 37.2, 35.4, 33.3, 31.9, 31.8, 31.3, 30.9, 30.19, 30.16, 29.3, 26.8, 18.4, 17.7, 15.2 pm. FTIR (film): 3487, 2925, 2872, 1737, 1188, 1119, 1074, 1019, 996, 896, 735 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{61}\text{H}_{86}\text{O}_{19}\text{Na}$, 1145.5656; found, 1145.5631. **C38-*epi*-19** was epimerized to Homohalichondrin B (**19**) by the following procedure:



[00694] To a solution of **C38-*epi*-19** (14.4 mg, 0.0128 mmol, 1 eq.) in CH_2Cl_2 (6.4 mL) was added TMSOTf (0.13 mL, 0.719 mmol, excess) at -78°C . After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. After being stirred for 1 h at 0°C , the organic layer was separated and the aqueous layer was extracted with

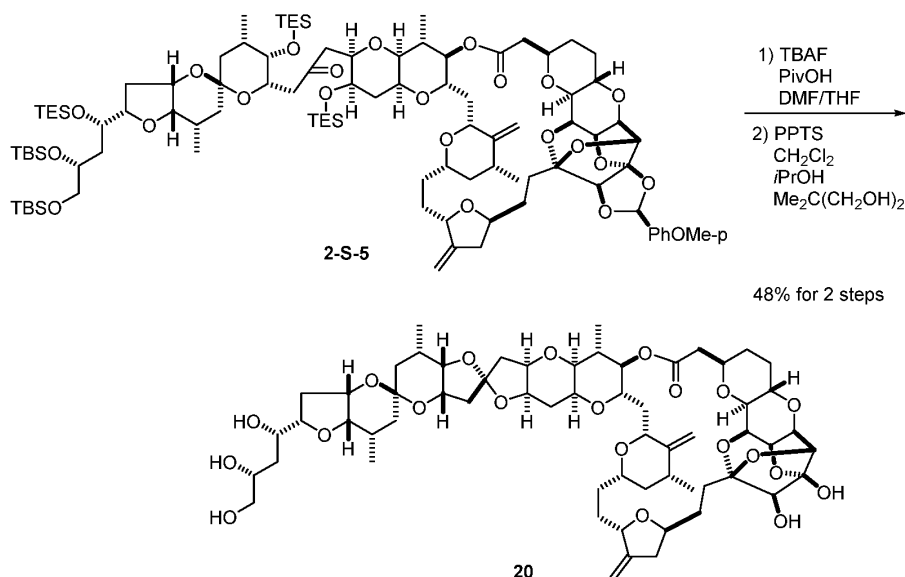
CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H_2O to 100% MeCN) to give Homohalichondrin B (**19**) (8.9 mg, 0.00792 mmol, 62%) as a colorless solid.

Halichondrin A (20)



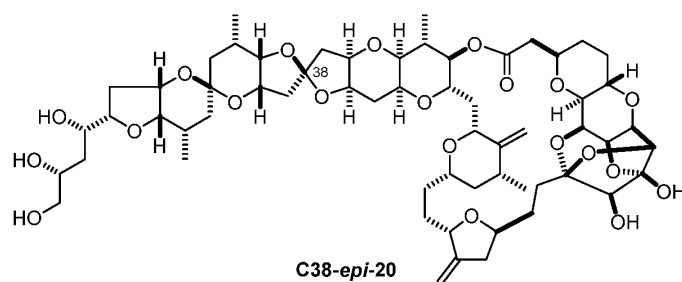
[00695] In a glove box, to a solution of iodide **2-12** (100.0 mg, 0.0883 mmol, 1 eq.) and thioester **2-14** (109.5 mg, 0.115 mmol, 1.3 eq.) in DMI (0.75 mL) and EtOAc (0.15 mL) were added DTBMP (72.5 mg, 0.353 mmol, 4 eq.), Zn powder (34.6 mg, 0.529 mmol, 6 eq.), Cp_2ZrCl_2 (77.4 mg, 0.265 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (12.9 mg, 0.027 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by YAMAZEN purification system on neutral silica gel (0%, 9%, then 20% EtOAc in Hexanes) to give ketone **2-S-4** (140.0 mg, 0.0756 mmol, 86%) as a colorless amorphous solid. (**2-S-4**): $[\alpha]_D^{20}$ -64.8 (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.35 (2H, d, $J = 8.4$ Hz), 6.72 (2H, d, $J = 8.4$ Hz), 6.07 (1H, s), 5.19 (1H, s), 5.08 (1H, s), 4.94 (1H, s), 4.84-4.79 (3H, m), 4.66 (1H, d, $J = 10.8$ Hz), 4.51-4.46 (2H, m), 4.39 (1H, br s), 4.35 (1H, dd, $J = 5.4, 1.2$ Hz), 4.33 (1H, dd, $J = 8.4, 1.2$ Hz), 4.23-4.21 (1H, m), 4.14-4.11 (1H, m), 3.83-3.80 (3H, m), 4.05 (1H, s), 4.03-3.95 (4H, m), 3.91-3.87 (1H, m), 3.82-3.79 (3H, m), 3.76-3.71 (3H, m), 3.46 (1H, dd, $J = 8.4, 4.8$ Hz), 3.41 (1H, s), 3.31 (1H, s), 3.24 (3H, s), 3.18-3.13 (2H, m), 3.07-2.99 (2H, m), 2.76 (1H, dd, $J = 16.8, 4.8$ Hz), 2.74-2.70 (1H, m), 2.52 (1H, d, $J = 9.6$ Hz), 2.47-2.41 (1H, m), 2.37-2.29 (6H, m), 2.27-2.18 (5H, m), 2.13-1.92 (11H, m), 1.83-1.46 (12H, m), 1.38-1.27 (1H, m), 1.18 (3H, d, $J = 6.6$ Hz),

1.12 (3H, d, $J = 7.2$ Hz), 1.09 (9H, t, $J = 7.2$ Hz), 1.08 (9H, s), 1.07-1.03 (21H, m), 1.02 (9H, s), 0.94 (3H, d, $J = 7.2$ Hz), 0.76 (6H, q, $J = 7.8$ Hz), 0.69-0.64 (12H, m), 0.26 (6H, s), 0.13 (3H, s), 0.13 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ 206.7, 171.1, 161.4, 152.9, 152.7, 119.0, 114.1, 109.3, 109.0, 105.0, 103.8, 97.0, 90.2, 83.8, 81.5, 80.3, 78.4, 78.1, 77.9, 76.0, 76.0, 75.5, 74.8, 74.7, 74.2, 74.0, 73.6, 72.9, 72.0, 71.8, 71.5, 70.1, 69.8, 68.3, 68.3, 65.9, 64.6, 54.8, 46.8, 46.2, 44.0, 41.3, 39.5, 38.7, 38.2, 37.7, 36.4, 35.4, 35.4, 32.4, 31.0, 30.9, 30.7, 30.3, 27.6, 26.6, 26.3($\times 6$), 26.3($\times 6$), 18.7, 18.6, 18.5, 18.4, 18.1, 16.4, 7.4($\times 6$), 7.3($\times 6$), 7.3($\times 3$), 7.3($\times 3$), 6.0($\times 6$), 5.7($\times 6$), 5.3($\times 6$) ppm. FTIR (film): 2955, 2917, 2876, 1736, 1648, 1519, 1253, 1096, 1032, 1009, 851 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{98}\text{H}_{164}\text{O}_{23}\text{Si}_5\text{Na}$, 1849.0510; found, 1849.0490.

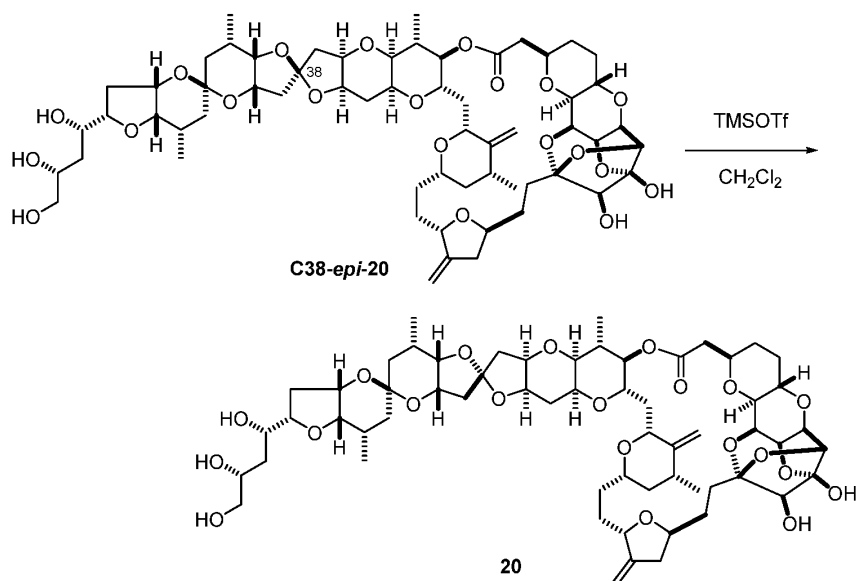


[00696] To a stirred solution of **2-S-5** (138 mg, 0.0735 mmol, 1 equiv.) in DMF (3.7 mL, 0.02M) was added the buffered TBAF solution (0.74 mL, 10 equiv., freshly prepared by 1.48 mL TBAF solution (1 M in THF) and 75 mg PivOH) at room temperature. After being stirred for 3 h at the same temperature, CaCO_3 (3.0 g) and DOWEX 50WX8-400 (3.0 g) were added.¹ After being stirred for 2 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude material, which was used in the next step without further purification. To a stirred solution of the crude material (calculated as 0.0735 mmol, 1 eq.) in CH_2Cl_2 (3.7 mL, 0.02M) was added PPTS (184.6 mg, 0.735 mmol, 10 equiv.) at room temperature. After being stirred for 1.5 h at the same temperature, TLC analysis indicated the disappearance of starting material. $i\text{PrOH}$ (1.2

mL) and additional PPTS (184.6 mg, 0.735 mmol, 10 eq.) were added to the resulted solution at the same temperature. After being stirred for 20 h, the reaction mixture was directly subjected to column chromatography on amino silica gel (CH₂Cl₂ then 100% EtOAc then 16% MeOH in EtOAc) to give a crude halichondrin A with its C-38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 50% MeCN in H₂O) to give halichondrin A (**20**) (40.0 mg, 0.035 mmol, 48% for 3 steps) as a white crystalline solid and 38-*epi*-halichondrin A (**C38-*epi*-20**) (13.5 mg, 0.0118 mmol, 16% for 3 steps) as a white solid. **Halichondrin A (20)**: $[\alpha]_D^{20}$ -73.2 (*c* 0.11, MeOH). MP: 168-170 °C (recrystallized from Hexanes-CH₂Cl₂) ¹H NMR (600 MHz, CD₃OD) δ : 5.08 (1H, s), 5.03 (1H, s), 4.88 (1H, s), 4.82 (1H, s), 4.62 (1H, dd, *J* = 7.2, 4.2 Hz), 4.45 (1H, d, *J* = 11.2 Hz), 4.37 (1H, dd, *J* = 4.8, 3.0 Hz), 4.32 (1H, ddd, *J* = 10.0, 10.0, 4.2 Hz), 4.32–4.28 (2H, m), 4.25 (1H, ddd, *J* = 11.2, 4.4, 2.4 Hz), 4.20 (1H, dd, *J* = 3.2, 2.1 Hz), 4.14–4.07 (4H, m), 4.05 (1H, ddd, *J* = 2.4, 2.4, 2.4 Hz), 3.99 (1H, ddd, *J* = 9.6, 4.8, 4.2 Hz), 3.91–3.84 (3H, m), 3.78 (1H, ddd, *J* = 8.8, 4.8, 4.4 Hz), 3.75–3.70 (1H, m), 3.69 (1H, dd, *J* = 2.3, 2.3 Hz), 3.61 (1H, d, *J* = 11.7 Hz), 3.56 (1H, dd, *J* = 2.3, 1.8 Hz), 3.53 (1H, s), 3.53 (1H, dd, *J* = 11.2, 4.7, Hz), 3.47 (1H, dd, *J* = 11.2, 6.5 Hz), 3.22 (1H, dd, *J* = 6.5, 4.7 Hz), 2.94 (1H, dd, *J* = 10.0, 2.3 Hz), 2.82 (1H, dddd, *J* = 15.8, 7.6, 4.7, 2.9 Hz), 2.57 (1H, dd, *J* = 17.9, 9.7 Hz), 2.45 (1H, dd, *J* = 17.9, 1.8 Hz), 2.40 (1H, dd, *J* = 13.2, 6.2 Hz), 2.36–2.24 (8H, m), 2.20–2.13 (1H, m), 2.10–1.97 (6H, m), 1.92–1.79 (4H, m), 1.78–1.67 (4H, m), 1.60 (1H, ddd, *J* = 14.2, 8.4, 8.4 Hz), 1.56–1.42 (4H, m), 1.42–1.28 (5H, m), 1.10 (3H, d, *J* = 6.5 Hz), 1.06 (3H, d, *J* = 7.6 Hz), 1.02 (3H, d, *J* = 7.0 Hz), 1.04–0.98 (1H, m), 0.97 (3H, d, *J* = 7.0 Hz) ppm. ¹³C NMR (125 MHz, ¹²CD₃OD) δ 172.8, 153.3, 153.1, 114.8, 113.4, 112.9, 105.7, 104.8, 98.4, 85.5, 82.3, 81.3, 81.2, 80.8, 79.0, 78.0, 77.9, 77.6, 77.4, 76.3, 76.0, 75.8, 75.5, 75.2, 75.1, 75.1, 73.8, 73.7, 73.3, 73.1, 73.0, 71.6, 69.6, 67.2, 65.6, 45.6, 45.0, 44.9, 41.1, 39.8, 37.9, 37.9, 37.8, 37.5, 37.5, 37.2, 36.3, 33.0, 31.8, 31.3, 31.3, 30.9, 30.8, 28.4, 27.1, 27.1, 18.4, 18.3, 18.1, 15.9 ppm. FTIR (film): 3429, 2925, 2872, 1736, 1454, 1372, 1269, 1191, 1129, 1109, 1073, 1020, 753 cm⁻¹. HRMS (ESI) *m/z*: [M+H]⁺ calcd for C₆₀H₈₇O₂₁, 1143.5734; found, 1143.5720.

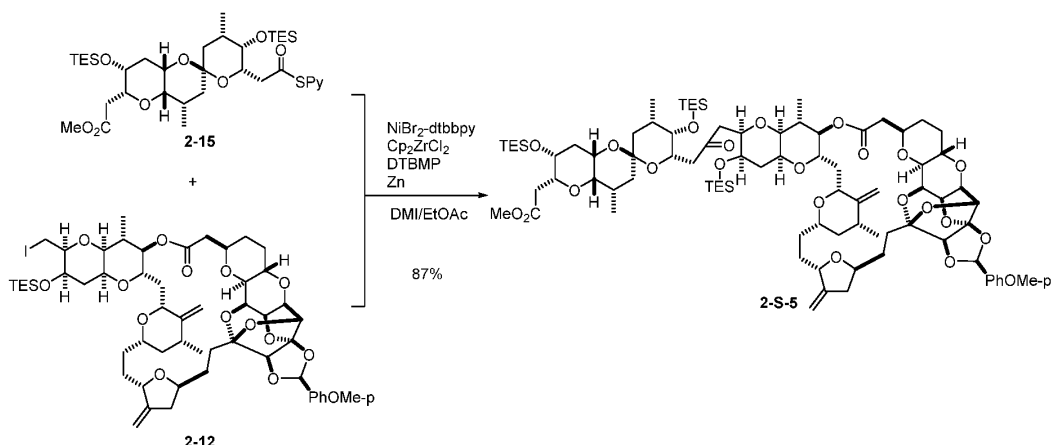


[00697] 38-*epi*-Halichondrin A (C38-*epi*-20): $[\alpha]_D^{20}$ -74.3 (*c* 0.50, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.04 (1H, s), 4.96 (1H, s), 4.87 (1H, s), 4.81 (1H, s), 4.72 (1H, dd, J = 10.2, 6.6 Hz), 4.43 (1H, d, J = 10.2 Hz), 4.38-4.31 (4H, m), 4.37 (1H, ddd, J = 12.0, 12.0, 4.8 Hz), 4.20 (1H, m), 4.18-4.06 (7H, m), 3.99 (1H, ddd, J = 9.6, 5.4, 4.2 Hz), 3.91-3.83 (4H, m), 3.78 (1H, ddd, J = 14.4, 4.8, 4.2 Hz), 3.64 (1H, d, J = 9.6 Hz), 3.59-3.56 (2H, m), 3.53 (1H, dd, J = 10.8, 4.5 Hz), 3.47 (1H, dd, J = 11.4, 6.0 Hz), 3.17 (1H, dd, J = 9.0, 6.6 Hz), 2.95 (1H, dd, J = 9.6, 1.8 Hz), 2.86-2.80 (1H, m), 2.56 (1H, dd, J = 17.4, 9.6 Hz), 2.47 (1H, dd, J = 17.4, 2.4 Hz), 2.36-2.18 (9H, m), 2.12-2.07 (3H, m), 2.04-1.96 (4H, m), 1.88-1.81 (3H, m), 1.77-1.73 (2H, m), 1.69-1.65 (2H, m), 1.62-1.34 (10H, m), 1.09 (3H, d, J = 6.6 Hz), 1.04 (3H, d, J = 8.4 Hz), 1.05-0.99 (1H, m), 1.02 (3H, d, J = 7.8 Hz), 1.00 (3H, d, J = 6.6 Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.9, 153.3, 152.8, 115.6, 113.3, 113.0, 105.1, 104.7, 98.3, 85.5, 82.4, 81.3, 81.1, 80.0, 79.1, 78.9, 78.9, 78.5, 77.9, 76.7, 76.1, 76.0, 75.9, 75.5, 75.3, 74.8, 73.9, 73.5, 73.3, 73.2, 73.2, 71.7, 69.5, 68.3, 67.2, 45.6, 44.9, 44.8, 41.2, 39.7, 38.3, 38.3, 38.1, 37.5, 37.5, 37.2, 36.2, 33.3, 31.8, 31.3, 31.0, 30.9, 29.9, 28.3, 27.1, 26.8, 18.4, 18.3, 18.3, 15.2 ppm. FTIR (film): 3439, 2925, 2872, 1736, 1454, 1372, 1279, 1192, 1119, 1073, 1020, 753 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{60}\text{H}_{87}\text{O}_{21}$, 1143.5734; found, 1143.5721. **C38-*epi*-20** was epimerized to Halichondrin A (**20**) by the following procedure:



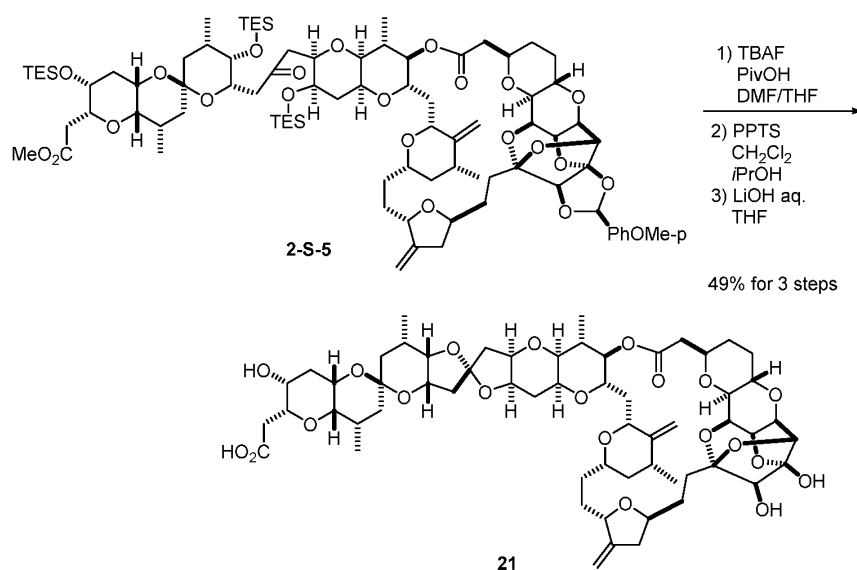
[00698] To a stirred solution of **C38-*epi*-20** (13.0 mg, 0.0114 mmol, 1 eq.) in CH₂Cl₂ (5.7 mL) was added TMSOTf (0.114 mL, 0.631 mmol, excess) at -78 °C. After being stirred for 15 min, the reaction was quenched with sat. NaHCO₃ aq. After being stirred for 1 h at 0 °C, the organic layer was separated and the aqueous layer was extracted with CH₂Cl₂. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 50% MeCN in H₂O) to give halichondrin A (**20**) (11.3 mg, 0.00988 mmol, 86%) as a white solid.

Norhalichondrin A (21)



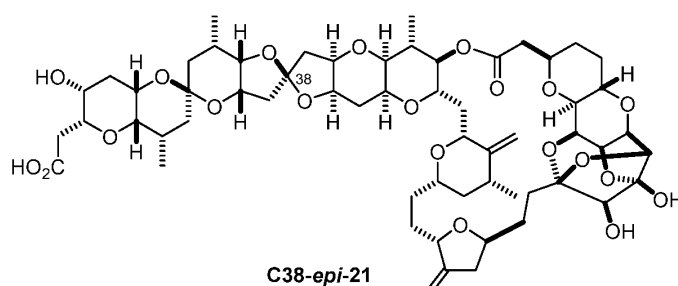
[00699] In a glove box, to a solution of iodide **2-5** (55 mg, 0.0487 mmol, 1 eq.) and thioester **2-16** (45.9 mg, 0.0634 mmol, 1.3 eq.) in DMI (0.4 mL) and EtOAc (0.08 mL) were added

DTBMP (40 mg, 0.195 mmol, 4 eq.), Zn powder (19.0 mg, 0.292 mmol, 6 eq.), Cp_2ZrCl_2 (42.7 mg, 0.146 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (7.1 mg, 0.0146 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et_2O and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with Et_2O . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 15%, 25% EtOAc in Hexanes) to give ketone **2-S-5** (68.4 mg, 0.0423 mmol, 87%) as a colorless amorphous solid. **2-S-5**: $[\alpha]_D^{20}$ -61.5 (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.36 (2H, d, $J = 8.4$ Hz), 6.73 (2H, d, $J = 8.4$ Hz), 6.08 (1H, s), 5.22 (1H, s), 5.11 (1H, s), 4.96 (1H, s), 4.84-4.80 (3H, m), 4.69 (1H, d, $J = 10.2$ Hz), 4.50 (1H, ddd, $J = 9.6, 9.3, 2.0$ Hz), 4.41 (1H, s), 4.39-4.34 (2H, m), 4.06 (1H, s), 4.05-3.96 (4H, m), 3.84-3.77 (2H, m), 3.77-3.70 (3H, m), 3.58 (1H, s), 3.45 (1H, dd, $J = 8.4, 4.2$ Hz), 3.37 (3H, s), 3.35 (1H, s), 3.22 (3H, s), 3.20-3.14 (1H, m), 3.11 (1H, s), 3.07 (1H, dd, $J = 17.4, 5.7$ Hz), 2.99 (1H, dd, $J = 17.4, 6.9$ Hz), 2.84 (1H, dd, $J = 14.4, 7.2$ Hz), 2.78 (1H, dd, $J = 14.8, 7.2$ Hz), 2.76-2.70 (1H, m), 2.57 (1H, dd, $J = 15.0, 5.4$ Hz), 2.53 (1H, d, $J = 9.6$ Hz), 2.43-2.20 (10H, m), 2.16-2.06 (5H, m), 2.06-2.01 (1H, m), 1.98 (1H, dd, $J = 12.0, 12.0$ Hz), 1.87-1.80 (1H, m), 1.78-1.63 (4H, m), 1.62-1.44 (7H, m), 1.38-1.30 (3H, m), 1.20 (3H, d, $J = 7.2$ Hz), 1.13-1.02 (34H, m), 0.98 (3H, d, $J = 7.2$ Hz), 0.71-0.60 (18H, m) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 206.9, 171.8, 171.2, 161.4, 153.0, 152.6, 128.8, 128.7, 119.0, 114.1, 109.3, 109.0, 105.1, 103.8, 96.9, 90.2, 83.8, 78.3, 78.1, 78.0, 77.2, 76.5, 76.1, 76.0, 75.5, 74.8, 74.7, 74.2, 74.0, 73.8, 73.6, 73.0, 70.1, 69.5, 68.4, 65.9, 65.7, 64.6, 64.5, 54.8, 50.9, 46.9, 46.3, 44.0, 41.3, 39.50, 38.7, 37.5, 37.2, 36.4, 36.3, 35.5, 32.4, 31.1, 31.0, 30.9, 30.3, 30.2, 29.2, 27.6, 18.6, 18.2, 17.3, 16.5, 7.5, 7.3, 7.2, 6.0, 5.4, 5.3 ppm. FTIR (film): 2953, 2934, 2876, 2104, 1738, 1518, 1458, 1373, 1304, 1154, 1090, 1033, 1014, 855, 830, 740 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{86}\text{H}_{134}\text{O}_{23}\text{Si}_3\text{Na}$, 1641.8521; found, 1641.8591.



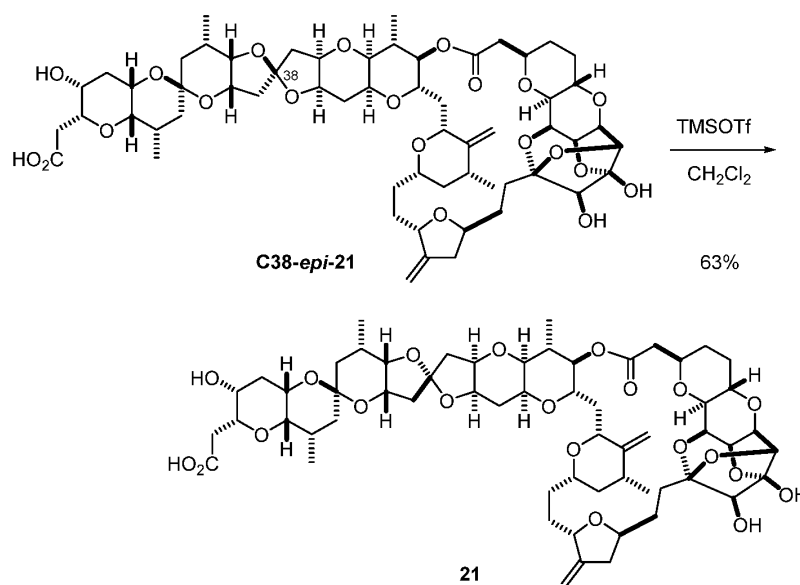
[00700] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.37 mL of 1 M in THF, 0.37 mmol, 10 eq.) and PivOH (18.5 mg, 0.182 mmol, 5 eq.). To a stirred solution of ketone **2-S-5** (59 mg, 0.0364 mmol, 1 eq.) in DMF (2.0 mL) was added the buffered TBAF solution at room temperature. After being stirred for 6 h at the same temperature, CaCO₃ (1.0 g) and DOWEX 50WX8-400 (1.0 g) were added after diluting with 5 mL EtOAc. After being stirred for 1 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of the crude tetraol (calculated as 0.0364 mmol, 1 eq.) in CH₂Cl₂ (8.5 mL) was added PPTS (36.8 mg, 0.146 mmol, 4 eq.) at room temperature. After 1 h, TLC analysis indicated the disappearance of starting material. iPrOH (0.4 mL) and additional PPTS (46.0 mg, 0.183 mmol, 5 eq.) were added to the resulted solution at the same temperature. After being stirred for 12 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (CH₂Cl₂ then 25%, 50%, 75%, then 100% EtOAc in Hexanes then 2% MeOH in EtOAc) to give a crude Norhalichondrin A methyl ester with its C38 epimer. The compound was used in the next step after concentration without further purification. To a stirred solution of the crude methyl ester (calculated as 0.0364 mmol, 1 eq.) in THF (5 mL) was added 1M LiOH aq. (1.5 mL) at room temperature.³ After being stirred for 2 h at the same temperature, the reaction mixture was diluted with water (3 mL). The THF was then removed from the mixture by evaporator. After the reaction was cooled down to 0 °C, 1 M HCl aq. (1.5 mL) was added and the reaction mixture was allowed for further 2

min stirring. The resulting mixture was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The resulting mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give Norhalichondrin A (**21**) (20.2 mg, 0.0179 mmol, 49% for 3 steps) as a colorless solid and 38-*epi*-Norhalichondrin A (**C38-*epi*-21**) (9.8 mg, 0.0087 mmol, 24% for 3 steps) as a colorless solid. **Norhalichondrin A (21)** [α]_D²⁰ -70.3 (*c* 0.37, MeOH) ¹H NMR (600 MHz, CD₃OD) δ : 5.06 (1H, s), 5.02 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.62 (1H, dd, *J* = 7.3, 4.7 Hz), 4.45 (1H, d, *J* = 9.6 Hz), 4.39–4.35 (1H, m), 4.35–4.29 (3H, m), 4.25 (1H, ddd, *J* = 11.2, 4.2, 1.8 Hz), 4.21 (1H, dd, *J* = 3.2, 2.4 Hz), 4.13–4.07 (3H, m), 3.98 (1H, d, *J* = 2.1 Hz), 3.91–3.86 (2H, m), 3.82–3.76 (2H, m), 3.74–3.69 (2H, m), 3.63–3.58 (2H, m), 3.53 (1H, s), 3.33–3.28 (1H, m), 3.22 (1H, dd, *J* = 6.6, 4.7 Hz), 2.93 (1H, dd, *J* = 9.6, 1.8 Hz), 2.83 (1H, ddd, *J* = 16.0, 8.0, 2.1 Hz), 2.58 (1H, dd, *J* = 16.2, 9.6 Hz), 2.53–2.47 (2H, brs), 2.44 (1H, dd, *J* = 17.6, 1.8 Hz), 2.39 (1H, dd, *J* = 13.2, 6.2 Hz), 2.36–2.23 (6H, m), 2.20–2.12 (2H, m), 2.11–1.99 (6H, m), 1.94 (1H, ddd, *J* = 14.8, 3.0, 3.0 Hz), 1.90–1.79 (2H, m), 1.76–1.66 (3H, m), 1.57–1.47 (4H, m), 1.43–1.31 (7H, m), 1.10 (3H, d, *J* = 6.6 Hz), 1.06 (3H, d, *J* = 7.0 Hz), 1.04–1.03 (1H, m), 0.98 (3H, d, *J* = 7.2 Hz), 0.96 (3H, d, *J* = 7.2 Hz) ppm. ¹³C NMR (150 MHz, CD₃OD) δ : 172.8 (2C), 153.3, 153.2, 114.8, 113.4, 112.9, 105.7, 104.8, 98.5, 85.5, 82.3, 80.7, 79.0, 78.8, 78.0 (2C), 77.6, 77.4, 77.3, 76.3, 76.0, 75.8, 75.5, 75.3, 75.1 (2C), 73.8, 73.7, 72.7, 69.6, 68.1, 68.0, 65.7, 45.5, 44.9 (2C), 41.1, 39.8, 38.2, 38.1, 37.8, 37.5, 37.1, 35.7, 33.0, 31.8, 31.34, 31.30, 30.8 (2C), 30.7, 30.1, 28.4, 27.3, 18.4, 18.1, 17.4, 15.8 ppm. FTIR (film): 3458, 2927, 2873, 1750, 1579, 1410, 1269, 1195, 1074, 1019, 991, 967 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₅₉H₈₂O₂₁Na, 1149.5246; found, 1149.5189.



[00701] 38-*epi*-Norhalichondrin A (C38-*epi*-21): [α]_D²⁰ -83.8 (*c* 0.277, MeOH) δ : 5.05 (1H, s), 4.96 (1H, s), 4.87 (1H, s), 4.80 (1H, s), 4.62 (1H, dd, *J* = 9.6, 6.6 Hz), 4.43 (1H, d, *J* = 9.6 Hz), 4.39–4.33 (2H, m), 4.32–4.29 (2H, m), 4.20 (1H, t, *J* = 3.2 Hz), 4.17–4.05 (4H, m), 4.00 (1H, dd, *J* = 5.4, 2.1 Hz), 3.93–3.83 (4H, m), 3.79 (1H, t, *J* = 6.4 Hz), 3.75–3.73 (1H, m),

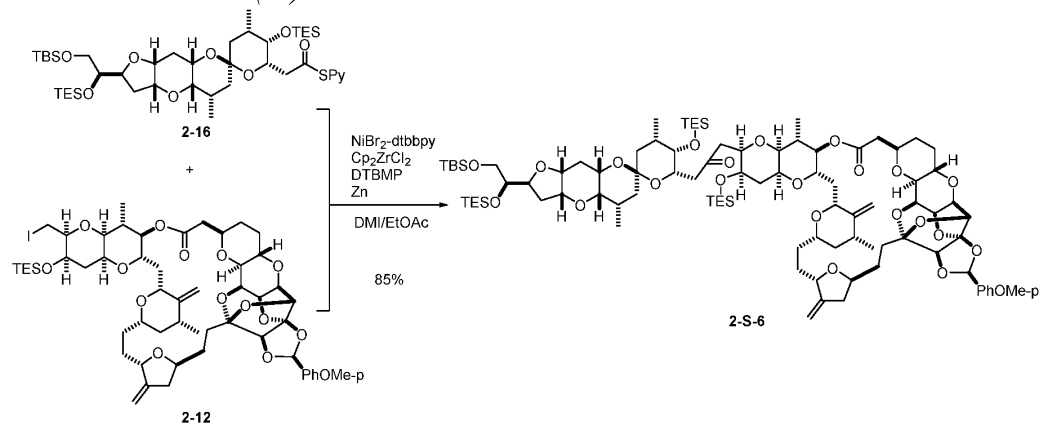
3.67–3.56 (2H, m), 3.52 (1H, s), 3.34–3.32 (1H, m), 3.17 (1H, dd, $J = 9.6, 6.0$ Hz), 2.94 (1H, dd, $J = 9.6, 1.8$ Hz), 2.83 (1H, ddd, $J = 16.0, 8.0, 2.1$ Hz), 2.56 (1H, dd, $J = 16.2, 9.6$ Hz), 2.53–2.47 (2H, brs), 2.47 (1H, dd, $J = 17.6, 1.8$ Hz), 2.35 (1H, d, $J = 15.6$ Hz), 2.33–2.17 (6H, m), 2.14–2.06 (4H, m), 2.05–1.95 (4H, m), 1.92–1.80 (m, 3H), 1.77–1.64 (3H, m), 1.59–1.46 (4H, m), 1.43–1.31 (7H, m), 1.09 (3H, d, $J = 6.6$ Hz), 1.03 (3H, d, $J = 7.0$ Hz), 1.01–0.99 (1H, m), 0.99 (3H, d, $J = 7.2$ Hz), 0.98 (3H, d, $J = 7.2$ Hz). ^{13}C NMR (125 MHz, CD_3OD) 172.9 (2C), 153.4, 152.7, 115.6, 113.4, 113.0, 105.2, 104.7, 98.4, 85.5, 82.4, 79.9, 79.1, 79.0, 78.9, 78.8, 78.6, 78.0, 77.0, 76.7, 76.1, 75.9, 75.8, 75.6, 75.3, 74.8, 74.0, 73.3, 73.2, 69.5, 68.4, 68.2, 68.0, 65.9, 45.5, 44.9, 44.8, 41.2, 39.7, 38.6, 38.2, 38.1, 37.5, 37.2, 35.7, 33.2, 31.8, 31.3, 31.0, 30.9, 30.1, 29.9, 27.0, 18.4, 18.3, 17.5, 15.1 ppm. FTIR (film): 3500 (br), 2927, 2873, 1736, 1579, 1191, 1075, 1020, 1009, 882, 756 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{59}\text{H}_{82}\text{O}_{21}\text{Na}$, 1149.5246; found, 1149.5156. **C38-*epi*-21** was epimerized to Norhalichondrin A (**21**) by the following procedure:



[00702] To a solution of **C38-*epi*-21** (9.8 mg, 0.0087 mmol, 1 eq.) in CH_2Cl_2 (3.5 mL) was added TMSOTf (0.11 mL, 0.435 mmol, excess) at -78°C . After being stirred for 15 min, the reaction was quenched with sat. NaHCO_3 aq. After being stirred for 1 h at 0°C , the organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient:

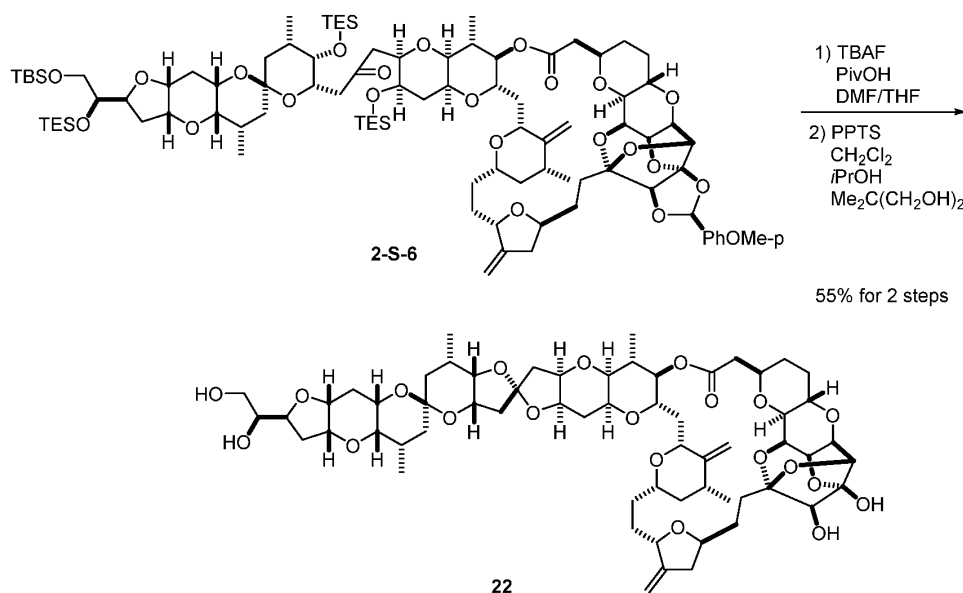
10% MeCN in H₂O to 100% MeCN) to give Norhalichondrin B (**21**) (6.2 mg, 0.0055 mmol, 63%) as a colorless solid.

Homohalichondrin A (22)



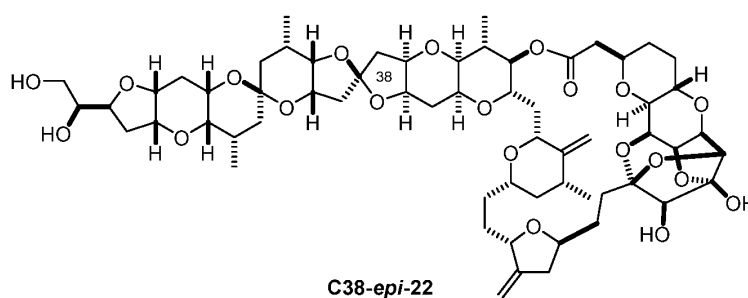
[00703] In a glove box, to a solution of iodide **2-12** (57.0 mg, 0.0503 mmol, 1 eq.) and thioester **2-16** (55.7 mg, 0.0653 mmol, 1.3 eq.) in DMI (0.42 mL) and EtOAc (80 μ L) were added DTBMP (41.3 mg, 0.201 mmol, 4 eq.), Zn powder (19.7 mg, 0.301 mmol, 6 eq.), Cp₂ZrCl₂ (44.1 mg, 0.151 mmol, 3 eq.), and NiBr₂-dtbbpy (7.3 mg, 0.0150 mmol, 30 mol%) at room temperature. After being stirred for 2 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et₂O and sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with Et₂O. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 9%, then 16% EtOAc in Hexanes) to give ketone **2-S-6** (74.8 mg, 0.0428 mmol, 85%) as a colorless amorphous solid. **2-S-6**: $[\alpha]_D^{20}$ -73.0 (*c* 1.07, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.36 (2H, d, *J* = 8.4 Hz), 6.73 (2H, d, *J* = 8.4 Hz), 6.08 (1H, s), 5.22 (1H, s), 5.11 (1H, s), 4.96 (1H, s), 4.84-4.80 (3H, m), 4.69 (1H, d, *J* = 10.2 Hz), 4.52-4.46 (2H, m), 4.41 (1H, s), 4.37 (1H, dd, *J* = 4.8, 1.2 Hz), 4.35 (1H, d, *J* = 10.2 Hz), 4.06-3.97 (5H, m), 3.83-3.80 (3H, m), 3.78-3.69 (69H, m), 3.67 (1H, s), 3.47 (1H, dd, *J* = 8.4, 4.2 Hz), 3.28 (1H, s), 3.22 (3H, s), 3.19-3.15 (2H, m), 3.06 (1H, dd, *J* = 17.4, 5.7 Hz), 3.00 (1H, dd, *J* = 17.4, 6.9 Hz), 2.92 (1H, d, *J* = 1.8 Hz), 2.79 (1H, dd, *J* = 17.1, 7.5 Hz), 2.77-2.73 (1H, m), 2.53 (1H, d, *J* = 9.6 Hz), 2.47 (1H, d, *J* = 15.6 Hz), 2.42-2.21 (10H, m), 2.15-2.03 (6H, m), 1.98 (1H, dd, *J* = 12.0, 12.0 Hz), 1.87-1.82 (2H, m), 1.76-1.67 (4H, m), 1.61-1.47 (6H, m), 1.38-1.27 (3H, m), 1.20 (3H, d, *J* = 7.8 Hz), 1.11 (9H, t, *J* = 8.1 Hz), 1.09 (3H, d, *J* = 6.6 Hz), 1.06-1.04 (21H, m), 1.00 (9H, s), 0.96 (3H, d, *J* = 6.0 Hz), 0.77 (6H, q, *J* = 7.8 Hz), 0.70-0.65 (12H, m), 0.14

(3H, s), 0.13 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.7, 171.1, 161.4, 153.0, 152.7, 128.73, 128.69, 119.0, 114.1, 109.3, 109.0, 105.1, 103.8, 96.9, 90.2, 83.8, 79.1, 78.4, 78.1, 77.99, 77.96, 76.8, 76.1, 76.0, 75.5, 74.8, 74.7, 74.2, 74.0, 73.8, 73.6, 73.1, 70.1, 69.3, 68.4, 66.2, 65.9, 64.6, 63.8, 54.8, 46.9, 46.3, 44.0, 41.3, 39.50, 39.45, 38.7, 37.7, 37.6, 36.5, 36.4, 35.5, 32.4, 31.6, 31.1, 30.9, 30.7, 30.3, 30.2, 29.6, 27.7, 26.2, 18.7, 18.6, 18.2, 17.6, 16.5, 7.5, 7.4, 7.3, 6.0, 5.7, 5.3, -5.1, -5.3 ppm. FTIR (film): 2953, 2927, 2875, 2104, 1724, 1615, 1518, 1459, 1373, 1306, 1251, 1092, 1032, 1010, 833, 742 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{93}\text{H}_{150}\text{O}_{23}\text{Si}_4\text{Na}$, 1769.9537; found, 1769.9316.



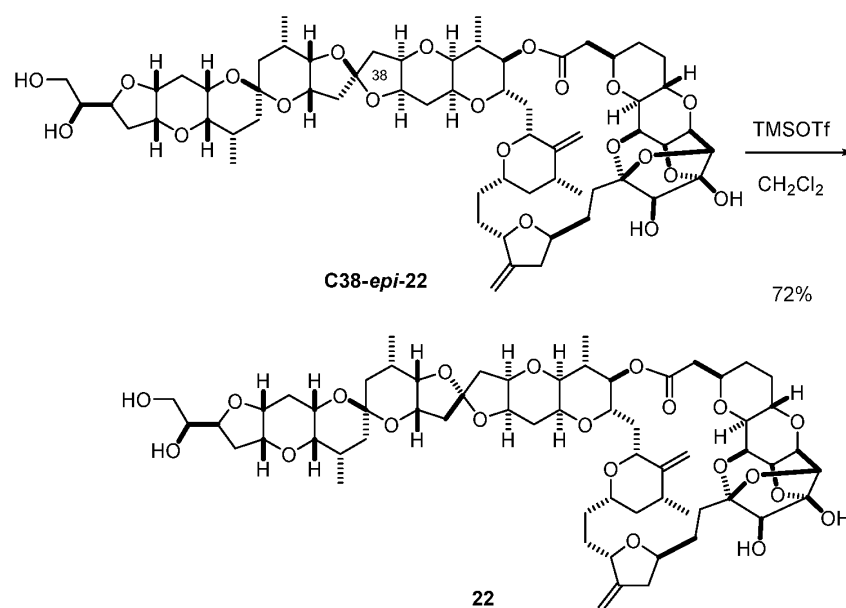
[00704] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.43 mL of 1 M in THF, 0.43 mmol, 10 eq.) and PivOH (22.0 mg, 0.215 mmol, 5 eq.). To a solution of ketone **S-6** (74.8 mg, 0.0428 mmol, 1 eq.) in DMF (2.1 mL) was added the buffered TBAF solution at room temperature. After being stirred for 7 h at the same temperature, CaCO_3 (1.5 g) and DOWEX 50WX8-400 (1.5 g) were added. After being stirred for 1 h at the same temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of above tetraol (calculated as 0.0428 mmol, 1 eq.) in CH_2Cl_2 (2.1 mL) was added PPTS (53.8 mg, 0.214 mmol, 5 eq.) at room temperature. After 1 h, TLC analysis indicated the disappearance of starting material. $i\text{PrOH}$ (0.43 mL) and additional PPTS (53.8 mg, 0.214 mmol, 5 eq.) were added to the resulted solution at the same temperature. After being stirred for 18 h, the reaction mixture was

directly subjected to column chromatography on amino silica gel (CH_2Cl_2 then 25%, 50%, 75%, then 100% EtOAc in Hexanes then 4% MeOH in EtOAc) to give a crude Homohalichondrin A with its C-38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H_2O to 100% MeCN) to give Homohalichondrin A (**22**) (27.2 mg, 0.0235 mmol, 55% for 2 steps) as a colorless solid and 38-*epi*-Homohalichondrin A (**C38-*epi*-22**) (10.7 mg, 0.00926 mmol, 22% for 2 steps) as a colorless solid. **Homohalichondrin A (22)**: $[\alpha]_D^{20}$ -80.3 (c 1.22, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.07 (1H, s), 5.01 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.63 (1H, dd, J = 7.8, 4.8 Hz), 4.44 (1H, d, J = 4.8 Hz), 4.37 (1H, s), 4.34-4.31 (3H, m), 4.25-4.22 (2H, m), 4.21 (1H, dd, J = 2.7, 2.7 Hz), 4.11-4.08 (3H, m), 4.03 (1H, s), 3.95 (1H, s), 3.89-3.87 (3H, m), 3.71 (1H, dd, J = 10.2, 10.2 Hz), 3.66 (1H, s), 3.61-3.57 (4H, m), 3.53 (1H, s), 3.50 (1H, dd, J = 9.6, 4.8 Hz), 3.22 (1H, dd, J = 6.0, 6.0 Hz), 3.12 (1H, s), 2.94 (1H, d, J = 10.2 Hz), 2.82 (1H, dd, J = 15.9, 5.7 Hz), 2.57 (1H, dd, J = 17.3, 9.3 Hz), 2.45 (1H, d, J = 17.3 Hz), 2.39 (1H, dd, J = 13.5, 5.7 Hz), 2.38-2.24 (6H, m), 2.20-2.14 (3H, m), 2.09-1.98 (8H, m), 1.91-1.81 (4H, m), 1.74-1.68 (3H, m), 1.51-1.32 (9H, m), 1.10 (3H, d, J = 6.6 Hz), 1.05 (3H, d, J = 7.2 Hz), 1.05-0.99 (1H, m), 0.95 (3H, d, J = 6.6 Hz), 0.94 (3H, d, J = 6.0 Hz) ppm. ^{13}C NMR (125 MHz, $^{12}\text{CD}_3\text{OD}$) δ : 172.8, 153.2, 153.0, 114.7, 113.3, 112.9, 105.7, 104.8, 98.1, 85.5, 82.3, 81.0, 79.8, 79.0, 78.4, 78.1, 77.9, 77.5, 76.3, 75.9, 75.8, 75.7, 75.5, 75.3, 74.5, 73.8, 73.7, 72.3, 69.6, 65.8, 65.2, 65.1, 49.9, 45.5, 44.9, 44.8, 41.1, 39.8, 38.1, 38.0, 37.8, 37.4, 37.23, 37.15, 33.0, 32.0, 31.8, 31.3, 30.8, 30.7, 30.1, 28.4, 27.1, 18.4, 18.2, 17.7 ppm. FTIR (film): 3398, 2927, 2873, 1736, 1372, 1269, 1187, 1129, 1074, 1020, 902, 753 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{NH}_4]^+$ calcd for $\text{C}_{61}\text{H}_{90}\text{NO}_{21}$, 1172.6000; found, 1172.5982.



[00705] 38-*epi*-Homohalichondrin A (C38-*epi*-22): $[\alpha]_D^{20}$ -92.3 (c 0.573, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.04 (1H, d, J = 1.8 Hz), 4.96 (1H, d, J = 1.8 Hz), 4.87 (1H, s), 4.80 (1H, s), 4.72 (1H, dd, J = 10.5, 6.6 Hz), 4.43 (1H, d, J = 10.2 Hz), 4.37 (1H, ddd, J = 9.6, 9.6, 4.2 Hz), 4.34 (1H, s), 4.32-4.30 (2H, m), 4.23 (1H, ddd, J = 10.2, 5.7, 4.5 Hz), 4.20 (1H, dd, J = 2.7, 2.7 Hz), 4.16-4.06 (4H, m), 4.02 (1H, s), 3.98 (1H, dd, J = 4.5, 2.1 Hz), 3.91-3.84 (3H,

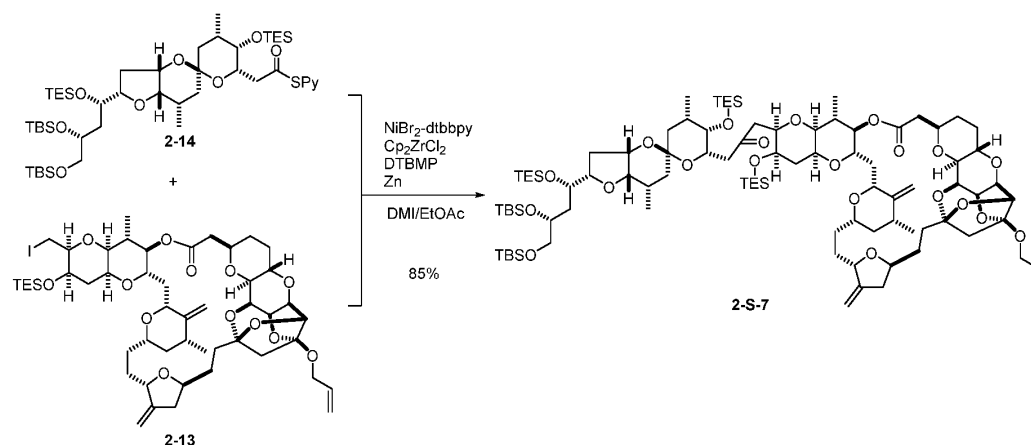
m), 3.81 (1H, dd, $J = 2.4, 2.4$ Hz), 3.64 (1H, dd, $J = 10.2, 10.2$ Hz), 3.60-3.56 (3H, m), 3.54 (1H, d, $J = 3.0$ Hz), 3.52 (1H, s), 3.51-3.47 (1H, m), 3.17 (1H, dd, $J = 8.4, 6.0$ Hz), 3.15 (1H, d, $J = 1.8$ Hz), 2.95 (1H, dd, $J = 9.6, 1.8$ Hz), 2.85-2.81 (1H, m), 2.56 (1H, dd, $J = 17.1, 9.0$ Hz), 2.47 (1H, dd, $J = 17.1, 2.4$ Hz), 2.36-1.97 (19H, m), 1.91 (1H, ddd, $J = 15.9, 4.8, 4.8$ Hz), 1.88-1.81 (2H, m), 1.79-1.65 (3H, m), 1.57-1.52 (1H, m), 1.47 (1H, dd, $J = 12.9, 12.9$ Hz), 1.42-1.35 (6H, m), 1.29 (1H, dd, $J = 12.6, 4.8$ Hz), 1.09 (3H, d, $J = 6.6$ Hz), 1.06-0.99 (1H, m), 1.003 (3H, d, $J = 7.2$ Hz), 0.997 (3H, d, $J = 6.6$ Hz), 0.96 (3H, d, $J = 7.2$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.9, 153.3, 152.7, 115.6, 113.3, 113.0, 105.1, 104.7, 97.9, 85.5, 82.4, 80.2, 79.8, 79.1, 78.88, 78.86, 78.5, 78.4, 77.8, 76.7, 76.1, 75.9, 75.83, 75.76, 75.5, 75.3, 75.1, 74.8, 74.4, 73.9, 73.2, 72.9, 69.5, 68.3, 65.3, 65.1, 45.6, 45.0, 44.8, 41.2, 39.7, 38.6, 38.3, 38.1, 37.5, 37.2, 33.3, 31.9, 31.7, 31.3, 31.0, 30.9, 30.1, 29.9, 28.3, 26.8, 18.4, 17.7, 15.2 ppm. FTIR (film): 3445, 2926, 2873, 1737, 1435, 1373, 1267, 1193, 1108, 1075, 1018, 897, 736 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{61}\text{H}_{86}\text{O}_{21}\text{Na}$, 1177.5536; found, 1177.5554. **C38-*epi*-22** was epimerized to Homohalichondrin A (**22**) by following procedure:



[00706] To a solution of **C38-*epi*-22** (9.7 mg, 0.00840 mmol, 1 eq.) in CH_2Cl_2 (4.2 mL) was added TMSOTf (0.084 mL, 0.465 mmol, excess) at -78°C . After being stirred for 15 min, the reaction was quenched with sat. NaHCO_3 aq. After being stirred for 1 h at 0°C , the organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The crude material was purified by YAMAZEN purification system with ODS column (Rf gradient:

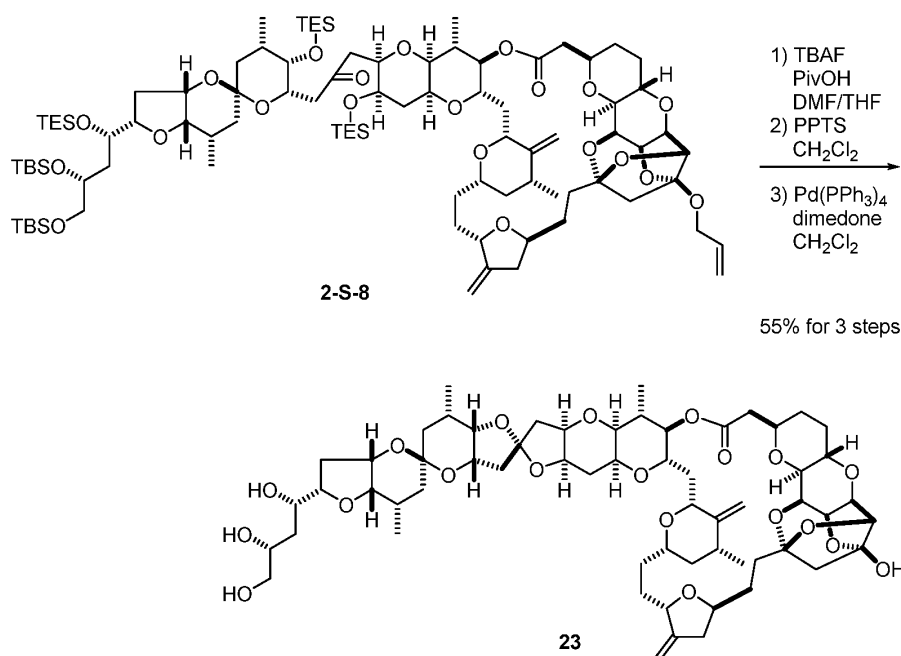
10% MeCN in H₂O to 100% MeCN) to give Homohalichondrin A (**22**) (7.0 mg, 0.00606 mmol, 72%) as a colorless solid.

Halichondrin C (23)



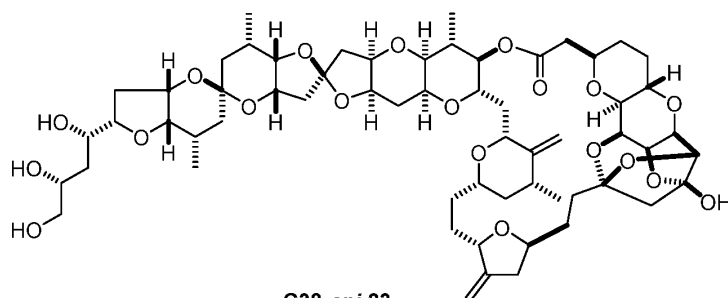
[00707] In a glove box, to a solution of iodide **2-13** (100 mg, 0.0962 mmol, 1 eq.) and thioester **2-14** (119 mg, 0.1246 mmol, 1.3 eq.) in DMI (0.80 mL) and EtOAc (0.16 mL) were added DTBMP (79 mg, 0.385 mmol, 4 eq.), Zn powder (37.7 mg, 0.576 mmol, 6 eq.), Cp₂ZrCl₂ (84.4 mg, 0.289 mmol, 3 eq.), and NiBr₂-dtbbpy (14.0 mg, 0.0288 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with EtOAc and sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by YAMAZEN purification system on neutral silica gel (0%, 9%, then 15% EtOAc in Hexanes) to give ketone **S-7** (143.2 mg, 0.0818 mmol, 85%) as a white amorphous solid. **2-S-8**: $[\alpha]^{20}_{\text{D}} -59.4$ (*c* 1.25, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 5.75 (1H, dddd, *J* = 17.3, 10.5, 5.1, 5.1 Hz), 5.21 (1H, s), 5.15 (1H, ddd, *J* = 17.3, 1.8, 1.8 Hz), 5.12 (1H, s), 4.97 (1H, ddd, *J* = 10.5, 1.8, 1.8 Hz), 4.94 (1H, s), 4.85 (1H, dd, *J* = 7.2, 7.2 Hz), 4.81-4.79 (2H, m), 4.67 (1H, d, *J* = 10.2 Hz), 4.54-4.50 (2H, m), 4.37-4.36 (2H, m), 4.32 (1H, d, *J* = 4.2 Hz), 4.26-4.23 (1H, m), 4.17-4.14 (1H, m), 4.11 (1H, dd, *J* = 6.6, 4.8 Hz), 4.06-3.99 (3H, m), 3.93-3.88 (2H, m), 3.84-3.74 (7H, m), 3.70 (1H, dd, *J* = 13.2, 6.0 Hz), 3.44-3.43 (2H, m), 3.33 (1H, s), 3.19 (1H, dd, *J* = 17.6, 6.3 Hz), 3.16 (1H, dd, *J* = 17.6, 6.6 Hz), 3.11-3.02 (2H, m), 2.77 (1H, dd, *J* = 10.8, 2.4 Hz), 2.57 (1H, d, *J* = 9.6 Hz), 2.49-2.44 (1H, m), 2.40-2.23 (9H, m), 2.18-2.02 (9H, m), 1.96 (1H, dd, *J* = 13.8, 6.0 Hz), 1.86-1.66 (7H, m), 1.61 (1H, ddd, *J* = 15.0, 4.8, 4.8 Hz), 1.55-1.47 (5H, m), 1.36-1.32 (3H, m),

1.19 (3H, d, $J = 7.2$ Hz), 1.15 (3H, d, $J = 6.6$ Hz), 1.11 (9H, t, $J = 8.0$ Hz), 1.10 (9H, s), 1.09 (9H, t, $J = 8.4$ Hz), 1.07 (9H, t, $J = 8.4$ Hz), 1.04 (9H, s), 1.01 (3H, d, $J = 7.2$ Hz), 0.96 (3H, d, $J = 7.2$ Hz), 0.77 (6H, q, $J = 7.8$ Hz), 0.70-0.65 (12H, m), 0.28 (6H, s), 0.15 (6H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.8, 171.3, 152.9, 152.6, 134.6, 116.6, 116.2, 109.3, 104.9, 103.8, 97.0, 83.3, 81.5, 80.4, 78.4, 77.9, 77.6, 75.9, 75.4, 75.1, 75.0, 75.0, 74.9, 74.7, 74.1, 74.0, 73.8, 72.8, 72.0, 71.8, 71.5, 70.4, 69.9, 68.4, 68.3, 65.9, 64.6, 51.7, 46.8, 46.3, 43.9, 41.2, 39.4, 39.3, 38.5, 38.2, 37.8, 36.4, 35.9, 35.4, 35.3, 32.5, 31.1, 30.7, 30.6, 30.5, 28.5, 26.6, 26.3($\times 6$), 26.3($\times 6$), 18.7, 18.6, 18.5, 18.4, 18.1, 16.4, 7.4($\times 6$), 7.4($\times 6$), 7.3($\times 6$), 6.0($\times 3$), 5.7($\times 3$), 5.3($\times 3$), -4.0, -4.2, -5.1, -5.2 ppm. FTIR (film): 2953, 2928, 2876, 1732, 1461, 1410, 1372, 1253, 1079, 1034, 1006, 834, 740 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{93}\text{H}_{162}\text{O}_{21}\text{Si}_5\text{Na}$, 1778.0347; found, 1778.0332.

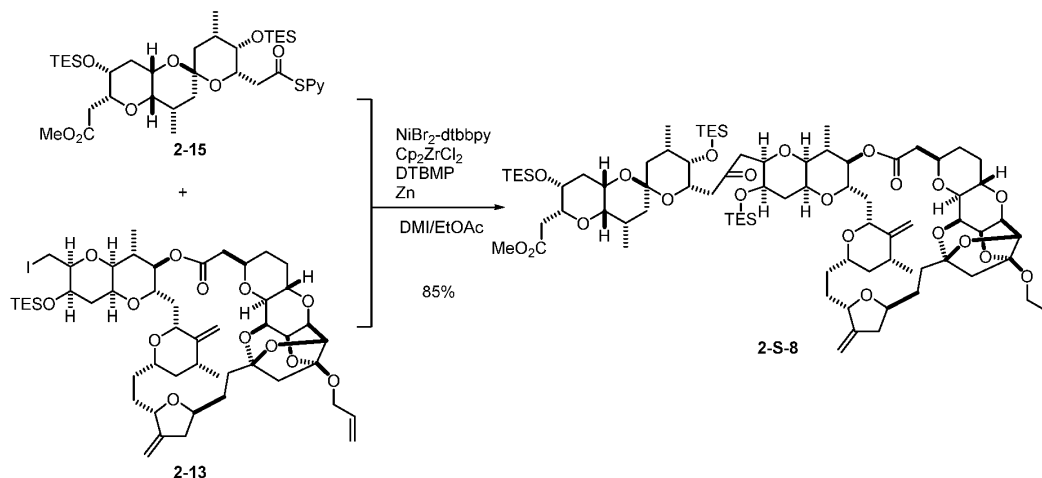


[00708] To a stirred solution of **2-S-8** (141 mg, 0.0803 mmol, 1 equiv.) in DMF (4.0 mL, 0.02M) was added the buffered TBAF solution (0.80 mL, 10 equiv., freshly prepared by 1.60 mL TBAF solution (1 M in THF) and 81.6 mg PivOH) at room temperature. After being stirred for 3 h at the same temperature, CaCO_3 (3.0 g) and DOWEX 50WX8-400 (3.0 g) were added.¹ After being stirred for 2 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude material, which was used in the next step without further purification. To a stirred solution of above

tetraol (calculated as 0.0803 mmol, 1 eq.) in CH₂Cl₂ (8 mL) was added PPTS (100 mg, 0.402 mmol, 5 eq.) at room temperature. After being stirred for 3 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (CH₂Cl₂ then 50%, 75%, then 100% EtOAc in Hexanes then 9% MeOH in EtOAc) to give a crude spiro ketal, which was used in the next step without further purification. To a mixture of above crude spiro ketal (calculated as 0.0803 mmol, 1 eq.), dimedone (22.5 mg, 0.160 mmol, 2 equiv.), and Pd(PPh₃)₄ (9.3 mg, 0.00803 mmol, 10 mol%) was added degassed CH₂Cl₂ (8 mL) at room temperature. After being stirred for 4 h at the same temperature, the resulted solution was directly subjected to column chromatography on amino silica gel (CH₂Cl₂ then 50%, 100% EtOAc in Hexanes then 9% MeOH in EtOAc) to give a crude halichondrin C with its C-38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give Halichondrin C (**23**) (48.1 mg, 0.0427 mmol, 55% for 3 steps) as a white crystalline solid and 38-*epi*-halichondrin C (**C38-*epi*-23**) (9.5 mg, 0.00843 mmol, 11% for 3 steps) as a colorless solid. **Halichondrin C (23)**: $[\alpha]_D^{20}$ -66.8 (c 0.25, MeOH). ¹H NMR (600 MHz, CD₃OD) δ : 5.07 (1H, s), 5.01 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.62 (1H, dd, *J* = 7.2, 4.4 Hz), 4.43 (1H, d, *J* = 11.2 Hz), 4.42-4.38 (1H, m), 4.34-4.27 (3H, m), 4.24 (1H, ddd, *J* = 11.2, 4.4, 2.0 Hz), 4.17 (1H, dd, *J* = 3.8, 1.0 Hz), 4.14-4.06 (4H, m), 4.05 (1H, ddd, *J* = 2.0, 2.0, 2.0 Hz), 3.99 (1H, ddd, *J* = 9.4, 4.3, 4.3 Hz), 3.92-3.82 (3H, m), 3.77 (1H, ddd, *J* = 8.4, 4.4, 4.2 Hz), 3.74-3.66 (2H, m), 3.61 (1H, d, *J* = 11.2 Hz), 3.58-3.55 (1H, m), 3.53 (1H, dd, *J* = 11.2, 4.4 Hz), 3.46 (1H, dd, *J* = 11.2, 6.4 Hz), 3.22 (1H, dd, *J* = 6.4, 4.8 Hz), 2.94 (1H, dd, *J* = 9.8, 2.0 Hz), 2.84-2.75 (1H, m), 2.55 (1H, dd, *J* = 17.6, 9.2 Hz), 2.45 (1H, dd, *J* = 17.6, 2.4 Hz), 2.39 (1H, dd, *J* = 13.2, 6.3 Hz), 2.36-2.21 (9H, m), 2.20-2.02 (5H, m), 2.01-1.94 (2H, m), 1.89-1.78 (3H, m), 1.78-1.56 (6H, m), 1.56-1.20 (11H, m), 1.10 (3H, d, *J* = 6.3 Hz), 1.06 (3H, d, *J* = 6.8 Hz), 1.05-0.99 (1H, m), 1.02 (3H, d, *J* = 7.2 Hz), 0.97 (3H, d, *J* = 6.8 Hz). ¹³C NMR (125 MHz, CD₃OD) δ : 172.8, 153.3, 153.1, 114.8, 114.2, 110.4, 105.7, 104.7, 98.4, 86.2, 81.3, 81.2, 80.8, 79.0, 78.0, 78.0, 77.4, 77.2, 76.4, 76.3, 76.1, 75.8, 75.3, 75.0, 74.9, 74.7, 73.8, 73.3, 73.1, 73.0, 71.6, 69.5, 67.2, 65.7, 53.7, 45.5, 44.9, 44.9, 41.2, 39.7, 38.0, 37.9, 37.8, 37.5, 37.5, 37.2, 36.3, 36.2, 33.0, 31.8, 31.2, 31.0, 30.8, 29.0, 27.1, 27.0, 18.4, 18.3, 18.1, 15.8. FTIR (film): 3422, 2926, 2873, 1736, 1436, 1310, 1186, 1117, 1074, 1021, 995, 910, 755 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₆₀H₈₆O₂₀Na, 1149.5605; found, 1149.5614.

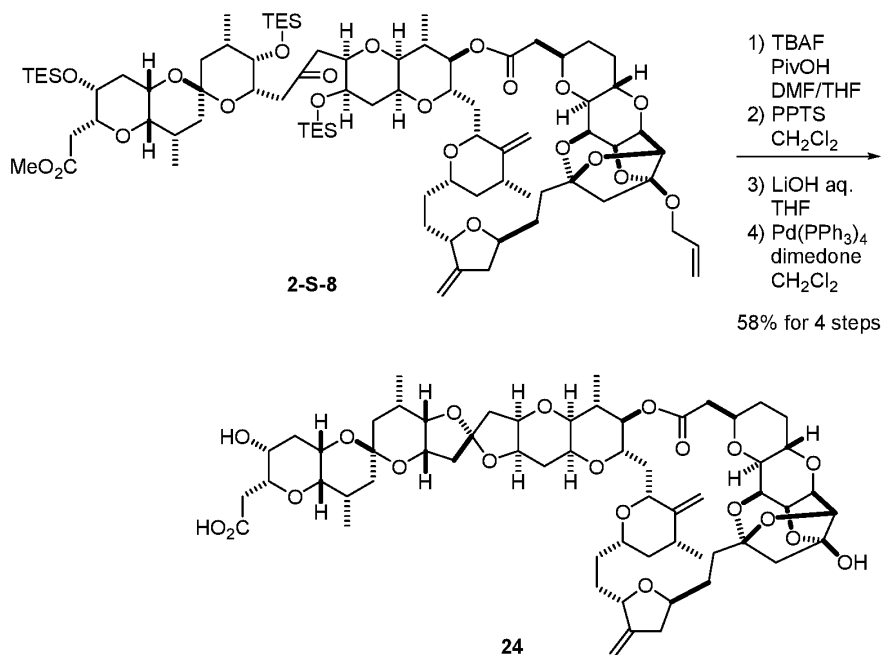
**C38-epi-23**

[00709] 38-*epi*-Halichondrin C (C38-*epi*-23): $[\alpha]_D^{20}$ -69.6 (*c* 0.46, MeOH). ^1H NMR (600 MHz, CD_3OD) δ : 5.03 (1H, s), 4.95 (1H, s), 4.87 (1H, s), 4.81 (1H, s), 4.72 (1H, dd, J = 10.2, 6.6 Hz), 4.41 (1H, d, J = 11.4 Hz), 4.38-4.32 (2H, m), 4.32-4.28 (2H, m), 4.17-4.06 (6H, m), 3.99 (1H, ddd, J = 9.6, 5.4, 4.2 Hz), 3.90-3.83 (4H, m), 3.78 (1H, ddd, J = 14.4, 4.8, 4.2 Hz), 3.63-3.56 (3H, m), 3.53 (1H, dd, J = 11.6, 4.5 Hz), 3.47 (1H, dd, J = 10.8, 6.0 Hz), 3.17 (1H, dd, J = 9.0, 6.0 Hz), 2.95 (1H, dd, J = 9.6, 1.8 Hz), 2.86-2.80 (1H, m), 2.54 (1H, dd, J = 16.8, 8.4 Hz), 2.47 (1H, dd, J = 16.8, 2.4 Hz), 2.34-2.08 (12H, m), 2.12-2.07 (3H, m), 2.04-1.96 (4H, m), 1.88-1.81 (2H, m), 1.79-1.29 (15H, m), 1.10 (3H, d, J = 6.0 Hz), 1.04 (3H, d, J = 8.4 Hz), 1.05-0.99 (1H, m), 1.02 (3H, d, J = 6.6 Hz), 1.00 (3H, d, J = 6.6 Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 172.9, 153.3, 152.8, 115.6, 114.2, 110.5, 105.1, 104.7, 98.3, 86.2, 81.3, 81.1, 80.0, 79.2, 78.9, 78.9, 78.4, 77.9, 76.6, 76.4, 76.1, 76.0, 75.9, 75.2, 74.9, 74.8, 73.6, 73.3, 73.2, 73.1, 71.7, 69.4, 68.3, 67.2, 53.7, 45.6, 44.9, 44.7, 41.2, 39.6, 38.3, 38.3, 38.1, 37.5, 37.5, 37.2, 36.2, 36.0, 33.3, 31.7, 31.2, 30.9, 30.2, 28.9, 27.1, 26.8, 18.4, 18.3, 18.3, 15.2. ppm. FTIR (film): 3427, 2925, 2872, 1736, 1662, 1553, 1436, 1311, 1188, 1117, 1075, 1023, 996, 898, 735 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{60}\text{H}_{86}\text{O}_{20}\text{Na}$, 1149.5605; found, 1149.5618.

Norhalichondrin C (24)

[00710] In a glove box, to a solution of iodide **2-13** (50 mg, 0.0482 mmol, 1 eq.) and thioester **2-15** (45.4 mg, 0.0627 mmol, 1.3 eq.) in DMI (0.4 mL) and EtOAc (0.08 mL) were added DTBMP (39.5 mg, 0.193 mmol, 4 eq.), Zn powder (18.8 mg, 0.289 mmol, 6 eq.), Cp_2ZrCl_2 (42.2 mg, 0.144 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (7.0 mg, 0.0144 mmol, 30 mol%) at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et_2O and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with Et_2O . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 15%, 25% EtOAc in Hexanes) to give ketone **2-S-8** (62.4 mg, 0.0409 mmol, 85%) as a colorless amorphous solid. (**S-8**): $[\alpha]_D^{20}$ -64.8 (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 5.75 (1H, dddd, $J = 17.3, 10.5, 5.1, 5.1$ Hz), 5.21 (1H, s), 5.15 (1H, ddd, $J = 17.3, 1.8, 1.8$ Hz), 5.11 (1H, s), 4.97 (1H, ddd, $J = 10.5, 1.8, 1.8$ Hz), 4.94 (1H, s), 4.87-4.82 (1H, m), 4.82-4.77 (2H, m), 4.67 (1H, d, $J = 10.8$ Hz), 4.52 (1H, ddd, $J = 10.2, 10.2, 4.8$ Hz), 4.38-4.33 (2H, m), 4.32 (1H, $J = 4.6$ Hz), 4.11 (1H, dd, $J = 6.9, 5.1$ Hz), 4.07-3.97 (4H, m), 3.84-3.78 (4H, m), 3.77-3.68 (3H, m), 3.58 (1H, d, $J = 1.2$ Hz), 3.43 (1H, ddd, $J = 4.5, 4.5, 4.5$ Hz), 3.37 (4H, s), 3.36 (1H, s), 3.21-3.11 (3H, m), 2.86-2.81 (1H, m), 2.80-2.74 (2H, m), 2.59-2.54 (2H, m), 2.42-2.35 (3H, m), 2.34-2.22 (8H, m), 2.20-2.04 (7H, m), 1.97 (1H, dd, $J = 13.8, 13.5$ Hz), 1.87-1.80 (1H, m), 1.78-1.63 (4H, m), 1.62-1.44 (7H, m), 1.38-1.30 (2H, m), 1.19 (3H, d, $J = 7.2$ Hz), 1.13-1.02 (31H, m), 1.01 (3H, d, $J = 7.2$ Hz), 0.99 (3H, d, $J = 7.2$ Hz), 0.71-0.60 (18H, m) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.9, 171.8, 171.3, 152.9, 152.6, 134.6, 116.6, 116.2, 109.3, 105.0, 103.8, 96.9, 83.3, 78.3, 77.9, 77.7, 77.2, 76.5, 76.0,

75.4, 75.1, 75.0, 74.9, 74.6, 74.1, 74.0, 73.8, 73.0, 70.4, 69.6, 68.4, 66.0, 65.9, 64.6, 64.5, 51.7, 50.9, 46.8, 46.3, 43.9, 41.2, 39.5, 39.3, 38.6, 37.7, 37.2, 36.4, 36.3, 35.9, 35.4, 32.5, 31.1, 31.0, 30.7, 30.5, 29.2, 28.5, 18.6, 18.2, 17.3, 16.5, 7.5, 7.3, 7.2, 6.0, 5.4, 5.3 ppm. FTIR (film): 2953, 2913, 2876, 1737, 1458, 1372, 1337, 1312, 1280, 1208, 1186, 1155, 1119, 1085, 1072, 1035, 1012, 823, 736 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{81}\text{H}_{132}\text{O}_{21}\text{Si}_3\text{Na}$, 1547.8467; found, 1547.8524.



[00711] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.33 mL of 1 M in THF, 0.33 mmol, 10 eq.) and PivOH (16.7 mg, 0.164 mmol, 5 eq.). To a stirred solution of ketone **2-S-8** (50 mg, 0.0328 mmol, 1 eq.) in DMF (2.0 mL) was added the buffered TBAF solution at room temperature. After being stirred for 6 h at the same temperature, CaCO_3 (1.0 g) and DOWEX 50WX8-400 (1.0 g) were added after diluting with 5 mL EtOAc. After being stirred for 1 h at room temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step without further purification. To a stirred solution of the crude tetraol (calculated as 0.0328 mmol, 1 eq.) in CH_2Cl_2 (8.5 mL) was added PPTS (33.1 mg, 0.132 mmol, 4 eq.) at room temperature. After 1 h, TLC analysis indicated the disappearance of starting material. The reaction mixture was directly subjected to column chromatography on amino silica gel (CH_2Cl_2 then 25%, 50%, 75%, then 100% EtOAc in

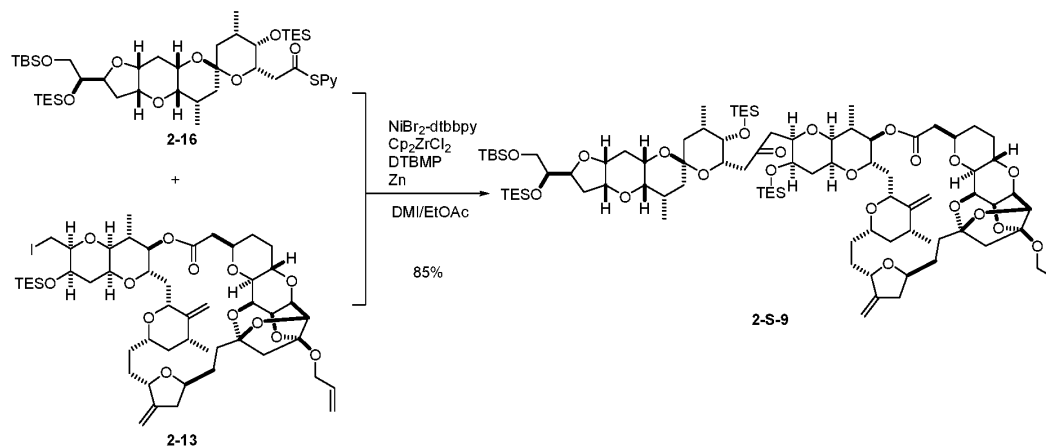
Hexanes then 2% MeOH in EtOAc) to give a crude allyl protected Norhalichondrin C methyl ester with its C38 epimer. The compound was used in the next step after concentration without further purification. To a stirred solution of the crude methyl ester (calculated as 0.0328 mmol, 1 eq.) in THF (5 mL), 1.3 mL of aqueous 1M LiOH was added at room temperature. After being stirred for 1.5 h at the same temperature, the reaction mixture was diluted by 3 mL of water. The THF was then removed from the mixture by evaporator. After the reaction was cooled down to 0 °C, 1.5 mL of 1 M aqueous HCl was added immediately followed by 10 mL of PH 7 aqueous buffer solution. The resulting mixture was extracted with EtOAc. The combined organic layers were dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The resulting mixture was moved to next step without further purification.

To a mixture of above crude acid (calculated as 0.0328 mmol, 1 eq.), dimedone (9.2 mg, 0.0656 mmol, 2 eq.), and Pd(PPh₃)₄ (5.7 mg, 0.00492 mmol, 15 mol%) was added CH₂Cl₂ (4.0 mL) at room temperature. After being stirred for 3 h at the same temperature, the resulted solution was diluted with DMF (3 mL). After removal of DCM by evaporator, the mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H₂O to 100% MeCN) to give Norhalichondrin C (**24**) (21.0 mg, 0.019 mmol, 58% for 4 steps) as a colorless solid. 38-*epi*-Homohalichondrin C (**C38-*epi*-24**) was decomposed while concentrating with inseparable reagent residue after ODS column. **Norhalichondrin C (24)**:

[α]_D²⁰ -61.2 (c 0.300, MeOH) ¹H NMR (600 MHz, CD₃OD) δ : 5.06 (1H, s), 5.01 (1H, s), 4.88 (1H, s), 4.81 (1H, s), 4.62 (1H, dd, *J* = 7.2, 4.8 Hz), 4.43 (1H, d, *J* = 10.8 Hz), 4.42–4.39 (1H, m), 4.34–4.27 (3H, m), 4.24 (1H, ddd, *J* = 11.2, 4.2, 1.8 Hz), 4.17 (1H, dd, *J* = 4.2 Hz), 4.11 (2H, brs), 4.09–4.05 (1H, m), 3.98 (1H, brs), 3.91–3.85 (2H, m), 3.82–3.76 (2H, m), 3.73–3.67 (2H, m), 3.62–3.57 (2H, m), 3.53 (1H, s), 3.31 (1H, m), 3.21 (1H, dd, *J* = 6.6, 4.7 Hz), 2.94 (1H, d, *J* = 9.6 Hz), 2.81 (1H, dd, *J* = 15.0, 6.6 Hz), 2.55 (1H, dd, *J* = 16.2, 9.6 Hz), 2.52–2.48 (2H, brs), 2.45 (1H, d, *J* = 17.6), 2.39 (1H, dd, *J* = 13.2, 6.2 Hz), 2.33 (2H, brs), 2.31–2.23 (5H, m), 2.19–1.99 (6H, m), 1.99–1.94 (2H, m), 1.92 (1H, d, *J* = 13.2 Hz), 1.90–1.79 (2H, m), 1.76–1.63 (3H, m), 1.54–1.46 (4H, m), 1.46–1.26 (7H, m), 1.10 (3H, d, *J* = 6.0 Hz), 1.06 (3H, d, *J* = 6.6 Hz), 1.02 (1H, d, *J* = 12.0 Hz), 0.98 (3H, d, *J* = 7.2 Hz), 0.96 (3H, d, *J* = 7.2 Hz). ¹³C NMR (150 MHz, CD₃OD) δ : 172.8 (2C), 153.3, 153.2, 114.8, 114.2, 110.4, 105.7, 104.8, 98.5, 86.2, 80.7, 80.6, 79.1, 79.0, 78.0 (2C), 77.2 (2C), 76.3 (2C), 76.0, 75.8, 75.3, 75.0, 74.9, 74.7, 73.8, 73.7, 72.7 (2C), 69.5, 68.1, 68.0, 65.7, 53.6, 45.5, 44.9, 41.1, 39.7, 38.2, 38.1, 37.8, 37.5, 37.1, 36.7, 35.7, 33.0, 31.8, 31.2, 32.0, 30.8, 30.0, 29.0, 27.3,

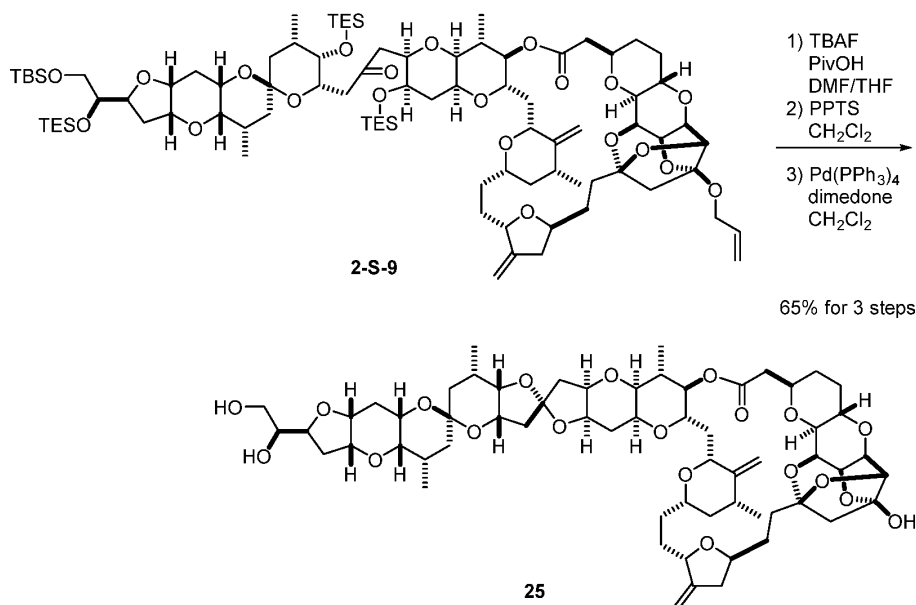
18.4, 18.1, 17.4, 15.8. FTIR (film): 3480, 2953, 2932, 2923, 1733, 1317, 1189, 1119, 1073, 1011, 995, 964, 914, 555 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{59}\text{H}_{82}\text{O}_{20}\text{Na}$, 1113.5297; found, 1113.5248.

Homohalichondrin C (25)



[00712] In a glove box, to a solution of iodide **2-13** (52.1 mg, 0.0501 mmol, 1 eq.) and thioester **2-16** (55.5 mg, 0.0651 mmol, 1.3 eq.) in DMI (0.40 mL) and EtOAc (80 μL) were added DTBMP (39.5 mg, 0.192 mmol, 4 eq.), Zn powder (18.9 mg, 0.289 mmol, 6 eq.), Cp_2ZrCl_2 (42.2 mg, 0.144 mmol, 3 eq.), and $\text{NiBr}_2\text{-dtbbpy}$ (7.0 mg, 0.0144 mmol, 30 mol%) at room temperature. After being stirred for 4 h at the same temperature, the reaction mixture was removed from glove box and diluted with Et_2O and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with Et_2O . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 9%, 13%, then 17% EtOAc in Hexanes) to give ketone **2-S-9** (70.5 mg, 0.0426 mmol, 85%) as a colorless amorphous solid. **2-S-9**: $[\alpha]_D^{20}$ -64.0 (c 0.800, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 5.75 (1H, dddd, J = 17.3, 10.5, 5.1, 5.1 Hz), 5.21 (1H, s), 5.15 (1H, ddd, J = 17.3, 1.8, 1.8 Hz), 5.12 (1H, s), 4.97 (1H, ddd, J = 10.5, 1.8, 1.8 Hz), 4.94 (1H, s), 4.86-4.79 (3H, m), 4.67 (1H, d, J = 10.8 Hz), 4.52 (1H, ddd, J = 10.2, 10.2, 4.8 Hz), 4.48 (1H, ddd, J = 10.2, 5.4, 5.4 Hz), 4.37 (1H, dd, J = 3.9, 3.0 Hz), 4.35 (1H, dd, J = 10.2, 1.8 Hz), 4.32 (1H, d, J = 3.6 Hz), 4.11 (1H, dd, J = 6.9, 5.1 Hz), 4.06-3.98 (4H, m), 3.84-3.79 (5H, m), 3.78-3.75 (2H, m), 3.73-3.67 (4H, m), 3.45 (1H, ddd, J = 4.5, 4.5, 4.5 Hz), 3.28 (1H, s), 3.21-3.16 (2H, m), 3.07 (1H, dd, J = 17.6, 6.3 Hz), 3.01 (1H, dd, J = 17.6, 6.6 Hz), 2.93 (1H, d, J = 2.4 Hz), 2.80-2.76 (2H, m), 2.57 (1H, dd, J = 9.9, 2.1 Hz), 2.47 (1H, d, J = 16.2 Hz), 2.40-2.23 (9H, m), 2.23-

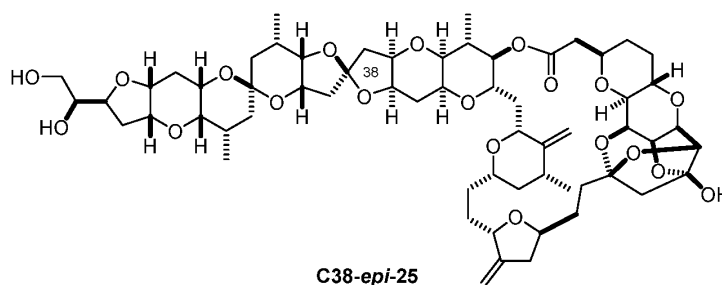
2.05 (6H, m), 1.98 (1H, dddd, $J = 14.1, 11.7, 4.0, 4.0$ Hz), 1.89-1.82 (2H, m), 1.77-1.68 (4H, m), 1.61 (1H, ddd, $J = 14.4, 4.8, 4.8$ Hz), 1.58 (1H, dd, $J = 12.9, 3.9$ Hz), 1.55-1.47 (5H, m), 1.36-1.32 (3H, m), 1.19 (3H, d, $J = 7.8$ Hz), 1.11 (9H, t, $J = 8.0$ Hz), 1.09 (3H, d, $J = 6.6$ Hz), 1.07 (9H, t, $J = 8.4$ Hz), 1.05 (9H, t, $J = 8.4$ Hz), 1.02-1.01 (12H, m), 0.96 (3H, d, $J = 7.2$ Hz), 0.77 (6H, q, $J = 7.8$ Hz), 0.69-0.65 (12H, m), 0.14 (3H, s), 0.13 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 206.8, 171.3, 153.0, 152.6, 134.6, 116.6, 116.2, 109.3, 104.39, 103.8, 96.9, 83.3, 79.1, 78.4, 78.0, 77.9, 77.7, 76.8, 76.0, 75.5, 75.1, 75.0, 74.9, 74.7, 74.22, 74.16, 74.1, 73.8, 73.6, 73.0, 70.4, 69.4, 68.4, 66.2, 65.9, 64.6, 63.8, 51.7, 46.8, 46.3, 43.9, 41.2, 39.5, 39.3, 38.6, 37.7, 37.6, 36.5, 36.4, 35.9, 35.3, 32.5, 31.6, 31.1, 30.72, 30.67, 30.5, 29.6, 28.6, 26.2, 18.7, 18.6, 18.1, 17.6, 16.4, 7.5, 7.4, 7.3, 6.0, 5.7, 5.3, -5.1, -5.3 ppm. FTIR (film): 2953, 2928, 2875, 1730, 1554, 1459, 1372, 1310, 1238, 1185, 1155, 1077, 1034, 1012, 834, 740 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{88}\text{H}_{149}\text{O}_{21}\text{Si}_4$, 1653.9663; found, 1653.9702.



[00713] A buffered TBAF solution was prepared by mixing TBAF solution (TCI #T1125; 0.43 mL of 1 M in THF, 0.43 mmol, 10 eq.) and PivOH (22.0 mg, 0.215 mmol, 5 eq.). To a stirred solution of ketone **2-S-9** (70.5 mg, 0.0426 mmol) in DMF (2.1 mL) were added the buffered TBAF solution at room temperature. After being stirred for 8 h at the same temperature, CaCO_3 (1.5 g) and DOWEX 50WX8-400 (1.5 g) were added. After being stirred for 1 h at the same temperature, the resulted mixture was diluted with EtOAc and filtered through a pad of Celite. The filter cake was washed with EtOAc thoroughly. The filtrate was concentrated under reduced pressure to give a crude tetraol, which was used in the next step

without further purification. To a stirred solution of above tetraol (calculated as 0.0426 mmol, 1 eq.) in CH_2Cl_2 (4.3 mL) was added PPTS (53.5 mg, 0.213 mmol, 5 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was directly subjected to column chromatography on amino silica gel (CH_2Cl_2 then 25%, 50%, 75%, then 100% EtOAc in Hexanes then 2% MeOH in EtOAc) to give a crude spiro ketal, which was used in the next step without further purification.

To a mixture of above crude spiro ketal (calculated as 0.0426 mmol, 1 eq.), dimedone (11.9 mg, 0.0849 mmol, 2 eq.), and $\text{Pd}(\text{PPh}_3)_4$ (4.9 mg, 0.00424 mmol, 10 mol%) was added CH_2Cl_2 (4.3 mL) at room temperature. After being stirred for 8 h at the same temperature, the resulted solution was directly subjected to column chromatography on amino silica gel (CH_2Cl_2 then 25%, 50%, 77%, then 100% EtOAc in Hexanes then 3% MeOH in EtOAc) to give a crude Homohalichondrin C with its C-38 epimer. The mixture was purified by YAMAZEN purification system with ODS column (Rf gradient: 10% MeCN in H_2O to 100% MeCN) to give Homohalichondrin C (**25**) (31.7 mg, 0.0278 mmol, 65% for 3 steps) as a colorless solid and 38-*epi*-Homohalichondrin C (**C38-*epi*-25**) (5.1 mg, 0.00448 mmol, 11% for 3 steps) as a colorless solid. **Homohalichondrin C (25)**: $[\alpha]_D^{20}$ -57.9 (*c* 0.53, CH_2Cl_2). ^1H NMR (600 MHz, CD_3OD) δ : 5.06 (1H, d, J = 1.8 Hz), 5.01 (1H, s), 4.88 (1H, s), 4.81 (1H, d, J = 1.2 Hz), 4.63 (1H, dd, J = 7.8, 4.8 Hz), 4.43 (1H, d, J = 9.6 Hz), 4.40 (1H, s), 4.34-4.28 (3H, m), 4.25-4.22 (2H, m), 4.17 (1H, d, J = 4.2 Hz), 4.13-4.09 (2H, m), 4.07 (1H, dd, J = 7.8, 7.8 Hz), 4.02 (1H, s), 3.95 (1H, s), 3.90-3.86 (3H, m), 3.70 (1H, dd, J = 10.5, 10.5 Hz), 3.66 (1H, dd, J = 2.7, 2.7 Hz), 3.60 (1H, d, J = 11.4 Hz), 3.59-3.57 (3H, m), 3.50 (1H, dd, J = 10.5, 5.1 Hz), 3.21 (1H, dd, J = 7.2, 4.8 Hz), 3.12 (1H, d, J = 2.4 Hz), 2.95 (1H, d, J = 10.2, 1.8 Hz), 2.82-2.79 (1H, m), 2.55 (1H, dd, J = 17.9, 9.3 Hz), 2.45 (1H, dd, J = 17.9, 2.4 Hz), 2.39 (1H, dd, J = 13.2, 6.0 Hz), 2.37-2.24 (8H, m), 2.20-1.96 (11H, m), 1.90 (1H, ddd, J = 15.6, 4.8, 4.8 Hz), 1.84-1.80 (2H, m), 1.74 (1H, s), 1.72 (1H, s), 1.68-1.64 (2H, m), 1.51-1.29 (9H, m), 1.10 (3H, d, J = 6.0 Hz), 1.05 (3H, d, J = 7.2 Hz), 1.04-0.99 (1H, m), 0.95 (3H, d, J = 7.2 Hz), 0.94 (3H, d, J = 6.6 Hz) ppm. ^{13}C NMR (125 MHz, CDCl_3) δ : 171.2, 151.7, 151.4, 113.2, 112.4, 109.3, 104.5, 104.2, 96.6, 84.4, 79.8, 79.4, 77.6, 77.1, 76.3, 76.2, 75.3, 75.1, 74.8, 74.7, 74.4, 73.8, 73.7, 73.5, 72.8, 72.0, 71.2, 70.8, 68.1, 66.1, 65.3, 63.6, 52.4, 43.4, 42.5, 40.4, 38.7, 37.2, 36.94, 36.89, 36.8, 36.5, 36.0, 34.9, 32.0, 31.3, 30.6, 30.0, 29.4, 29.0, 28.9, 27.7, 25.8, 18.0, 17.7, 17.1, 15.0 ppm. FTIR (film): 3422, 2926, 2873, 1736, 1436, 1310, 1186, 1117, 1074, 1021, 995, 910, 755 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{61}\text{H}_{86}\text{O}_{20}\text{Na}$, 1161.5605; found, 1161.5587.



[00714] 38-*epi*-Homohalichondrin C (C38-*epi*-25): $[\alpha]_D^{20}$ -88.2 (*c* 0.34, CH₂Cl₂). ¹H NMR (600 MHz, CD₃OD) δ : 5.04 (1H, d, *J* = 1.8 Hz), 4.95 (1H, d, *J* = 2.4 Hz), 4.87 (1H, s), 4.81 (1H, s), 4.72 (1H, dd, *J* = 10.2, 6.6 Hz), 4.42 (1H, d, *J* = 10.8 Hz), 4.38 (1H, s), 4.36 (1H, ddd, *J* = 10.4, 10.4, 4.5 Hz), 4.32-4.28 (2H, m), 4.23 (1H, ddd, *J* = 10.1, 5.9, 4.2 Hz), 4.17-4.09 (4H, m), 4.08-4.05 (1H, m), 4.03 (1H, s), 3.98 (1H, dd, *J* = 4.8, 1.8 Hz), 3.89-3.84 (3H, m), 3.81 (1H, dd, *J* = 2.4, 2.4 Hz), 3.63-3.56 (4H, m), 3.54 (1H, d, *J* = 3.0 Hz), 3.50 (1H, dd, *J* = 10.2, 5.4 Hz), 3.17 (1H, dd, *J* = 9.0, 6.6 Hz), 3.15 (1H, s), 2.96 (1H, dd, *J* = 9.9, 2.1 Hz), 2.83-2.79 (1H, m), 2.54 (1H, dd, *J* = 17.4, 8.4 Hz), 2.47 (1H, dd, *J* = 17.4, 2.7 Hz), 2.36-1.89 (23H, m), 1.83 (1H, ddd, *J* = 12.0, 12.0, 2.4 Hz), 1.78-1.75 (1H, m), 1.72-1.67 (2H, m), 1.64 (1H, ddd, *J* = 12.0, 3.0, 3.0 Hz), 1.58-1.53 (1H, m), 1.49-1.28 (8H, m), 1.10 (3H, d, *J* = 6.0 Hz), 1.05-0.98 (1H, m), 1.00 (3H, d, *J* = 7.2 Hz), 0.998 (3H, d, *J* = 6.6 Hz), 0.96 (3H, d, *J* = 7.2 Hz) ppm. ¹³C NMR (125 MHz, CD₃OD) δ : 172.9, 153.3, 152.8, 115.6, 114.2, 110.5, 105.1, 104.7, 98.0, 86.2, 80.3, 79.8, 79.2, 78.9, 78.5, 78.4, 77.8, 76.6, 76.4, 76.1, 76.0, 75.9, 75.8, 75.2, 75.1, 74.9, 74.8, 74.4, 73.2, 72.9, 69.4, 68.4, 65.3, 65.1, 53.7, 45.6, 45.0, 44.8, 41.2, 39.6, 38.6, 38.3, 38.2, 37.5, 37.2, 36.0, 33.3, 31.9, 31.7, 31.2, 30.9, 30.2, 28.9, 26.8, 18.4, 17.7, 15.2 ppm. FTIR (film): 34.7, 2925, 2872, 1736, 1662, 1553, 1436, 1311, 1188, 1117, 1075, 1023, 996, 898, 735 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₆₁H₈₆O₂₀Na, 1161.5605; found, 1161.5618.

Synthesis of Right Halves

[00715] The synthesis of C27-C37 building block is summarized in *Figure 9A*. First, a catalytic, asymmetric Ni/Cr-mediated reaction was used to couple aldehyde **4-2** with methyl β -iodoacrylate (**4-3**) in the presence of the Cr-catalyst (10 mol%) prepared from (*R*)-**4-E** (*Figure 9B*), to furnish the allylic alcohol in 93% yield with 19:1 stereoselectivity (see, *e.g.*, Namba, K.; Kishi, Y. *Org. Lett.* **2004**, *6*, 5031; Guo, H.; Dong, C.-G.; Kim, D.-S.; Urabe, D.; Wang, J.; Kim, J. T.; Liu, X.; Sasaki, T.; Kishi, Y. *J. Am. Chem. Soc.* **2009**, *131*, 15387). Second, an oxy-Michael reaction was used to construct the tetrahydropyran ring with the

desired stereochemistry at C29. The previous work suggested that this process could be achieved in a highly stereoselective manner; indeed, on treatment of **4-4** with K₃PO₄/18-Crown-6/toluene, oxy-Michael cyclization smoothly took place, to furnish the desired stereoisomer exclusively (see, *e.g.*, Aicher, T. D.; Buszek, K. R.; Fang, F. G.; Forsyth, C. J.; Jung, S. H.; Kishi, Y.; Scola, P. M. *Tetrahedron Lett.* **1992**, 33, 1549). The methyl ester was then reduced with DIBAL, to give aldehyde **5**, the substrate for the next catalytic, asymmetric Ni/Cr-mediated coupling reaction to form the C19-C20 bond.

[00716] *Figure 10A* summarizes the synthesis of C20-C37 building block from **4-5**. The crucial transformation in this sequence was the catalytic, asymmetric Ni/Cr-mediated coupling to introduce the chiral center at C27, followed by reductive cyclization to introduce the chiral center at C23. The overall stereochemistry-outcome of the proposed transformation deserves a comment. The stereochemistry at C27 was introduced under the influence of a chiral Cr-catalyst prepared from a chiral sulfonamide. Via the toolbox approach, (*S*)-**4-F** was identified as the best ligand for the substrates closely related to **4-6** (see, *e.g.*, Guo, H.; Dong, C.-G.; Kim, D.-S.; Urabe, D.; Wang, J.; Kim, J. T.; Liu, X.; Sasaki, T.; Kishi, Y. *J. Am. Chem. Soc.* **2009**, 131, 15387). The same Cr-catalyst was found equally effective for the present case; the (**4-5** → **4-6**)-coupling was conducted in the presence of 10 mol% Cr-catalyst and 2 mol% (Et)₂Phen(H)₂•NiCl₂, to furnish the expected allylic alcohol with >40:1 stereoselectivity (¹H NMR). To facilitate ¹H NMR analysis, the authentic sample of undesired allylic alcohol was prepared with the Cr-catalyst prepared from (*R*)-sulfonamide.

[00717] The next reductive cyclization was expected stereoselectively to yield the desired product, as demonstrated on the closely related substrates (see, *e.g.*, Lewis, M. D.; Cha, J. K.; Kishi, Y. *J. Am. Chem. Soc.* **1982**, 104, 4976; Dong, C.-G.; Henderson, J. A.; Kaburagi, Y.; Sasaki, T.; Kim, D.-S.; Kim, J. T.; Urabe, D.; Guo, H.; Kishi, Y. *J. Am. Chem. Soc.* **2009**, 131, 15642). Nevertheless, there was a concern on the proposed transformation, because of the presence of dioxasilinane group. Experimentally, it was found that the reductive cyclization did give the desired product as the major product, but accompanied with by-products apparently derived through reactions on the dioxasilinane group. In order to avoid the complication due to the undesired side-reaction(s), the dioxasilinane group was first deprotected with TBAF-AcOH treatment and then subjected to reductive-cyclization, to furnish practically the single product. The undesired allylic alcohol was prepared via the coupling of **4-5** with **4-6** in the presence of the Cr-catalyst derived from (*S*)-**4-A**. The product at this stage was a triol, as the C30 MPM group was cleaved off during the process. With a

standard procedure, the alcohols at C35 and C37 were selectively protected, to give six-membered acetonide **4-7**.

[00718] The observed stereochemistry-outcome was explained by a stereoelectronic effect, coupled with conformational analysis. Due to a stereoelectronic effect, the reducing-reagent approached the oxonium ion preferentially from the direction resulting in a trans-diaxial relationship between the newly formed bond and a lone pair of oxygen-electrons, the oxonium ion **4-A** (*Figure 10B*). For clarity, the C27-olefinic carbon is replaced with a saturated carbon in the oxonium ions **4-A-C**, and the oxonium ion **4-C** is shown its antipode. Similarly, the approach from the bottom-direction also met with the stereoelectronic effect. However, the top-approach was favored over the bottom-approach, because the former approach led directly to the product in chair-conformation, whereas the latter approach led to the product in boat-conformation. Apparently, there was no serious steric hindrance for the reagent to approach from the top-face. Reductive cyclization of the undesired allylic alcohol gave a 2:1 mixture of tetrahydropyrans. The observed result was again explained again by a stereoelectronic effect, coupled conformational analysis, cf., oxonium ions **4-B** and **4-C** in *Figure 10B*. Two modes of reduction depicted on **4-B** and **4-C** led directly to a product in chair conformation, but both approaches suffer from the 1,3-diaxial-like interaction with either C25-Me or C29-CH₂ group. This analysis explained the poor stereoselectivity in reductive cyclization in the undesired allylic alcohol series. Because of the poor-selectivity observed in the reductive cyclization of undesired allylic alcohol, the overall stereoselectivity of **4-5** + **4-6** → **4-7** became higher than the stereoselectivity achieved by the catalytic, asymmetric Ni/Cr-mediated coupling. Finally, the ethyl ester in **4-7** was reduced with DIBAL, to furnish aldehyde **4-8**, the synthetic intermediate for the next Ni/Cr-mediated coupling reaction to form the C19-C20 bond of halichondrins A-C. Notably, without protection of the C30 hydroxyl group, the DIBAL reduction smoothly and cleanly proceeded to give the desired **4-8** in 89% yield, along with ~5% of the over-reduced primary alcohol.

[00719] The C19-C20 bond with a catalytic, asymmetric Ni/Cr-mediated coupling of **4-8** with **4-9-B** was formed (*Figure 11*) (see, e.g., Yan, W.; Li, Z.; Kishi, Y. *J. Am. Chem. Soc.* **2015**, *137*, 6219; Li, Z.; Yan, W.; Kishi, Y. *Am. Chem. Soc.* **2015**, *137*, 6226). It should be noted that aldehyde **4-8** bore a free hydroxyl group at C30. There was no precedent for demonstrating that the catalytic cycle of Ni/Cr-mediated reaction could function with a substrate with a free hydroxyl group. Nevertheless, this possibility was pursued, because two synthetic steps, i.e., protection and deprotection of the C30 hydroxyl group, could be saved in

this manner. In this connection, it should be noted that catalytic asymmetric Ni/Cr-mediated coupling uses Cp_2ZrCl_2 as the agent dissociating a product from a Cr-complex (see, *e.g.*, Namba, K.; Kishi, Y. *Org. Lett.* **2004**, 6, 5031) thereby suggesting a possibility of utilizing Cp_2ZrCl_2 as a masking agent for the free hydroxyl group *in situ*. This possibility was experimentally tested; the catalytic, asymmetric coupling of **4-8** with **4-9-B** smoothly proceeded in addition of Cp_2ZrCl_2 (2.5 equiv.) and 2,6-di-*t*-butyl-4-methylpyridine (2.5 equiv.), to furnish the desired product in an excellent yield.

[00720] Being encouraged with the successful coupling of **4-8** bearing a free hydroxyl group, a coupling with the vinyl iodide bearing a free carboxylic acid at C1 was tested. Amazingly, the catalytic, asymmetric Ni/Cr-mediated coupling of **4-8** did give the desired product in ~50% yield. The coupling with **4-9-B** was used for further studies.

[00721] Adopting the toolbox approach, a satisfactory sulfonamide ligand was identified. The ligand screening was conducted in the presence of Cr-catalyst, prepared from CrCl_2 (10 mol%), sulfonamide (13 mol%), proton scavenger (12 mol%), and $(\text{Me})_6\text{Phen}\bullet\text{NiCl}_2$ in MeCN at room temperature. Through this screening, three exemplary sulfonamides emerged, *i.e.*, (*S*)-**4-I** (*dr* = 19:1), (*S*)-**4-G** (*dr* = 29:1), and (*S*)-**4-H** (*dr* = 24:1). The coupling yield with these three ligands was then estimated from the overall yield of **4-8** → **4-11-B**, *i.e.*, 73% with (*S*)-**4-I**, 65% with (*S*)-**4-G**, and 67% with (*S*)-**4-H**. These overall yields were based on the experiments starting with 1.65 g, 250 mg, and 250 mg of **7** with (*S*)-**4-I**, (*S*)-**4-G**, and (*S*)-**4-H**, respectively. Based on this result, sulfonamide (*S*)-**4-I** was used for preparative purpose. It is noteworthy that, unlike the first and second couplings, this Ni/Cr-mediated coupling utilized the structurally complex nucleophile. Remarkably, the coupling efficiency was excellent even with use of the molar ratio **4-7**:**4-8** = 1.0:1.1.

[00722] The next task was an $\text{S}_{\text{N}}2$ -cyclization of the C20 alcohol to the C17 chloride to form the methylenetetrahydrofuran ring. Various bases cleanly achieved this five-membered-ring forming cyclization, unless there was a base-labile functional group(s) in the substrate (see, *e.g.*, Lee, J. H.; Li, Z.; Osawa, A.; Kishi, Y. *J. Am. Chem. Soc.* **2016**, 138, 16248). After the base-induced cyclization, the methyl ester at C1 was hydrolyzed with aqueous base, to furnish seco-acid **4-10-B**. For example, in the halichondrin-A synthesis, this transformation was achieved in 2 separate steps, *i.e.*, $\text{AgOTf}/\text{Ag}_2\text{O}$ in THF and LiOH in aqueous MeOH. In this work, this transformation was carried out in one-pot. The seco-acid thus obtained was subjected to macrolactonization with Shiina's reagent, to furnish crystalline C1-C37 mactrolactone acetone **4-11-B** in an excellent yield (see, *e.g.*, Shiina, I.; Kubota, M.; Ibuka,

R. *Tetrahedron Lett.* **2002**, 43, 7535; Shiina, I.; Mukaiyama, T. *Chem. Lett.* **1994**, 677; Shiina, I. *Bull Chem. Soc. Jpn.* **2014**, 87 196). The structure of **4-11-B** was confirmed via an X-ray analysis of its C35/C37-diol, *i.e.*, the product at step c-1 (*Figure 12*).

[00723] For the preparative purpose, the transformation of **4-8** + **4-9-B** → **4-10-B** → **11-B** was carried out without purification/isolation of the intermediates. The macrolactone **4-11-B** was isolated by silica gel flash-chromatography (neutral silica gel) in 73% overall yield from **4-8** in multi-gram scales.

[00724] A ¹H NMR analysis indicated that **4-11-B** thus obtained was contaminated with ~5% of its C20-epimer, thereby showing that the overall stereoselectivity for this transformation was ~20:1. The material was carried on of this purity; namely, **4-11-B** was transformed to iodide **4-12-B** with 4-steps/2-pots procedure, *i.e.*, (1) *p*-TsOH/MeOH-CH₂Cl₂, (2) Tf₂O/lutidine/ CH₂Cl₂, followed by addition of TESOTf and then NaI in DMF. The product was isolated by silica gel flash-chromatography (neutral silica gel), to furnish **4-12-B** in 92% overall yield. ¹H NMR analysis indicated that **4-12-B** thus obtained was contaminated with ~5% of the C20-epimer. Although **4-12-B** was crystalline, it was again difficult to remove the minor stereoisomer by recrystallization. Therefore, the minor C20 diastereomer was removed by preparative HPLC (ZORBAX SIL; 300~500 mg injection), to furnish **4-12-B** (mp: 158~160 °C), which was used for the synthesis of halichondrins in the B-series.

[00725] *Figure 13* summarizes the synthesis of right half in the halichondrin-A series. The synthesis followed the synthetic route developed in the halichondrin B series, with two modifications. First, **4-9-A**, instead of **4-9-B**, was used. Second, the C35/C37-protecting group in **8** was switched to the corresponding bis-TBS before the Ni/Cr-mediated coupling reaction, because the anisylidene group in **4-9-A** was acid-labile and could not survive under the aqueous acidic condition; step c-1 in *Figure 11*. Once again, the minor stereoisomer that originated from the Ni/Cr-mediated coupling was removed by preparative HPLC.

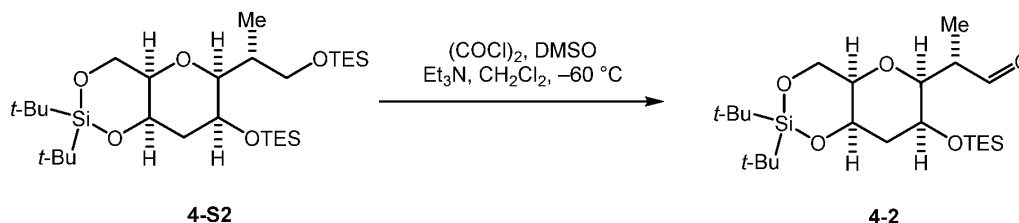
[00726] *Figure 14* summarizes the synthesis of right half in the halichondrin C series. The synthesis followed the synthetic route developed in the halichondrin B series, except one; **4-9-C**, instead of **4-9-B**, was used for the Ni/Cr-mediated coupling. The synthesis proceeded without any unexpected difficulty, to furnish the right half **4-12-C** (mp: 84-6 °C) in the overall efficiency very similar to that in the halichondrin B series. Once again, the minor stereoisomer that originated from the Ni/Cr-mediated coupling was removed by preparative HPLC.

[00727] In the syntheses reported, seven chiral centers were introduced at C17, C20, C23, C25, C27, C29, and C30. The availability of the authentic sample of undesired stereoisomer(s) should facilitate the analysis of purity and stereoselectivity for a given step. Among them, the chiral centers at C20, C27, and C30 were introduced under the influence of chiral Cr-catalysts derived from (*R*)-**4-E**, (*R*)-**4-F**, and (*S*)-**4-I**. Therefore, the authentic sample of undesired minor stereoisomers formed in each of catalytic, asymmetric Ni/Cr-mediated couplings was readily prepared via the coupling in the presence of (*S*)-**4-E**, (*S*)-**4-F**, and (*R*)-**4-I**. In practice, the antipode was prepared with use of the Cr-catalyst prepared from (*S*)-**4-E** or (*R*)-**4-E** for each coupling. The chiral centers at C17 and C25 originated from the chiral centers present in C1-C19 and C20-C26 building blocks. Thus, use of C17-*epi*-C1-C19 building block gave the stereoisomer at C17, whereas use of the antipode of C20-C26 building block gave the minor stereoisomers at C25. The stereochemistry analysis was carried out in reference to these authentic samples.

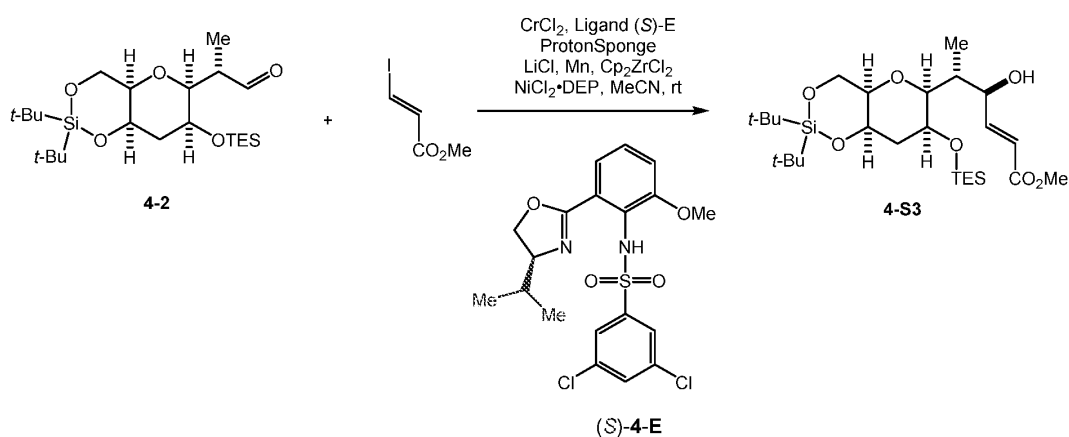
[00728] Right halves of halichondrins A-C were synthesized by coupling the common C20-C37 building block **4-8** with the C1-C19 building blocks **4-9-A**, **4-9-B**, and **4-9-C**, respectively. Catalytic, asymmetric Ni/Cr-mediated coupling was used for three C-C bond formations. For all the cases, the stereochemistry was introduced under the influence of Cr-catalysts prepared from chiral sulfonamides, identified via the toolbox approach. For (**4-2** + **4-3**)-, (**4-5** + **4-6**)-, and (**4-7** + **4-8**)-couplings, the stereoselectivity of 19:1, >40:1, and ~20:1 was achieved by the Cr-catalysts prepared from (*R*)-**4-E**, (*R*)-**4-F**, and (*S*)-**4-I**, respectively. Unlike the first and second couplings, the third coupling utilized the structurally complex nucleophile. It was demonstrated that the coupling efficiency was excellent even with use of the molar ratio **4-8** : **4-9A~C** = 1.0:1.1. In addition, third coupling was achieved with the substrate bearing a free-hydroxyl group. The products obtained in the Ni/Cr-mediated couplings were converted to the right halves of halichondrins A-C in excellent overall yields. The right halves of halichondrins A-C (**4-12A** through **4-12C**) were synthesized in 28, 24, and 24 steps from commercial D-galactal in 13.4%, 21.1%, and 16.7% overall yields, respectively.

321

CDCl₃) δ : 81.1, 77.1, 68.2, 67.8, 65.5, 64.9, 38.8, 37.4, 27.8, 27.2, 23.3, 20.6, 13.2, 7.0, 6.8, 5.1, 4.4 ppm. FTIR (film): 2954, 2875, 1465, 1239, 1168, 1104, 1082, 1036, 1007, 927, 826, 772, 723, 649, 442 cm⁻¹. HRMS (ESI) m/z : [M+Na]⁺ calcd for C₂₉H₆₂O₅Si₃Na, 597.3797; found, 597.3807.

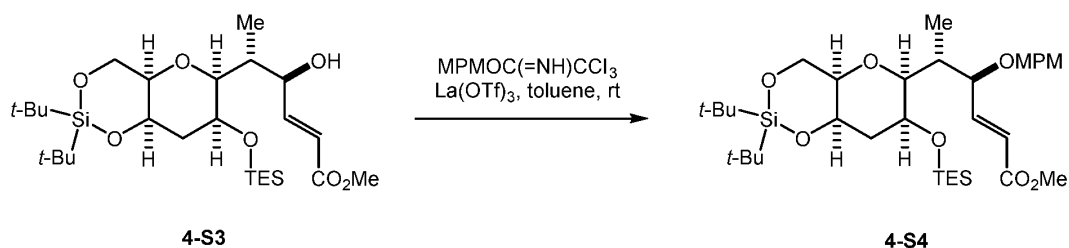


[00731] To a stirred solution of (COCl)₂ (7.5 mL, 88.6 mmol, 5 eq.) in CH₂Cl₂ (250 mL) was added a solution of DMSO (12.4 mL, 174 mmol, 10 eq.) in CH₂Cl₂ (10 mL) at -78 °C. After being stirred for 30 min at the same temperature, a solution of **4-S2** (10.0 g, 17.4 mmol, 1 eq.) in CH₂Cl₂ (30 mL) was introduced to the reaction mixture. After being stirred for 2 h at -60 °C, to the mixture was added Et₃N (42 mL, 305 mmol, 17 eq.) at -78 °C and warmed up to 0 °C over 30 min. After being stirred for 15 min at 0 °C, the mixture was quenched with sat. NH₄Cl aq. The organic layer was separated and the aqueous phase was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, and concentrated under reduced pressure. The residue was filtrated though a pad of silica gel (hexanes/EtOAc = 1:1) to give a crude aldehyde **4-2** as pale yellow oil. The obtained crude material was used in the next reaction without further purification.



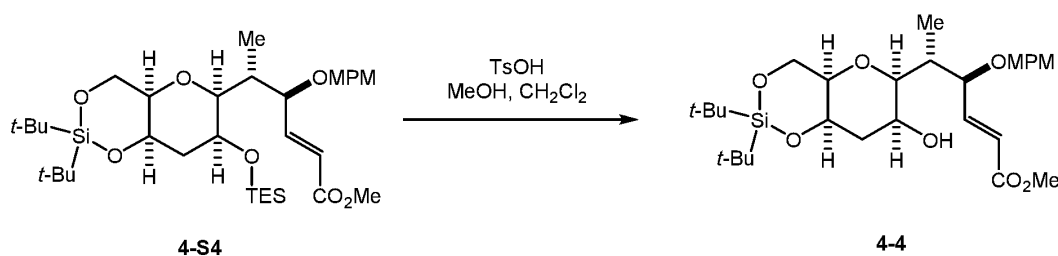
[00732] To a mixture of CrCl₂ (257 mg, 2.1 mmol, 12 mol%), (*S*)-sulfonamide ligand [*i*-Pr, PhCl₂, OMe] (1.08 g, 2.4 mmol, 14 mol%), and proton scavenger (522 mg, 2.4 mmol, 14 mol%) in a glove box was added MeCN (44 mL) and the resulting solution was stirred for 3 h

at room temperature. In a separate flask, the above crude **4-2** (calculated as 17.4 mmol, 1 eq.), methyl 3-iodoacrylate (3.7 g, 17.4 mmol, 1 eq.), $\text{NiCl}_2 \cdot \text{DMP}(\text{OMe})_2$ (138 mg, 0.348 mmol, 2 mol%), LiCl (1.48 mg, 34.8 mmol, 2 eq.), Mn (3.8 g, 70 mmol, 4 eq.), Cp_2ZrCl_2 (5.6 g, 19.1 mmol, 1.1 eq.), and 2,6-lutidine (4.1 mL, 34.8 mmol, 2 eq.) were mixed together and the Cr-complex solution (26 mL) was transferred to the flask. Additional methyl 3-iodoacrylate (1.8 g, 8.7 mmol, 0.5 eq.) and Cr-complex solution (11 mL) were added after 30 min. After being stirred for 45 min, to the mixture was added the remained Cr-complex solution (7 mL). After being stirred for 15 min, the reaction mixture was removed from the glove box and diluted with EtOAc. After florisil was added, the resultant suspension was stirred vigorously for 1 h. The mixture was filtered through a short pad of silica gel (EtOAc). After removal of solvent under reduced pressure, the residue was purified by flash column chromatography on silica gel (3% then 25% EtOAc in Hexanes) to give allylic alcohol **4-S3** (8.8 g, 16.2 mmol, 93% for 2 steps, *dr*: 16:1 based on integration ratio of ^1H NMR) as pale yellow oil. **4-S3**: $[\alpha]_D^{20} -1.3$ (*c* 1.0, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.18 (1H, dd, $J = 15.6, 3.6$ Hz), 6.32 (1H, dd, $J = 15.6, 2.4$ Hz), 4.39 (1H, brs), 4.15 (1H, dd, $J = 12.6, 1.2$ Hz), 3.94 (1H, dd, $J = 12.6, 2.4$ Hz), 3.89 (1H, brs), 3.66 (1H, ddd, $J = 4.2, 2.4, 1.2$ Hz), 3.43 (3H, s), 3.18 (1H, dd, $J = 6.6, 1.2$ Hz), 2.66 (1H, brs), 2.13 (1H, qdd, $J = 6.6, 6.6, 1.8$ Hz), 2.02 (1H, ddd, $J = 15.0, 2.4, 2.4$ Hz), 1.99 (1H, d, $J = 4.2$ Hz), 1.29 (1H, m), 1.28 (9H, s), 1.12 (9H, s), 1.10 (3H, d, $J = 6.6$ Hz), 1.06-1.00 (9H, m), 0.74-0.61 (6H, m) ppm. ^{13}C NMR (150 MHz, CDCl_3) δ : 166.6, 150.2, 128.3, 120.5, 82.2, 77.4, 72.1, 68.2, 67.7, 65.6, 51.1, 40.3, 38.7, 28.0, 27.6, 23.5, 20.9, 9.8, 7.3, 5.4 ppm. IR (film): 3501, 2952, 2876, 1725, 1706, 1659, 1474, 1167 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{27}\text{H}_{52}\text{O}_7\text{Si}_2\text{Na}$, 567.3144; found, 567.3157.



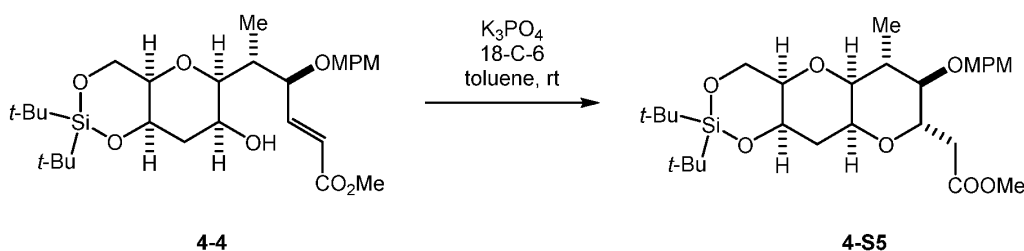
[00733] To a stirred suspension of allylic alcohol **4-S3** (5.65 g, 10.4 mmol, 1 eq.) and $\text{La}(\text{OTf})_3$ (640 mg, 1.1 mmol, 10 mol%) in toluene (21 mL) was added $\text{MPMOC}(\text{NH})\text{CCl}_3$ (6.0 g, 21 mmol, 2 eq.) by syringe pump over 5.5 h. After being stirred for additional 30 min, the mixture was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous phase was extracted with EtOAc. The combined organic layer was washed with

brine, dried over Na₂SO₄, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (3% then 17% EtOAc in Hexanes) to give MPM ether **S-3** as a partially separable mixture with reagent residues. **4-S4**: $[\alpha]_D^{20} +20.9$ (c 1.0, CHCl₃). ¹H NMR (600 MHz, CDCl₃) δ : 7.20 (2H, d, $J = 8.4$ Hz), 6.89 (1H, dd, $J = 16.2$, 6.0 Hz), 6.87 (2H, d, $J = 8.4$ Hz), 6.03 (1H, dd, $J = 16.2$, 1.2 Hz), 4.56 (1H, d, $J = 11.4$ Hz), 4.20 (1H, d, $J = 11.4$ Hz), 4.19 (1H, brs), 4.18 (1H, dd, $J = 12.0$, 2.4 Hz), 4.11 (1H, dd, $J = 12.0$, 1.2 Hz), 3.96 (1H, ddd, $J = 6.0$, 2.4, 1.2 Hz), 3.80 (3H, s), 3.77 (3H, s), 3.45 (1H, ddd, $J = 4.2$, 2.4, 1.2 Hz), 3.27 (1H, dd, $J = 7.2$, 1.2 Hz), 3.14 (1H, brs), 2.10 (1H, dqd, $J = 7.2$, 6.6, 2.4 Hz), 2.08 (1H, ddd, $J = 14.4$, 2.4, 2.4 Hz), 1.61 (1H, ddd, $J = 14.4$, 4.2, 4.2 Hz), 1.03 (9H, s), 1.01 (3H, d, $J = 6.6$ Hz), 1.00 (9H, s), 0.98-0.94 (9H, m), 0.65-0.50 (6H, m) ppm. ¹³C NMR (150 MHz, CDCl₃) δ : 166.7, 159.3, 147.7, 130.2, 129.3, 121.4, 113.7, 80.6, 77.9, 76.9, 70.5, 68.0, 67.5, 64.9, 55.2, 51.6, 38.6, 27.7, 27.2, 23.2, 20.5, 10.3, 7.0, 5.1 ppm. IR (film): 2950, 2875, 1726, 1659, 1612, 1513, 1168 cm⁻¹. HRMS (ESI) m/z : $[M+Na]^+$ calcd for C₃₅H₆₀O₈Si₂Na, 687.3719; found, 687.3706.



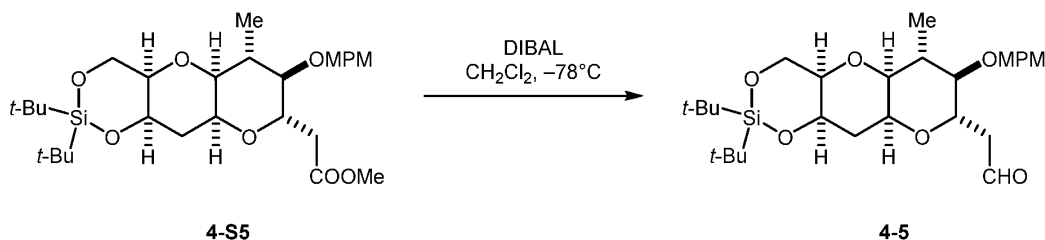
[00734] This selective deprotection of TES ether could be accomplished by following two different procedures. **Selective deprotection with TsOH·H₂O**: To a stirred solution of the above MPM-ether **4-S4** (calculated as 10.4 mmol, 1 eq.) in CH₂Cl₂ (20 mL) and MeOH (10 mL) was added TsOH·H₂O (10 mg, 0.0525 mmol, 0.5 mol%) at room temperature. After being stirred for 30 min at the same temperature, additional TsOH·H₂O (20 x 3 mg, 0.450 mmol, 3 mol%) was added every 30 min, and the resultant mixture was stirred for 4 h. The reaction mixture was quenched with sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (3% then 33% EtOAc in Hexanes) to give secondary alcohol **4-4** (5.04 g, 9.15 mmol, 88% for 2 steps) as colorless oil, which is contaminated with a small amount of reagent residue. **Selective deprotection with HFIP**: A solution of the above MPM-ether **4-S4** (~6 g) in 1,1,1,3,3,3-hexafluoro-2-propanol (40 mL)

and H₂O (4 mL) was stirred for 10 h at room temperature. The reaction mixture was concentrated and the resultant residue was purified by flash column chromatography on silica gel (3% then 33% EtOAc in Hexanes) to give secondary alcohol **4-4** (4.2 g, 7.63 mmol, ca. 79% in 2 steps) as colorless oil. **4-4**: $[\alpha]_D^{20} +18.8$ (*c* 1.0, CHCl₃). ¹H NMR (600 MHz, CDCl₃) δ : 7.20 (2H, d, *J* = 12.0 Hz), 6.95 (1H, dd, *J* = 16.2, 4.8 Hz), 6.86 (2H, d, *J* = 12.0 Hz), 6.07 (1H, dd, *J* = 16.2, 1.2 Hz), 4.54 (1H, d, *J* = 12.0 Hz), 4.35 (1H, dd, *J* = 3.0, 3.0 Hz), 4.22 (1H, dd, *J* = 12.6, 3.0 Hz), 4.19 (1H, d, *J* = 12.0 Hz), 4.18 (1H, m), 4.16 (1H, dd, *J* = 12.6, 1.2 Hz), 3.80 (3H, s), 3.76 (3H, s), 3.53 (1H, d, *J* = 10.8 Hz), 3.45 (1H, ddd, *J* = 10.8, 3.0, 3.0 Hz), 3.27 (1H, brs), 3.24 (1H, d, *J* = 8.4 Hz), 2.19 (1H, ddd, *J* = 14.4, 3.0, 3.0 Hz), 2.14 (1H, dqd, *J* = 8.4, 6.6, 2.4 Hz), 1.57 (1H, ddd, *J* = 14.4, 3.0, 3.0 Hz), 1.03 (9H, s), 1.02 (3H, d, *J* = 6.6 Hz), 1.02 (9H, s) ppm. ¹³C NMR (150 MHz, CDCl₃) δ : 166.7, 159.3, 148.1, 130.2, 129.4, 121.6, 113.8, 82.1, 77.1, 76.6, 70.6, 69.3, 68.6, 64.4, 55.3, 51.6, 38.3, 36.7, 27.7, 27.2, 23.1, 20.3, 10.9 ppm. IR (film): 3538, 2938, 2859, 1725, 1658, 1612, 1514, 1251 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₉H₄₆O₈SiNa, 573.2854; found, 573.2846.

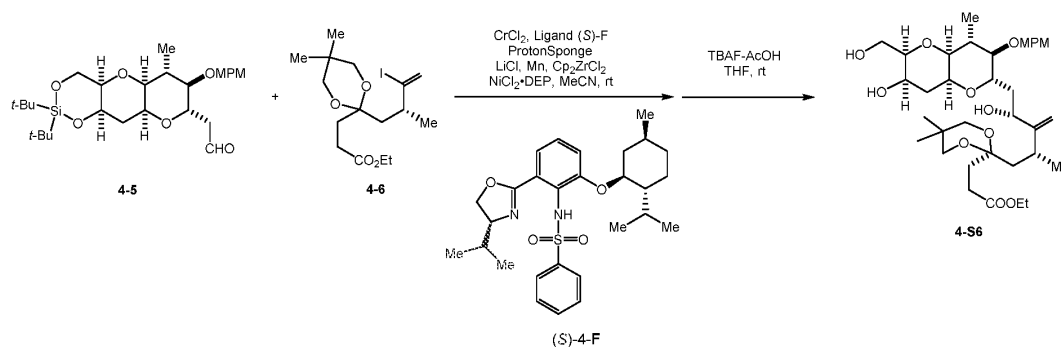


[00735] To a stirred solution of **4-4** (2.80 g, 5.08 mmol, 1 eq.) in toluene (340 mL) were added 18-crown-6 (2.69 g, 10.2 mmol, 2 eq.) and K₃PO₄ (21.6 g, 102 mmol, 20 eq.) at room temperature. After being stirred for 14 h at the same temperature, additional 18-crown-6 (671 mg, 2.54 mmol, 0.5 eq.) was added. After being stirred for 5 h at the same temperature, the reaction was quenched with sat. NH₄Cl aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography to give **4-S5** (2.64 g, 4.79 mmol, 94%) as colorless oil. **4-S5**: $[\alpha]_D^{20} -25.8$ (*c* 1.0, CHCl₃). ¹H NMR (600 MHz, CDCl₃) δ : 7.25 (2H, d, *J* = 8.4 Hz), 6.88 (1H, d, *J* = 8.4 Hz), 4.56 (1H, d, *J* = 10.2 Hz), 4.45 (1H, d, *J* = 10.2 Hz), 4.29 (1H, ddd, *J* = 7.8, 4.2, 4.2 Hz), 4.14 (1H, dd, *J* = 12.0, 6.0 Hz), 4.07 (1H, dd, *J* = 12.0, 3.0 Hz), 3.98 (1H, ddd, *J* = 9.6, 7.8, 3.6 Hz), 3.82-3.77 (4H, m), 3.66 (3H, s), 3.60 (1H, ddd, *J* = 6.0, 4.2, 3.0

Hz), 3.40 (1H, dd, $J = 7.8, 4.8$ Hz), 3.14 (1H, dd, $J = 9.6, 9.6$ Hz), 2.67 (1H, dd, $J = 15.6, 3.6$ Hz), 2.48 (1H, dd, $J = 15.6, 7.8$ Hz), 2.11 (1H, ddd, $J = 13.2, 7.8, 7.8$ Hz), 2.00-1.92 (2H, m), 1.21 (3H, d, $J = 7.2$ Hz), 1.03 (9H, s), 1.01 (9H, s) ppm. ^{13}C NMR (150 MHz, CDCl_3) δ : 171.6, 159.4, 129.9, 129.7, 113.9, 80.4, 78.6, 73.7, 73.0, 72.4, 68.9, 66.9, 65.1, 55.3, 51.6, 39.5, 37.8, 31.9, 27.4, 27.1, 22.2, 20.8, 16.1 ppm. IR (film): 2934, 2858, 1740, 1612, 1514, 1250 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{29}\text{H}_{46}\text{O}_8\text{SiNa}$, 573.2854; found, 573.2847.

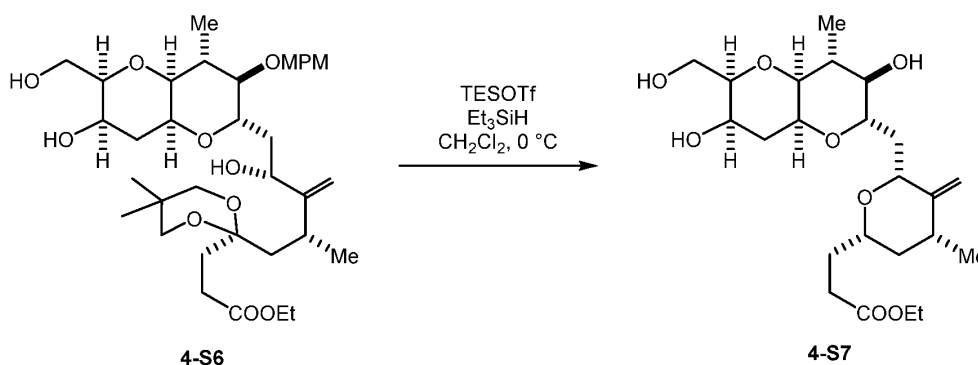


[00736] To a stirred solution of **4-S5** (5.0 g, 9.07 mmol, 1 eq.) in toluene (100 ml) was added DIBAL solution (11.3 mL of 1M in hexanes, 11.3 mmol, 1.2 eq.) dropwise at -78°C . After being stirred for 1.5 h at the same temperature, the reaction was quenched with acetone (0.3 mL). The mixture was stirred for 15 min and sat. Rochelle's salt aq. was added. After being stirred for 3 h at room temperature, the mixture was diluted with EtOAc. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (5%, 20% then 33% EtOAc in Hexanes) to give **4-5** (4.37 g, 8.49 mmol, 94%) as a colorless amorphous solid. **4-5**: $[\alpha]_D^{20} -37.3$ (c 1.0, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 9.51 (1H, t, $J = 2.4$ Hz), 7.21 (2H, m), 6.82 (2H, m), 4.33 (1H, d, $J = 10.6$ Hz), 4.24 (1H, d, $J = 10.6$ Hz), 4.17 (1H, dd, $J = 12.3, 3.5$ Hz), 4.11 (1H, m), 3.99-3.91 (2H, m), 3.31 (3H, s), 3.26 (1H, m), 3.0 (1H, dd, $J = 7.0, 4.1$ Hz), 2.91 (1H, m), 2.81 (1H, dd, $J = 10.6, 9.4$ Hz), 2.40 (1H, m), 2.34 (1H, m), 2.18-2.08 (2H, m), 1.54 (1H, dt, $J = 14.1, 5.3$ Hz), 1.24 (9H, s), 1.18 (3H, d, $J = 7.0$ Hz), 1.13 (9H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 200.1, 160.6, 131.2, 130.4, 114.7, 81.4, 80.8, 74.5, 74.4, 73.3, 69.1, 68.2, 66.0, 55.4, 47.9, 40.9, 34.2, 28.4, 28.1, 23.6, 21.6, 17.3 ppm. IR (film): 2933, 2857, 1725, 1514, 1250, 1078 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{28}\text{H}_{44}\text{O}_7\text{SiNa}$, 543.7272; found, 543.2750.



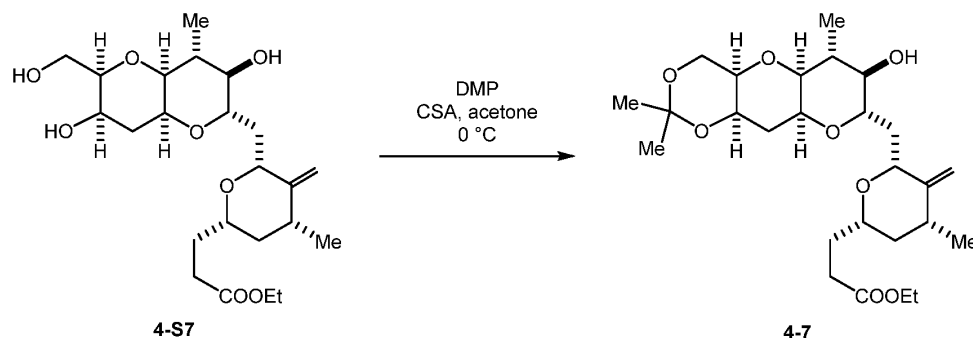
[00737] In a glove box, to a mixture of CrCl_2 (104 mg, 0.849 mmol, 10 mol%), (*S*)-sulfonamide ligand (465 mg, 0.934 mmol, 11 mol%), and proton scavenger (200 mg, 0.934 mmol, 11 mol%) was added MeCN (28 mL) and the resulted solution was stirred for 1 h at room temperature. In a separate flask, **4-5** (4.37 g, 8.49 mmol, 1 eq.), **4-6** (5.22 g, 12.7 mmol, 1.5 eq.), LiCl (720 mg, 17.0 mmol, 2 eq.), Mn (932 mg, 17.0 mmol, 2 eq.), Cp_2ZrCl_2 (2.48 g, 8.49 mmol, 1 eq.), and $\text{NiCl}_2\cdot\text{DEP}$ (331 mg, 0.849 mmol, 10 mol%) were mixed together. Then Cr complex solution was transferred to the flask, and the resulting mixture was stirred vigorously for 3 h at room temperature. The reaction mixture was removed from the glove box and diluted with EtOAc (100 mL). Aqueous potassium serinate (0.5 M, 100 mL) and sat. NaHCO_3 aq. (100 mL) were added, and the resulting mixture was stirred for 1 h. The suspension was filtered through a pad of Celite. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude allylic alcohol, which was used in the next reaction without further purification. A buffered TBAF solution was prepared by mixing TBAF solution (21 ml of 1M in THF, 21.0 mmol, 2.5 eq.) and AcOH (0.608 mL, 10.6 mmol, 1.25 eq.) at room temperature. To a stirred solution of the crude alcohol (calculated as 8.49 mmol) in THF (100 mL) was added the TBAF-AcOH solution at 0 °C. The reaction mixture was stirred for 1 h at the same temperature, and then quenched with sat. NH_4Cl aq. The mixture was diluted with EtOAc. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (20%, 50% then 100% EtOAc in Hexanes) to give **4-S6** (4.45 g, 6.69 mmol, 79% for 2 steps) as colorless oil. **4-S6**: $[\alpha]_D^{20}$ -23.9 (*c* 1.0, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.29 (2H, m), 6.86 (2H, m), 5.45 (1H, s), 5.01 (1H, s), 4.52 (1H, brd, J = 5.9 Hz), 4.47 (1H, d, J = 11.2 Hz), 4.43 (1H, m), 4.38 (1H, d, J = 11.2 Hz), 4.02-3.95 (3H, m), 3.87 (2H, m), 3.57 (1H, brs), 3.43 (1H, brs),

3.37-3.30 (4H, m), 3.25 (2H, m), 3.07 (1H, m), 2.98 (1H, dd, $J = 7.6, 6.5$ Hz), 2.93 (1H, m), 2.61 (1H, m), 2.55 (2H, t, $J = 7.9$ Hz), 2.29 (1H, m), 2.24-2.14 (2H, m), 2.12 (1H, ddd, $J = 14.2, 3.2, 3.2$ Hz), 2.02 (2H, m), 2.07-1.98 (2H, m), 1.90-1.80 (3H, m), 1.30 (3H, d, $J = 7.0$ Hz), 1.18 (1H, m), 0.99 (3H, t, $J = 7.0$ Hz), 0.89 (3H, d, $J = 7.6$ Hz), 0.74 (3H, s), 0.69 (3H, s) ppm, ^{13}C NMR (125 MHz, C_6D_6) δ : 173.7, 159.8, 159.3, 131.1, 129.5, 114.1, 107.9, 100.0, 81.0, 80.1, 79.5, 75.7, 74.5, 72.2, 70.12, 70.11, 64.8, 64.4, 63.5, 60.2, 54.8, 41.5, 38.9, 38.8, 35.2, 31.3, 30.5, 29.4, 28.9, 22.9, 22.8, 22.7, 17.8, 14.3 ppm. IR (film): 3470, 2956, 2871, 1732, 1514, 1248, 1081, 1035 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{36}\text{H}_{56}\text{O}_{11}\text{Na}$, 687.3715; found, 687.3715.

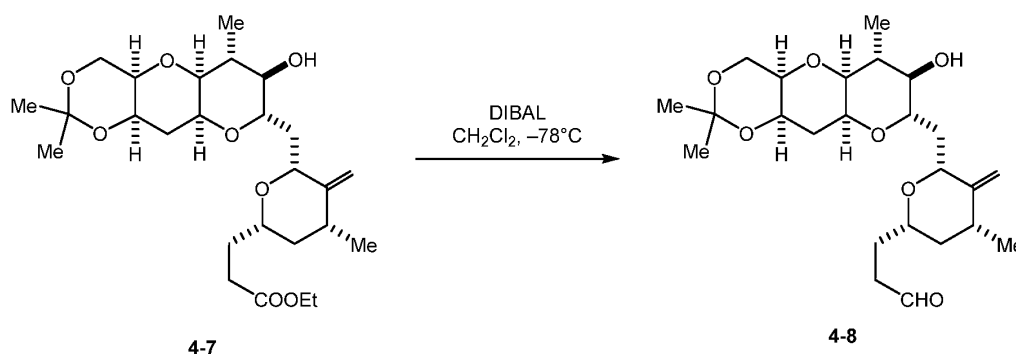


[00738] To a stirred solution of **4-S6** (4.45 g, 6.69 mmol, 1 eq.) and TESH (10.7 mL, 66.9 mmol, 10 eq.) in CH_2Cl_2 (90 mL) was added TESOTf (7.58 mL, 33.5 mmol, 5 eq.) dropwise at 0°C . After being stirred for 3 h, the mixture was poured into sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was washed with brine, dried over Na_2SO_4 , filtrated, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (10%, 50% then 100% EtOAc in Hexanes) to give **4-S7** (2.57 g, 5.81 mmol, 87%) as colorless oil **4-S7**: $[\alpha]_D^{20} -29.1$ (c 1.0, MeOH). ^1H NMR (600 MHz, C_6D_6) δ : 4.77 (1H, s), 4.75 (1H, d, $J = 1.8$ Hz), 4.39 (1H, m), 4.27-4.14 (2H, m), 4.03 (1H, m), 3.95 (1H, m), 3.88 (1H, ddd, $J = 11.9, 7.8, 4.4$ Hz), 3.78 (1H, dd, $J = 8.2, 4.1$ Hz), 3.57 (1H, m), 3.52 (1H, m), 3.46 (1H, ddd, $J = 7.5, 3.6, 3.6$ Hz), 3.41 (1H, m), 3.03 (1H, dd, $J = 5.0$ Hz), 2.91 (1H, brs), 2.88 (1H, brs), 2.44 (2H, m), 2.25 (1H, m), 2.07-1.85 (4H, m), 1.76-1.65 (2H, m), 1.31 (1H, ddd, $J = 12.8, 4.3, 1.8$ Hz), 1.16 (1H, ddd, $J = 15.0, 3.1, 3.1$ Hz), 1.02 (3H, t, $J = 7.0$ Hz), 0.95-0.87 (4H, m), 0.85 (3H, d, $J = 7.6$ Hz) (one OH proton is missing) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 173.5, 151.2, 104.6, 80.9, 78.6, 77.2, 76.7, 76.5, 71.0, 65.5, 63.5, 62.6, 60.3, 42.7, 38.7, 35.9, 35.0, 32.4, 31.3, 30.7, 18.0, 16.2, 14.3 ppm. IR (film): 3435, 2928, 1729, 1420, 1253, 1165,

1092, 1048, 639 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{23}\text{H}_{38}\text{O}_8\text{Na}$, 465.2459; found, 465.2460.

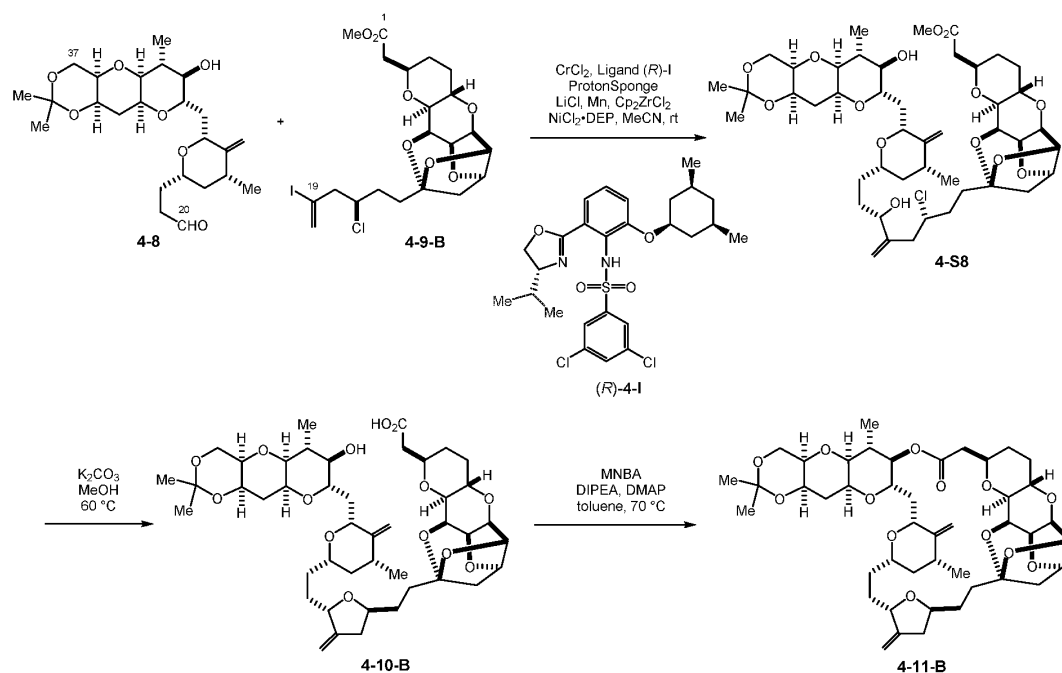


[00739] To a stirred solution of **4-S7** (1.83 g, 4.14 mmol, 1 eq.) and 2,2-dimethoxypropane (37.6 mL, 306 mmol, 30 eq.) in acetone (100 mL) was added CSA (236 mg, 1.02 mmol, 10 mol%) at 0 °C. After being stirred for 30 min, the mixture was warmed to room temperature and stirred for 4 h. The reaction was quenched with Et_3N (2 mL), diluted with EtOAc and stirred for 30 min. Then the mixture was poured into sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtrated, and concentrated under reduced pressure. The residue was passed through a pad of silica gel (EtOAc) to give a crude acetone **4-7** as colorless oil. The obtained crude material was used in the next reaction without further purification. Pure acetone **4-7** (239 mg, 0.495 mmol) was isolated in 91% yield from diol (241 mg, 0.545 mmol). The product was obtained as colorless oil. **4-7**: $[\alpha]_{\text{D}}^{20}$ -19.0 (c 1.05, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 4.91 (1H, s), 4.72 (1H, s), 4.39-4.36 (1H, m), 4.09 (1H, dd, J = 7.2, 4.2 Hz), 4.01-3.98 (2H, m), 3.75 (1H, dd, J = 12.6, 2.4 Hz), 3.68 (1H, dd, J = 7.8, 4.5 Hz), 3.63 (1H, dd, J = 12.3, 2.7 Hz), 3.52 (1H, s), 3.45-3.40 (2H, m), 3.37 (1H, d, J = 7.2 Hz), 3.17 (1H, dd, J = 3.6, 3.6 Hz), 2.71 (1H, dd, J = 5.4, 3.0 Hz), 2.47-2.44 (2H, m), 2.28-2.13 (4H, m), 1.96-1.92 (1H, m), 1.74-1.66 (2H, m), 1.48 (3H, s), 1.41 (1H, ddd, J = 14.4, 4.5, 4.5 Hz), 1.32-1.30 (1H, m), 1.21 (3H, s), 1.17 (3H, d, J = 7.2 Hz), 1.00 (3H, t, J = 6.9 Hz), 0.94-0.87 (1H, m), 0.87 (3H, d, J = 6.6 Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 173.2, 151.3, 104.7, 98.4, 78.3, 76.7, 75.8, 73.9, 70.2, 64.1, 63.7, 62.9, 60.0, 42.9, 40.6, 36.0, 34.5, 32.8, 31.2, 30.6, 28.8, 19.7, 18.0, 16.5, 14.3 ppm. FTIR (film): 3502, 2957, 2926, 2874, 1733, 1457, 1373, 1250, 1181, 1084, 1057, 1039, 977, 906, 836 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{NH}_4]^+$ calcd for $\text{C}_{26}\text{H}_{46}\text{NO}_8$, 500.3218; found, 500.3248.



[00740] To a stirred solution of **4-7** (calculated as 10.2 mmol) in CH_2Cl_2 (200 mL) was added DIBAL (22 mL, 22.4 mmol, 2.2 eq.) dropwise at -78°C . After being stirred for 1.5 h at the same temperature, the reaction was quenched with acetone (0.30 mL) and stirred for 15 min at -78°C . Then sat. Rochelle's salt aq. was added, and the mixture was warmed to room temperature. After being stirred for 3h, the mixture was diluted with EtOAc. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtrated, and concentrated under reduced pressure. The residue was combined with another crude material (ca. 400 mg) and purified by flash column chromatography on neutral silica gel (5%, 20% then 33% EtOAc in Hexanes) to give aldehyde **4-8** (4.1 g, 82-90% for 2 steps) as a colorless amorphous solid. **One batch reaction:** To a solution of **4-7** (206 mg, 0.427 mmol, 1 eq.) in CH_2Cl_2 (8.5 mL) was added DIBAL (0.94 mL of 1.0 M in Hexanes, 0.939 mmol, 2.2 eq.) dropwise at -78°C . After being stirred for 50 min at the same temperature, the reaction was quenched with acetone (0.30 mL) and MeOH (0.30 mL). Then 20% Rochelle's salt aq. was added, and the mixture was stirred for 2 h at room temperature. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtrated, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (5%, 20% then 33% EtOAc in Hexanes) to give **4-8** (167 mg, 0.381 mmol, 89%) as a colorless amorphous solid. **4-8:** $[\alpha]_D^{20} -17.3$ (c 1.0, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 9.46 (1H, t, $J = 1.5$ Hz), 4.92 (1H, s), 4.73 (1H, d, $J = 1.8$ Hz), 4.41 (1H, dd, $J = 5.9$ Hz), 4.02 (1H, dd, $J = 7.6, 4.7$ Hz), 3.74 (1H, dd, $J = 12.3, 2.9$ Hz), 3.66 (1H, dd, $J = 7.6, 4.1$ Hz), 3.60 (1H, dd, $J = 12.3, 3.5$ Hz), 3.48 (1H, m), 3.39 (1H, m), 3.32 (1H, m), 3.27 (1H, m), 3.12 (1H, dd, $J = 3.5, 3.5$ Hz), 2.64 (1H, m), 2.34-2.11 (6H, m), 1.91 (1H, m), 1.53-1.45 (4H, m), 1.38 (1H, ddd, $J = 14.1, 4.7, 4.7$ Hz), 1.24 (1H, ddd, $J = 12.6, 4.4, 2.3$ Hz), 1.19 (3H, s), 1.11 (3H, d, $J = 7.6$ Hz), 0.89 (3H, d, $J = 6.5$ Hz), 0.86 (1H, dd, $J = 24.6, 12.3$ Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 201.0, 151.2, 104.8, 98.4, 78.2,

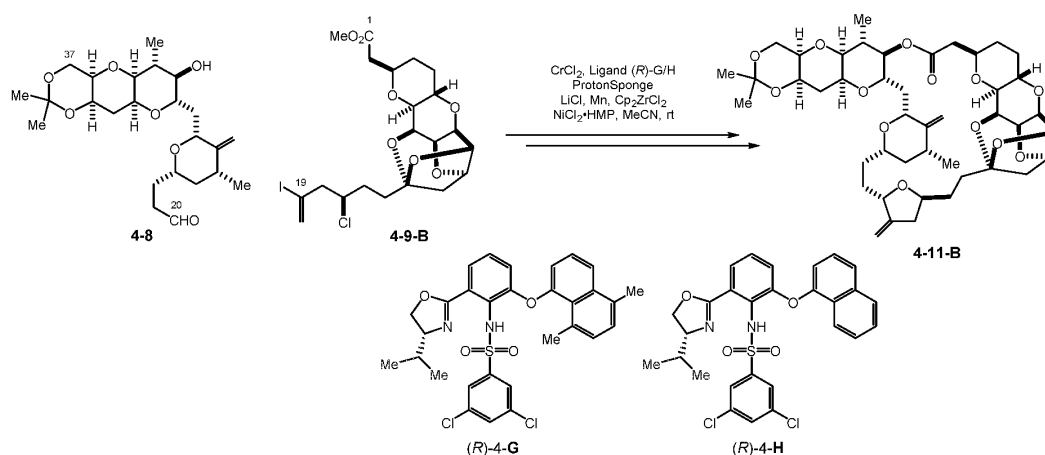
76.6, 75.9, 74.2, 73.8, 70.2, 63.9, 63.2, 63.0, 42.9, 40.38, 40.36, 36.0, 34.5, 33.0, 29.0, 28.4, 19.5, 18.0, 16.4 ppm. IR (film): 3506, 2923, 2852, 1723, 1374, 1112, 1086, 908 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{24}\text{H}_{38}\text{O}_7\text{Na}$, 461.2510; found, 461.2512.



[00741] In a glove box, to a mixture of CrCl_2 (46.2 mg, 0.376 mmol, 10 mol%), (R)-sulfonamide ligand (264 mg, 0.491 mmol, 13 mol%), and proton scavenger (97 mg, 0.453 mmol, 12 mol%) was added MeCN (9.4 mL) and the resulting solution was stirred for 1 h at room temperature. In a separate flask, 4-8 (1.65 g, 3.76 mmol, 1 eq.), 4-9-B (2.30 g, 4.14 mmol, 1.1 eq.), DTBMP (1.93 g, 9.40 mmol, 2.5 eq.), LiCl (319 mg, 7.52 mmol, 2 eq.), Mn (826 mg, 15.0 mmol, 4 eq.), Cp_2ZrCl_2 (2.75 g, 9.40 mmol, 2.5 eq.), and $\text{NiCl}_2\cdot\text{DEP}$ (27.5 mg, 0.0752 mmol, 2 mol%) were mixed and then the solution of Cr complex was transferred to this flask. After being stirred for 1 h at room temperature, the reaction was removed from the glove box and diluted with EtOAc (15 mL). Potassium serinate aq. (0.5M, 15 mL) and sat. NaHCO_3 aq. (15 mL) were added. After being stirred for 1 h, the resulting suspension was filtered through a pad of Celite. The organic layer was separated and the aqueous layer was extracted with EtOAc and washed with brine. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude diol, which was used in the next reaction without further purification. To a stirred solution of the crude diol (calculated as 3.76 mmol, 1 eq.) in MeOH (75 mL) was added K_2CO_3 (5.2 g, 37.6 mmol, 10

eq.). The reaction was heated to 60 °C and stirred for 15 h. Then H₂O (7.5 mL) was added. After being stirred for additional 3 h at the same temperature, the mixture was cooled to room temperature, filtered through a pad of Celite, and concentrated under reduced pressure. To the residue were added EtOAc, sat. NH₄Cl aq., and brine. The organic layer was separated and the aqueous phase was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtrated, and concentrated under reduced pressure to give a crude seco acid **10-B**, which was used in the next reaction without further purification. The macrolactonization was tested under Shiina's condition and Yamaguchi's condition. **Shiina macrolactonization**: To a stirred solution of MNBA (7.8 g, 22.6 mmol, 6 eq.) in toluene (2.5 L) was added DMAP (5.5 g, 45.1 mmol, 12 eq.) at 70 °C. A solution of the crude **4-10-B** (calculated as 3.76 mmol, 1 eq.) and DIPEA (3.9 mL, 22.6 mmol, 6 eq.) in toluene (200 mL) was added to the MNBA solution via a syringe pump over 15 h. After completion of addition, the syringe was rinsed with toluene (40 mL). After being stirred for additional 30 min, the reaction was cooled to room temperature and concentrated under reduced pressure. The residue was dissolved in CH₂Cl₂ and washed with 0.5 M HCl and sat. NaHCO₃ aq., successively. The organic layer was dried over Na₂SO₄, filtrated, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (13%, 25%, 33%, 50%, 67%, then 100% EtOAc in Hexanes) to give **4-11-B** (2.19 g, 2.74 mmol, 73% for 3 steps) as a colorless solid. The obtained product was a mixture of diastereomer at C-20 stereocenter (dr = 19:1, determined by ¹H NMR) **Yamaguchi macroloctonization**: To a stirred solution of the crude **4-10-B** (calculated as 0.456 mmol) in THF (4.6 mL) were added NEt₃ (0.159 mL, 1.14 mmol, 2.5 eq.) and 2,4,6-trichlorobenzoyl chloride (139 mg, 0.57 mmol, 1.25 eq.). The mixture was stirred for 2 h, and then filtered through a pad of Celite, which was washed with toluene (35.4 mL). In a separate flask, DMAP (334 mg, 2.74 mmol, 6 eq.) was dissolved in toluene (290 mL, 1.6 mM) at 80 °C. The solution of mixed anhydride was transferred into DMAP solution via a syringe pump over 13 h. The reaction was cooled to room temperature and quenched with sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated. The residue was purified by flash column chromatography on neutral silica gel (13%, 25%, 33%, 50%, 67%, then 100% EtOAc in Hexanes) to give **4-11-B** (124 mg, 0.155 mmol, 34% for 3 steps) as a colorless solid. **4-11-B**: [α]_D²⁰ -72.5 (c 1.09, CHCl₃). MP: 188-189 °C (recrystallized from CH₂Cl₂-EtOAc). ¹H NMR (600 MHz, C₆D₆) δ : 5.22 (1H, brs), 5.08 (1H, d, *J* = 1.8 Hz), 5.00 (1H, ddd, *J* = 10.6, 7.6, 2.9 Hz), 4.94 (1H, s), 4.82 (1H, dd, *J* =

7.6, 5.3 Hz), 4.76 (1H, brs), 4.68 (1H, brd, $J = 10.0$ Hz), 4.55 (1H, ddd, $J = 10.3, 10.3, 4.1$ Hz), 4.30 (1H, dd, $J = 4.1, 1.8$ Hz), 4.17-4.07 (3H, m), 3.98 (1H, m), 3.90 (1H, dd, $J = 6.5, 4.7$ Hz), 3.85 (1H, brd, $J = 10.6$ Hz), 3.81 (1H, dd, $J = 11.7, 4.1$ Hz), 3.73 (1H, m), 3.69 (1H, dd, $J = 12.0, 3.8$ Hz), 3.65 (1H, dd, $J = 6.5, 4.1$ Hz), 3.58-3.50 (2H, m), 3.05 (1H, dd, $J = 3.3, 3.3$ Hz), 2.85 (1H, dd, $J = 16.4, 7.0$ Hz), 2.76 (1H, m), 2.72 (1H, m), 2.62 (1H, dd, $J = 9.4, 1.8$ Hz), 2.50-2.36 (3H, m), 2.34-2.22 (3H, m), 2.22-2.03 (5H, m), 1.97-1.85 (3H, m), 1.72 (1H, dddd, $J = 9.8, 9.8, 9.8, 4.7$ Hz), 1.62-1.27 (11H, m), 1.26-1.21 (6H, m), 1.06 (1H, dd, $J = 23.5, 12.3$ Hz), 0.97 (3H, d, $J = 6.5$ Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 171.3, 152.9, 152.7, 110.0, 105.0, 103.8, 98.7, 82.3, 81.0, 78.2, 78.1, 77.3, 76.9, 76.4, 76.1, 75.1, 74.7, 74.3 (x2), 74.0, 70.4, 69.9, 68.5, 64.1, 63.9, 62.4, 48.5, 43.8, 41.5, 39.4, 39.3, 38.8, 36.3, 35.6, 32.5, 32.4, 31.11, 31.09, 30.62, 29.3, 28.5, 20.3, 18.1, 16.4 ppm. IR (film): 2923, 2869, 1725, 1188, 1118, 1086, 755 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{44}\text{H}_{62}\text{O}_{13}\text{Na}$, 821.4083; found, 821.4084.

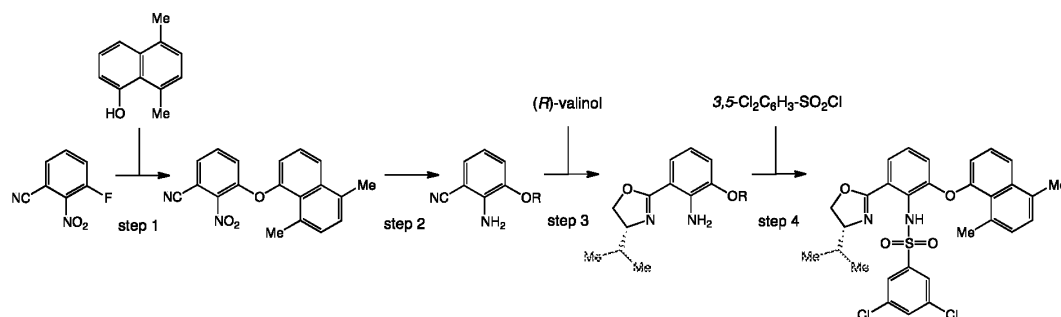


[00742] Ni/Cr-Mediated coupling of **4-8** with **4-9-B** in the presence of Cr-catalyst prepared from (*R*)-**4-G** and (*R*)-**4-H** was carried out as follows: In a glove box, to a mixture of CrCl_2 (7.0 mg, 10 mol%), sulfonamide (*R*)-**G** (43.2 mg, 13 mol%), proton scavenger (14.7 mg, 12 mol%) and LiCl (24.1 mg, 0.57 mmol, 1.0 eq) was added MeCN (1.4 mL, 0.4 M) and stirred for 1 h at r.t. In a separate flask, **4-8** (250 mg, 0.57 mmol, 1.0 eq), **4-9-B** (350 mg, 0.63 mmol, 1.1 eq), DTBMP (290 mg, 1.43 mmol, 2.5 eq), Mn (125 mg, 2.28 mmol, 4.0 eq) and Cp_2ZrCl_2 (0.42 g, 1.43 mmol, 2.5 eq) were mixed, and the solution of Cr complex was then transferred to this flask. After being stirred for 1 min, $\text{HMP}\cdot\text{NiCl}_2$ (2.2 mg, 1 mol%; doped in LiCl) was added. Additional $\text{HMP}\cdot\text{NiCl}_2$ (2.2 mg, 1 mol%; doped in LiCl) was added after 1 and 2 h, respectively. The reaction mixture was stirred for 3 h total at r.t. The reaction was removed from the glove box and diluted with EtOAc (10 mL). Aqueous potassium serinate

(0.5 M, 3 mL) and saturated aqueous NaHCO_3 (3 mL) were added. After being stirred for 1 h, the resulting suspension was filtered through a pad of celite. The filtrate was extracted with EtOAc, washed with sat. NaCl, dried over Na_2SO_4 , filtered, and concentrated. With use of the procedure given for the coupling with sulfonamide (*R*)-**4-G**, the crude product was converted to macrolactone **4-11-B**; thus, 296 mg **4-11-B** (65% overall yield; *dr* = 29:1) was obtained from 250 mg **4-8**.

[00743] The Ni/Cr-mediated coupling with the Cr-catalyst prepared from (*R*)-**4-H** was carried out with use of the same procedure; 305 mg **4-11-B** (67% overall yield; *dr* = 24:1) was obtained from 250 mg **4-8**.

[00744] Sulfonamides (*R*)-**4-G**, (*R*)-**4-H**, and (*R*)-**4-I** were synthesized via the general scheme shown below. As an example, the synthesis of (*R*)-**4-G** is given below.



[00745] **Step 1:** To a slurry of NaH (5.56 g, 60% in mineral oil, 140 mmol, 3.0 eq) in anhydrous THF (100 mL) was added 5,8-dimethylnaphthol (8.00 g, 46.5 mmol, 1.0 eq) at 0 °C with stirring for 1 h. Then 3-fluoro-2-nitrobenzonitrile (8.50 g, 51.2 mmol, 1.1 eq) was dissolved in THF (50 mL) added via syringe. The reaction was allowed to warm to room temperature and stirred for 3 h at room temperature before quenched with water at 0 °C. The organic solvent (THF) was removed in *vacuo* and the slurry was extracted with EtOAc (2 X 100 mL). The combined organic layers were washed with brine, water, dried over Na_2SO_4 and filtered. The solvent was removed in *vacuo* to afford 16.1 g of crude product which was purified with column chromatography using EtOAc: Hexanes (10:90) to afford pure red solid (12.20 g, 38.4 mmol, 82%). ^1H NMR (600 MHz, CDCl_3) δ : 7.96 (1H, dd, *J* = 8.5, 1.1 Hz), 7.48 (1H, dd, *J* = 8.4, 7.5 Hz), 7.46–7.40 (2H, m), 7.30–7.24 (1H, m), 7.19 (1H, d, *J* = 7.1 Hz), 7.08 (1H, dd, *J* = 7.5, 1.1 Hz), 6.97–6.91 (1H, m), 2.69 (3H, s), 2.66 (3H, s) ppm. ^{13}C NMR (125 MHz, CDCl_3) δ : 151.6, 151.5, 142.3, 135.6, 132.7, 130.9, 129.7, 127.7, 126.7, 126.5, 125.3, 123.5, 122.8, 117.6, 113.6, 108.0, 23.6, 20.1 ppm. IR (neat) ν 2963, 2929, 2242, 1599, 1573, 1442, 1357, 1263, 1135, 997, 908, 824, 793, 750, 729 cm^{-1} . HRMS (ESI) calcd. for $\text{C}_{19}\text{H}_{14}\text{N}_2\text{O}_3\text{Na}$ [$\text{M} + \text{Na}$] $^+$: 341.0897, found 341.0903.

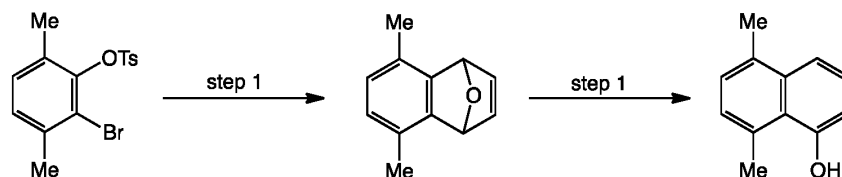
[00746] Step 2: To a solution of the above adduct (12.20 g, 38.4 mmol, 1.0 eq) in anhydrous EtOAc (150 mL) was added AcOH (17.5 mL, 307 mmol, 8.0 eq) followed by Pd/C (0.61 g, 5.8 mmol, 0.015 eq). A hydrogen balloon was attached. After the reaction was stirred for 4 h, the slurry was filtered through celite, and a mixture of brine and sat. NaHCO₃ (1:1) was added to filtrate, extracted carefully with EtOAc (2 X 50 mL) and the organic solvent was removed in *vacuo* and crude product was purified by column chromatography using EtOAc: Hexanes (5:95) to afford pure yellow solid (9.69 g, 33.6 mmol, 88% yield). ¹H NMR (600 MHz, CDCl₃) δ: 7.85 (1H, dd, *J* = 8.5, 1.2 Hz), 7.43 (1H, dd, *J* = 8.5, 7.5 Hz), 7.28–7.23 (1H, m), 7.18 (1H, dd, *J* = 7.1, 1.0 Hz), 7.13 (1H, dd, *J* = 7.9, 1.4 Hz), 6.96 (1H, dd, *J* = 7.6, 1.1 Hz), 6.67 (1H, dd, *J* = 8.0, 1.4 Hz), 6.59 (1H, t, *J* = 7.9 Hz), 4.74 (1H, s), 2.75 (3H, s), 2.68 (3H, s). ¹³C NMR (125 MHz, CDCl₃) δ: 153.3, 145.5, 141.7, 135.6, 132.5, 131.7, 129.1, 127.4, 126.6, 126.0, 125.4, 121.7, 120.9, 117.8, 117.3, 115.9, 96.7, 23.9, 20.1. IR (neat) ν 3477, 3366, 2963, 2930, 2218, 1621, 1481, 1232, 1140, 969, 824, 750, 731 cm⁻¹.

[00747] Step 3: To a solution of the above product (9.60 g, 33.3 mmol, 1.0 eq) in anhydrous chlorobenzene (80 mL) was added ZnCl₂ (9.09 g, 66.7 mmol, 2.0 eq) and (*R*)-valinol (6.9 g, 66.7, 2.0 eq) at room temperature. The solution was heated to reflux for 30 h and quenched with water. The slurry was treated with NH₄OH (50 mL) with stirring for 30 min and extracted with EtOAc (3 X 50 mL). The combined organic layers were washed with brine and water, dried over Na₂SO₄ and filtered. The solvent was removed in *vacuo* and the crude product was purified by column chromatography using EtOAc: Hexanes (15:85) to afford pure light yellow solid (10.43 g, 27.9 mmol, 84%). [α]_D²⁰ -22.9 (c = 1.0, CHCl₃). ¹H NMR (600 MHz, CDCl₃) δ: 7.78 (1H, dd, *J* = 8.5, 1.1 Hz), 7.51 (1H, dd, *J* = 8.1, 1.5 Hz), 7.39 (1H, t, *J* = 8.0 Hz), 7.25 (1H, dd, *J* = 7.1, 0.9 Hz), 7.18 (1H, dd, *J* = 7.2, 1.1 Hz), 6.97 – 6.92 (1H, m), 6.72 (1H, dd, *J* = 7.8, 1.4 Hz), 6.56 (2H, t, *J* = 7.9 Hz), 6.51 (1H, s), 4.38 (1H, dd, *J* = 9.5, 8.2 Hz), 4.15 (1H, ddd, *J* = 9.5, 8.1, 6.8 Hz), 4.05 (1H, t, *J* = 8.1 Hz), 2.87 (3H, s), 2.68 (3H, s), 1.89–1.78 (1H, m, *J* = 6.6 Hz), 1.08 (3H, d, *J* = 6.7 Hz), 0.98 (3H, d, *J* = 6.8 Hz) ppm. ¹³C NMR (125 MHz, CDCl₃) δ: 163.6, 154.8, 144.8, 140.9, 135.5, 132.4, 132.2, 128.7, 127.2, 126.5, 125.5, 124.2, 120.5, 120.1, 115.1, 114.4, 110.2, 73.1, 69.0, 33.3, 24.5, 20.2, 19.1, 18.7 ppm. IR (neat) ν 3478, 3282, 2958, 1634, 1592, 1551, 1472, 1362, 1228, 1192, 1075, 981, 823, 734 cm⁻¹. HRMS (ESI) calcd. for C₂₄H₂₆Cl₂N₂O₂ [M+H]⁺: 375.2067, found 375.2051.

[00748] Step 4: To a solution of the above product (10.41 g, 27.8 mmol, 1.0 eq) in anhydrous pyridine (70 mL) was added 3,5-dichlorobenzenesulfonyl chloride (10.26 g, 41.8 mmol, 1.5 equiv) and DMAP (36 mg, 0.3 mmol, 0.01 eq). The solution was stirred at room temperature overnight before quenched with water. The mixture was extracted with EtOAc (2 X 50 mL)

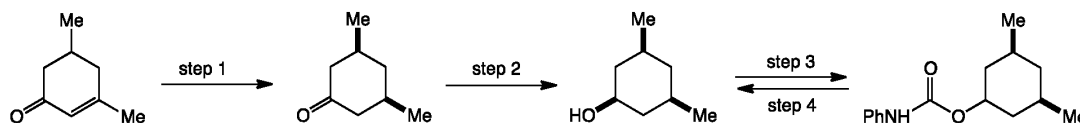
and combined organic layers were washed with 1N HCl (3 X 50 ml), brine and water. The organic layer was dried over Na₂SO₄ and filtered. The solvent was removed in *vacuo* to afford the crude product which was purified by column chromatography using EtOAc: Hexanes (15:85) to afford pure white solid (*R*)-**4-G** (12.82g, 22.0 mmol, 83%). Recrystallization from EtOAc/Hexanes gave 10.31 g white crystals. (*R*)-**4-G**: [α]_D²⁰ -12.1 (c = 1.0, CHCl₃). mp = 138-140 °C. ¹H NMR (600MHz, CDCl₃) δ : 12.71 (1H, s), 7.78 (2H, s), 7.56 (4H, brs), 7.45-7.42 (2H, m), 7.19-7.14 (3H, m), 7.02 (4H, s), 7.01-6.97 (4H, m), 6.90 (2H, brs), 6.85 (2H, s), 6.80 (2H, s), 4.48 (1H, dd, *J* = 9.6, 8.3 Hz), 4.25 (1H, brs), 4.17 (2H, t, *J* = 8.2 Hz), 2.64 (1.3H, s), 2.37 (1.7H, s), 2.18 (3H, s), 1.92 (1H, m), 1.15 (3H, s), 1.04 (3H, s) ppm. ¹³C NMR (125 MHz, CDCl₃) δ : 163.4, 152.1, 150.7, 145.0, 135.4, 134.7, 132.2, 131.3, 130.9, 128.7, 127.0, 126.5, 125.2, 124.8, 124.6, 123.6, 123.1, 122.3, 121.7, 121.0, 118.3, 116.9, 115.9, 72.7, 70.1, 33.3, 23.2, 20.1, 18.8 ppm. IR (neat) ν 3078, 2959, 2929, 1639, 1571, 1464, 1339, 1265, 1164, 1134, 989, 923, 801, 746, 669, 576 cm⁻¹. HRMS (ESI) calcd. for C₃₀H₂₈Cl₂N₂O₄S [M+H]⁺: 583.1220, found 583.1233;

[00749] Using the same procedure, sulfonamides (*R*)-**4-H** and (*R*)-**4-I** were synthesized. (*R*)-**4-H**: [α]_D²⁰ -9.1 (c = 1.0, CHCl₃). mp = 117-119 °C. ¹H NMR (500MHz, CDCl₃) δ : 12.77 (1H, s), 7.85 - 7.78 (1H, m), 7.67 - 7.59 (2H, m), 7.59 - 7.55 (2H, m), 7.47 (1H, ddd, *J* = 8.2, 6.9, 1.3 Hz), 7.40 - 7.30 (2H, m), 7.02 (1H, t, *J* = 8.0 Hz), 6.89 - 6.82 (3H, m), 4.49 (1H, dd, *J* = 9.6, 8.3 Hz), 4.25 (1H, ddd, *J* = 9.6, 8.1, 6.8 Hz), 4.17 (1H, t, *J* = 8.3 Hz), 1.97 - 1.86 (1H, m, *J* = 6.8 Hz), 1.15 (3H, d, *J* = 6.7 Hz), 1.04 (3H, d, *J* = 6.7 Hz) ppm. ¹³C NMR (125 MHz, CDCl₃) δ 163.5, 150.9, 150.2, 145.3, 134.9, 131.1, 127.7, 126.8, 126.6, 126.1, 125.7, 124.8, 124.7, 124.6, 124.3, 123.9, 121.8, 121.5, 118.4, 114.1, 72.8, 70.3, 33.4, 18.9 ppm. IR (neat) ν 3076, 2959, 2929, 1639, 1571, 1465, 1389, 1339, 1268, 1165, 1134, 1078, 980, 800, 739, 580 cm⁻¹. HRMS (ESI) calcd. for C₂₈H₂₅Cl₂N₂O₄S [M+H]⁺: 555.0907, found 555.0915. (*R*)-**4-I**: [α]_D²⁰ +13.4 (c = 1.0, CHCl₃). mp = 96-98 °C. ¹H NMR (600 MHz, CDCl₃) δ : 12.27 (1H, brs), 7.86-7.94 (2H, m), 7.50-7.54 (1H, m), 7.43 (1H, d, *J* = 7.8, Hz), 7.13 (1H, t, *J* = 8.1 Hz), 7.01 (1H, d, *J* = 8.3 Hz), 4.40-4.48 (1H, m), 4.06-4.20 (3H, m), 1.83-1.91 (1H, m), 1.79 (1H, brd, *J* = 12.2 Hz), 1.71 (1H, br d, *J* = 12.2 Hz), 1.55 (1H, m), 1.31-1.43 (2H, m), 1.10 (3H, d, *J* = 6.6 Hz), 0.99 (3H, d, *J* = 6.6 Hz), 0.85 (3H, d, *J* = 6.8 Hz), 0.87 (3H, d, *J* = 6.8 Hz), 0.40 (1H, q, *J* = 12.0 Hz), 0.25 (1H, q, *J* = 12.0 Hz), 0.20 (1H, q, *J* = 12.0 Hz); ¹³C NMR (125 MHz, CDCl₃) δ : 163.4, 150.5, 146.3, 135.3, 131.5, 128.9, 125.2, 124.8, 120.8, 119.1, 117.2, 76.7, 72.6, 69.8, 42.9, 39.1, 39.0, 33.1, 30.4, 30.4, 22.0, 18.7, 18.6. IR (neat) ν 3079, 2951, 2926, 2869, 1638, 1571, 1467, 1337, 1269, 1165, 1014, 940, 801, 744, 606, 577 cm⁻¹. HRMS (ESI) *m/z* 539.1556 [(M+H)⁺]; calcd. for C₂₆H₃₃Cl₂N₂O₄S: 539.1533].

5,8-Dimethyl-1-naphthol

[00750] Step 1: A stirred solution of bromotosylate (see, *e.g.*, Velder, J.; Robert, T.; Weidner, I.; Neudörfl, J. M. *Adv. Synth. Catal.* **2008**, 350, 1309) (20.0 g, 56.5 mmol, 1.0 equiv) and freshly distilled furan (34 mL, 452 mmol, 8.0 equiv) in 150 mL of THF was cooled under nitrogen to -78°C , and *n*-BuLi (34 mL, 84.7 mmol, 2.5 M, 1.5 equiv) was added dropwise. Stirring was continued for 10 h, during which time the solution was allowed to warm to room temperature. See, *e.g.*, Jung, K. Y.; Koreeda, M. *J. Org. Chem.* **1989**, 54, 5667. The reaction was quenched by the addition of a few drops of saturated aqueous ammonium chloride solution. The solvent was evaporated, and the resulting brownish residue was taken up in 200 mL of diethyl ether. The ethereal solution was washed with brine (100 mL) and water (100 mL), and dried over anhydrous Na_2SO_4 . Evaporation of the solvent under reduced pressure, followed by purification by flash column chromatography on silica gel with ethyl acetate/hexanes (1:9) as the eluent, gave the adduct as light yellow solid (8.64 g, 50.2 mmol, 88%). ^1H NMR (600 MHz, CDCl_3) δ : 7.03 (2H, dd, $J = 1.0, 1.0$ Hz), 6.72 (2H, s), 5.8 (2H, dd, $J = 1.0, 1.0$ Hz), 2.3 (6H, s) ppm.

[00751] Step 2: To a solution of $\text{Cu}(\text{OTf})_2$ (0.358 g, 0.99 mmol) in anhydrous DCE (10 mL) at 4°C , the above adduct (3.41 g, 19.8 mmol) in DCE (10 mL) was added under a nitrogen atmosphere. The resulting mixture was stirred at r.t. until the reaction was complete (TLC monitoring). The mixture was then quenched by the addition of H_2O (20 mL), extracted with CH_2Cl_2 (50 mL) and dried over Na_2SO_4 . The crude product thus obtained was purified by flash column chromatography on silica gel with methylene chloride/hexanes (2:3) as the eluent, to give the 5,8-dimethyl-1-naphthol as white crystalline solid (2.67g, 15.5 mmol, 77%). mp = $76\text{--}78^{\circ}\text{C}$; ^1H NMR (600 MHz, CDCl_3) δ : 7.55 (1H, dd, $J = 8.5, 1.1$ Hz), 7.33–7.23 (1H, m), 7.17 (1H, dd, $J = 7.1, 1.0$ Hz), 7.09 (1H, dd, $J = 7.1, 1.0$ Hz), 6.75 (1H, dd, $J = 7.4, 1.1$ Hz), 5.19 (1H, s), 2.92 (3H, s), 2.61 (3H, s) ppm. ^{13}C NMR (125 MHz, CDCl_3) δ : 154.3, 135.5, 132.9, 131.8, 127.7, 126.9, 125.4, 123.8, 117.6, 110.2, 24.8, 20.2 ppm. IR (neat) ν 3322, 3032, 2930, 2898, 1589, 1461, 1413, 1277, 1240, 1138, 897, 737 cm^{-1} .

syn,syn,syn-3,5-Dimethylcyclohexan-1-ol

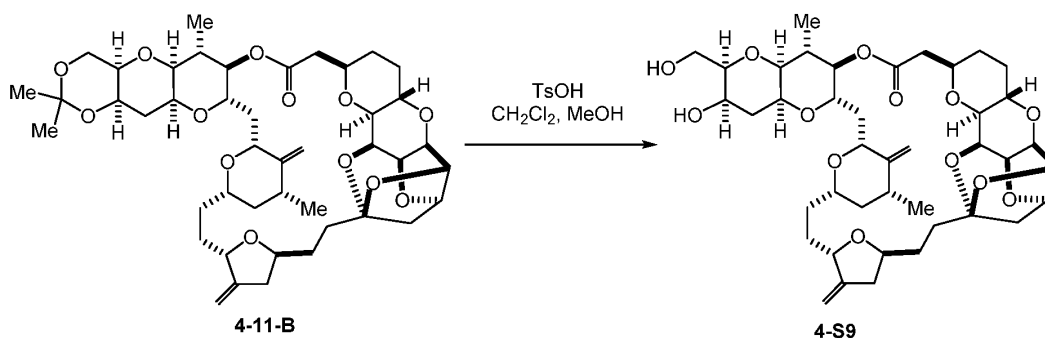
[00752] Step 1: Pd/C (wet Degussa, 5% wt%; 1.2 g, 8.46 mmol, 0.05 eq) was added to the solution of 3,5-dimethylcyclohexanone (21.0 g, 169 mmol, 1.0 eq) in isopropanol (210 mL). The internal atmosphere was replaced with H₂ (balloon). The reaction mixture was stirred under H₂ (balloon) at room temperature for 2 h. The crude reaction mixture was filtered through Celite pad, and the filtrate was diluted with 100 mL water. The ketone was extracted with pentane (3 x 150 mL). The combined organic layers were washed with water (2 x 100 mL), dried over Na₂SO₄, and concentrated under reduced pressure, to give 21.3 g of crude product (*syn:anti* ratio = 33:1 by ¹H NMR analysis). The crude product **2** was used for next step without further purification.

[00753] Step 2: To a solution of 3,5-dimethylcyclohexanone (~21.0 g, 0.17 mol, 1.0 eq) in anhydrous hexanes (315 mL) was added anhydrous isopropanol (129 mL, 1.70 mol, 10 eq), followed by sodium metal (24.4 g, 102 mmol, 6.00 eq) were added under N₂ atmosphere at 10 °C. The resulting solution was stirred vigorously at this temperature, slowly warmed to room temperature, and stirred for another 2 h. The reaction was quenched by the addition of an aqueous HCl (100 mL, 3.0 M) at 0 °C, diluted with Et₂O (100 mL), and the organic layer was separated. The aqueous layer was extracted by Et₂O (3 x 200 mL). Combined organic layers were dried over Na₂SO₄ and concentrated under reduced pressure to give 19.1 g of crude product (the diastereomeric ratio was >45:1 by ¹H NMR). The product thus obtained was purified by recrystallization of its phenylurethane.

[00754] Step 3: To the solution of crude alcohol (19.1 g, 149 mmol, 1 equiv) in CH₂Cl₂ (200 mL) was added phenyl isocyanate (19.4 mL, 179 mmol, 1.2 equiv) followed by DMAP (0.98 g, 7.46 mmol, 0.05 eq). The resulting mixture was stirred at room temperature for 15 h. After completion of the reaction (by TLC), the reaction mixture was filtered through silica gel pad and concentrated under reduced pressure. The crude residue was dissolved in EtOAc and diluted with large excess of hexanes. The solution was left standing for overnight at 0 °C and white crystals were precipitated out. Crystals were collected by filtration and washed with cold hexanes to give urethane (31.22 g; 75 % overall yield from 3,5-dimethylcyclohexanone; *dr* = ca. 200:1). m.p. 95-97 °C (ref. 106-107 °C)⁶. ¹H NMR (500MHz, CDCl₃) δ: 7.38-7.34 (2H, m), 7.28 (2H, t, *J* = 15 Hz), 7.05 (1H, t, *J* = 15 Hz), 6.52 (1H, br), 4.7 (1H, m), 2.04 (1H, dd, *J* = 3.4Hz, 3.2Hz), 1.63 (1H, dt, *J*₁ = 14 Hz, *J*₂ = 1.2 Hz), 1.57-1.52 (2H, m), 0.96 (2H,

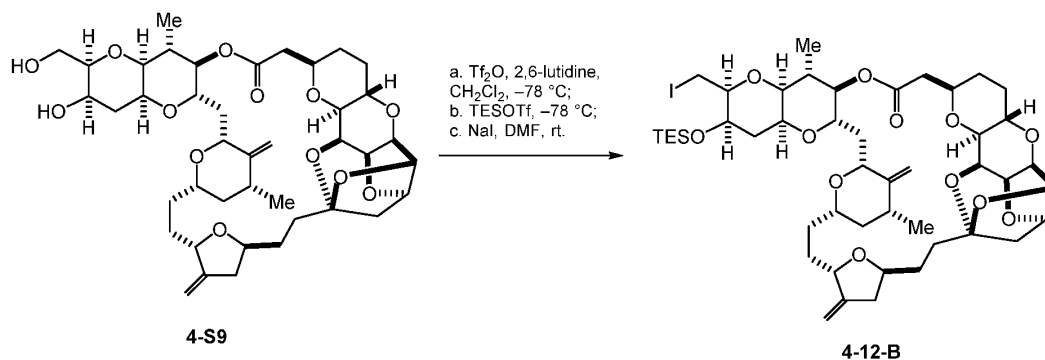
m), 0.94 (6H, d, $J = 6.6$ Hz), 0.56 (1H, q, $J = 12$ Hz). ^{13}C NMR (125MHz, CDCl_3) δ : 153.3, 138.1, 128.9, 123.2, 118.5, 73.9, 42.9, 40.2, 30.6, 22.1. IR (neat) ν 3313, 2949, 2925, 1702, 1598, 1539, 1442, 1312, 1221, 1052, 751, 691 cm^{-1} . HRMS (ESI) m/z : $(\text{M}+\text{Na})^+$ calculated for $\text{C}_{15}\text{H}_{21}\text{NO}_2\text{Na}$, 270.1484; found, 270.1464.

[00755] Step 4: A solution of urethane (31.2 g, 126.2mmol) in 10% NaOH/MeOH (320 mL) was heated up to 60 °C and stirred for 18 h at this temperature. After completion of the reaction, the reaction mixture was diluted by EtOAc/hexanes (1:1) and washed with H_2O , 1N HCl (3 x 100 mL), and brine, dried and concentrated. Distillation of the crude product gave pure *syn,syn,syn*-3,5-dimethylcyclohexan-1-ol (14.42 g; colorless liquid; 90% yield). ^1H NMR (500MHz, CDCl_3) δ : 3.60 (1H, m), 1.92 (2H, m), 1.58 (1H, m), 1.49-1.42 (3H, m), 0.92 (1H, d, $J = 7.8\text{Hz}$), 0.82 (2H, q, $J = 15$ Hz), 0.51 (1H, q, $J = 15$ Hz) ppm. ^{13}C NMR (125MHz, CDCl_3) δ : 70.3, 44.0, 43.1, 30.7, 22.2 ppm.



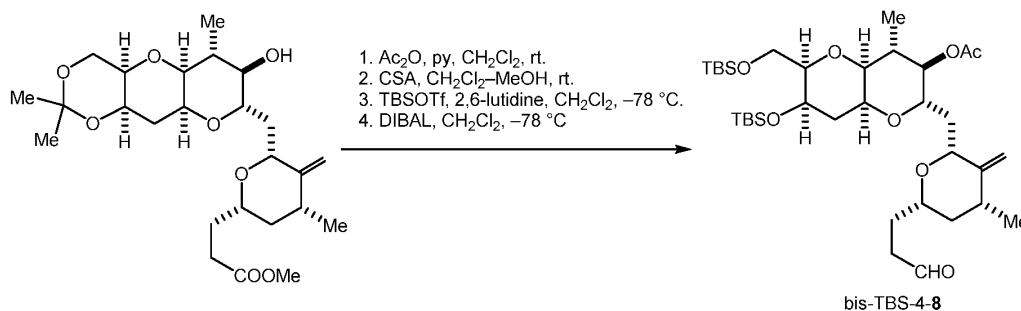
[00756] To a stirred solution of **4-11-B** (1.42 g, 1.78 mmol, 1 eq., dr = ~20:1) in CH_2Cl_2 (8.9 mL) and MeOH (8.9 mL, 0.2 M) was added TsOH· H_2O (16.9 mg, 0.089 mmol, 5 mol%) at room temperature. After being stirred for 1 h, the reaction was quenched with Et_3N (0.1 mL) and concentrated under reduced pressure. The residue was dissolved in EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude diol **4-S9**, which was used in the next reaction without further purification. For the analytical purpose, a single crystal of the **4-S9** was obtained by vapor diffusion method (CH_2Cl_2 - Et_2O) (See Figure 14B). **4-S9**: $[\alpha]_D^{20}$ -69.5 (c 1.02, CHCl_3). MP: 239-241 °C (recrystallized from CH_2Cl_2 - Et_2O). ^1H NMR (600 MHz, CD_2Cl_2) δ : 5.03 (1H, s), 4.95 (1H, s), 4.84 (1H, s), 4.81 (1H, s), 4.66 (1H, dd, $J = 4.5, 4.5$ Hz), 4.56 (1H, dd, $J = 4.5, 4.5$ Hz), 2.54 (1H, dd, $J = 3.9, 3.9$ Hz), 4.38 (1H, d, $J = 10.2$ Hz), 4.33 (1H, d, $J = 10.8$ Hz), 4.29 (1H, s), 4.23 (1H, ddd, $J = 9.9, 9.9, 4.2$ Hz), 4.16 (1H, dd, $J =$

5.7, 5.7 Hz), 4.09 (1H, dd, $J = 7.8, 7.8$ Hz), 4.03 (1H, dd, $J = 6.3, 3.9$ Hz), 3.87 (1H, s), 3.84-3.79 (2H, m), 3.71-3.62 (4H, m), 3.50-3.43 (2H, m), 3.64 (1H, dd, $J = 6.6, 4.2$ Hz), 3.32 (1H, s), 2.89 (1H, dd, $J = 9.3, 1.5$ Hz), 2.81-2.77 (1H, m), 2.51 (1H, dd, $J = 17.1, 9.9$ Hz), 2.37 (1H, dd, $J = 17.1, 2.1$ Hz), 2.35-2.32 (2H, m), 2.26 (1H, d, $J = 15.0$ Hz), 2.23-2.07 (4H, m), 2.05-2.03 (1H, m), 1.96-1.91 (2H, m), 1.88-1.78 (3H, m), 1.72-1.65 (2H, m), 1.56 (1H, ddd, $J = 12.3, 12.3, 4.8$ Hz), 1.51-1.31 (5H, m), 1.17-1.13 (1H, m), 1.15 (3H, d, $J = 7.2$ Hz), 1.10-1.03 (1H, m), 1.10 (3H, d, $J = 6.6$ Hz) ppm. ^{13}C NMR (125 MHz, CD_2Cl_2) δ : 171.2, 153.2, 152.4, 110.3, 104.7, 104.4, 82.9, 81.7, 80.2, 78.1, 77.6, 77.5, 77.1, 75.9, 75.5, 74.7, 74.5, 74.4, 74.3, 73.9, 73.8, 68.9, 66.1, 64.4, 63.9, 48.8, 44.2, 41.0, 39.2, 37.1, 36.4, 35.9, 35.3, 35.2, 31.9, 31.7, 31.3, 30.8, 29.1, 18.3, 16.9 ppm. FTIR (film): 3502, 2926, 2854, 1734, 1189, 1135, 1071, 1020, 995, 753 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{NH}_4]^+$ calcd for $\text{C}_{41}\text{H}_{62}\text{NO}_{13}$, 776.4216; found, 776.4230.



[00757] To a stirred solution of **4-S9**(crude, calculated as 1.78 mmol) in CH_2Cl_2 (17.8 mL) was added 2,6-lutidine (1.04 mL, 8.93 mmol, 5 eq.). The mixture was cooled to -78°C , and then TiF_2O (0.36 mL, 2.14 mmol, 1.2 eq.) was added. After being stirred for 15 min at the same temperature, TESOTf (0.60 mL, 2.65 mmol, 1.5 eq) was added. The reaction was warmed to 0°C with ice bath. After being stirred for additional 20 min, to the reaction were added DMF (35.6 mL) and NaI (1.33 g, 8.87 mmol, 5 eq). The resulting mixture was allowed to warm to room temperature and stirred for 2.5 h. Then the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with TBME. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (9%, 13%, 17%, then 25% EtOAc in Hexanes) to give iodide **4-12-B** (1.64 g, 1.67 mmol, 94% for 2 steps), which was a mixture of C20-epimer (dr = ~25:1), as a colorless solid. The C20-epimer was separated by HPLC purification (Column: DuPont Instruments ZORBAL SIL 21.2 mm x 25 cm (880952-101), Solvent: 10% *i*PrOH in Hexanes, Flow rate: 10.0

mL/min., Detection: UV at 254 nm and 220 nm, $t_{R(\text{desired})}$ = 19 min. $t_{R(\text{C20-epimer})}$ = 25 min.). **4-12-B**: $[\alpha]_D^{20}$ -33.2 (c 1.0, CHCl_3). MP: 158-160 °C (recrystallized from Hexanes-*i*PrOH). ^1H NMR (600 MHz, C_6D_6) δ : 5.17 (1H, s), 5.12 (1H, d, J = 1.8 Hz), 4.92 (1Hm s), 4.78 (3H, m), 4.67 (1H, brd, J = 10.0 Hz), 4.53 (1H, ddd, J = 10.0, 10.0, 4.1 Hz), 4.27 (1H, dd, J = 3.5, 1.8 Hz), 4.14 (1H, dd, J = 4.7, 4.7 Hz), 4.10 (1H, dd, J = 4.7, 4.7 Hz), 4.06 (1H, m), 4.00 (1H, m), 3.89 (1H, dd, J = 6.5, 4.7 Hz), 3.79 (1H, d, J = 11.2 Hz), 3.75-3.67 (2H, m), 3.64 (1H, dd, J = 6.5, 4.1 Hz), 3.36-3.28 (3H, m), 3.20 (1H, ddd, J = 6.7, 6.7, 1.8 Hz), 3.03 (1H, dd, J = 4.7, 3.5 Hz), 2.78 (1H, dd, J = 16.7, 7.3 Hz), 2.71 (1H, m), 2.60 (1H, dd, J = 9.7, 2.1 Hz), 2.41-2.23 (5H, m), 2.23-1.90 (8H, m), 1.84 (1H, m), 1.73 (1H, m), 1.58-1.27 (8H, m), 1.17 (3H, d, J = 7.0 Hz), 1.08 (1H, dd, J = 24.1, 12.3 Hz), 1.02 (9H, t, J = 7.9 Hz), 0.99 (3H, d, J = 6.5 Hz), 0.67 (6H, q, J = 8.2 Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 171.4, 153.0, 152.6, 110.0, 105.0, 103.6, 82.4, 81.0, 80.0, 78.7, 78.0, 77.7, 76.9, 76.1, 75.6, 74.8, 74.7, 74.08, 74.06, 73.9, 70.1, 68.5, 64.9, 64.0, 48.6, 43.9, 41.4, 39.4, 39.3, 39.0, 36.3, 36.2, 35.5, 32.5, 31.1, 30.7, 30.6, 29.0, 18.1 16.6, 73.2, 5.7, 5.4 ppm. IR (film): 2951, 2874, 1726, 1087, 1016, 996, 907, 753 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{47}\text{H}_{71}\text{IO}_{12}\text{SiNa}$, 1005.3652; found, 1005.3649.



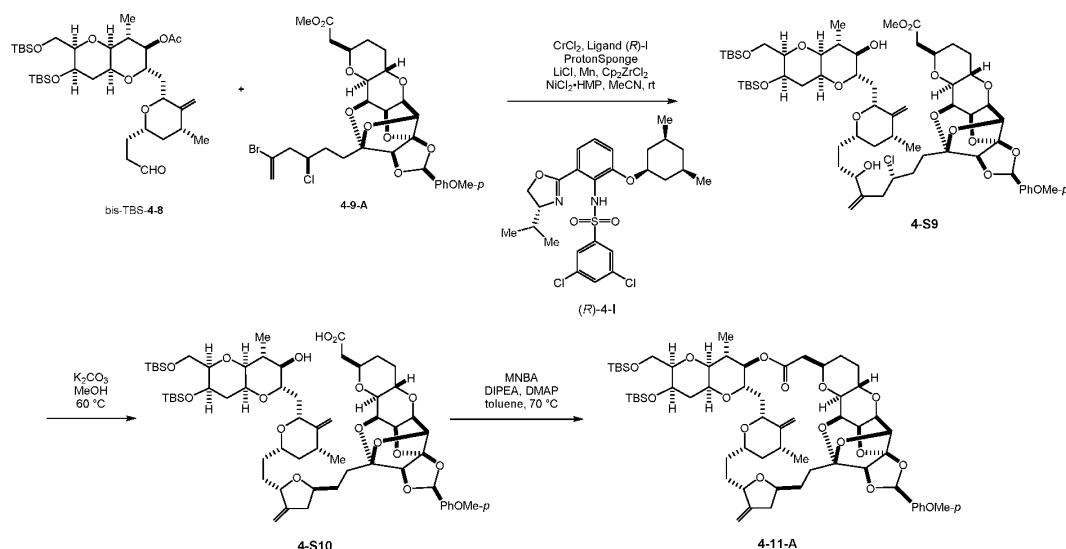
[00758] To a stirred solution of alcohol (1.30 g, 2.77 mmol) in CH_2Cl_2 (26 mL) was added pyridine (1.35 mL, 16.6 mmol), acetic anhydride (0.79 mL, 8.3 mmol) at 0°C . The reaction was stirred for 1 h at room temperature, added MeOH at 0°C , and further stirred for 5 h at room temperature. Sat. NaHCO_3 aq. was added to the reaction and the aqueous phase was extracted with CH_2Cl_2 three times. Combined organic phases were dried over Na_2SO_4 and evaporated. The crude product was purified by flash column chromatography on silica gel (3% EtOAc in hexanes) to provide acetate (1.35 g, 2.64 mmol, 96%) as a pale oil.

[00759] To a stirred solution of acetate (1.35 g, 2.64 mmol, 1 eq.) in CH_2Cl_2 (26 mL) and MeOH (26 mL) was added CSA (30.7 mg, 0.089 mmol, 5 mol%) at room temperature. After being stirred for 3 h, the reaction was quenched with Et_3N (0.2 mL) and concentrated under

reduced pressure. The residue was dissolved in EtOAc and sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, filtered, and concentrated under reduced pressure. This crude was used in the next reaction without further purification.

[00760] To a mixture of alcohol (2.644 mmol, 1 eq.) and 2,6-lutidine (2.2 mL, 18.7 mmol) in CH₂Cl₂ (30 mL) was added TBSOTf (2.15 mL, 9.3 mmol) dropwise at -78 °C and the reaction mixture was stirred for 30 min at the same temperature prior to the addition of sat. NaHCO₃ aq. The aqueous phase was extracted with CH₂Cl₂ twice and combined organic phases were dried over Na₂SO₄, concentrated under reduced pressure. The crude material was connected to high vacuum pump to remove remaining 2,6-lutidine. The obtained residue was purified by flash column chromatography on silica gel (10% then 20% EtOAc in hexanes) to give TBS ether (1.83 g, 99% for 2 steps) as a pale oil.

[00761] To a solution of bis-TBS ether (1.40 g, 2.0 mmol, 1 eq.) in CH₂Cl₂ (20 mL) was added DIBAL (5 mL of 1.0 M in Hexanes, 5 mmol, 2.5 eq.) dropwise at -78 °C. After being stirred for 30 min at the same temperature, the reaction was quenched with acetone (0.30 mL) and MeOH (0.30 mL). Then 20% Rochelle's salt aq. was added, and the mixture was stirred for 2 h at room temperature. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, filtrated, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (5%, 20% then 33% EtOAc in Hexanes) to give bis-TBS-4-8 (1.105 g, 1.76 mmol, 88%) as a colorless amorphous solid. bis-TBS-4-8: $[\alpha]_D^{20}$ -22.4 (c = 1.0, CDCl₃). ¹H NMR (600 MHz, C₆D₆) δ: 9.40 (1H, br s), 4.89 (1H, br s), 4.73 (1H, br d, *J* = 1.9 Hz), 4.29 (1H, q, *J* = 5.6 Hz), 3.88 (1H, t, *J* = 6.1 Hz), 3.83 (2H, d, *J* = 6.2 Hz), 3.68 (1H, td, *J* = 3.5, 1.6 Hz), 3.60 (1H, td, *J* = 3.7, 2.1 Hz), 3.59-3.50 (1H, m), 3.36 (1H, t, *J* = 5.6 Hz), 3.26 (1H, td, *J* = 6.2, 1.7 Hz), 3.24-3.18 (1H, m), 3.05 (1H, t, *J* = 2.4 Hz), 2.32-2.25 (1H, m), 2.17- 2.05 (4H, m), 2.00 (1H, dt, *J* = 14.6, 3.4 Hz), 1.94-1.83 (1H, m), 1.48-1.34 (2H, m), 1.27 (1H, dt, *J* = 14.5, 3.9 Hz), 1.22-1.17 (1H, m), 1.01 (9H, s), 1.0 (3H, d, *J* = 6.0 Hz), 0.97 (9H, s), 0.86 (3H, d, *J* = 6.5 Hz), 0.84-0.76 (1H, m), 0.11 (3H, s), 0.10 (6H, s), 0.05 (3H, s). ¹³C NMR(125 MHz, C₆D₆) δ: 200.2, 150.8, 128.2, 104.4, 80.9, 78.9, 76.4, 75.7, 74.4, 72.1, 64.2, 63.4, 62.6, 42.4, 39.8, 39.4, 35.6, 35.4, 32.6, 27.9, 25.8, 25.7, 18.2, 18.1, 17.7, 16.2, -4.3, -5.2, -5.3, -5.4. IR (neat) ν 3518, 2955, 2928, 2855, 1726, 1471, 1462, 1372, 1252, 1092, 836, 755 cm⁻¹. HRMS (ESI) calcd. for C₃₃H₆₃O₇Si₂ [M+H]⁺ : 627.4107, found 627.4108.



[00762] In a glove box, to a mixture of CrCl₂ (3.9 mg, 10 mol%), sulfonamide ligand (R)-I (24.1 mg, 13 mol%), proton scavenger (8.2 mg, 12 mol%) and LiCl (13.2 mg, 0.32 mmol, 1.0 eq) was added MeCN (0.8 mL, 0.4 M) and stirred for 1 h at r.t. In a separate flask, bis-TBS-4-8 (0.20 g, 0.32 mmol, 1.0 eq), 4-9-A (0.23 g, 0.35 mmol, 1.1 eq), DTBMP (0.16 g, 0.79 mmol, 2.5 eq), Mn (70.1 mg, 1.28 mmol, 4.0 eq) and Cp₂ZrCl₂ (0.23 g, 0.79 mmol, 2.5 eq) and the above Cr complex solution was transferred, and mixture was stirred for 1 min, then NiCl₂·HMP (1.3 mg, 2 mol%, 0.02 eq, doped in LiCl) was added. Additional NiCl₂·HMP (1.3 mg each, 2 mol%, 0.01 eq, twice, doped in LiCl) was added after 1 and 2 h respectively. In total the reaction mixture was stirred for 3.5 h at r.t. The reaction was removed from the glove box and diluted with EtOAc (10 mL). Aqueous potassium serinate (0.5M, 2 mL) and saturated aqueous NaHCO₃ (2 mL) were added. After being stirred for 1 h, the resulting suspension was filtered through a pad of celite. The filtrate was extracted with EtOAc, washed with sat. NaCl, dried over Na₂SO₄, filtered, and concentrated. Without further purification, the crude was used in next reaction.

[00763] To a solution of 4-S9 (estimated as 0.32 mmol, 1.0 eq) in MeOH (8 mL) was added K₂CO₃ (0.44 g, 3.2 mmol, 10.0 eq). The reaction was heated to 60 °C and stirred for 15 h. Then H₂O (0.4 mL) was added. After being stirred for additional 3 h at the same temperature, the mixture was cooled to room temperature, filtered through a pad of Celite, and concentrated. The residue was dissolved in EtOAc, sat. NH₄Cl, and sat. NaCl. The mixture was extracted for five times with EtOAc. The combined organic layers were dried over

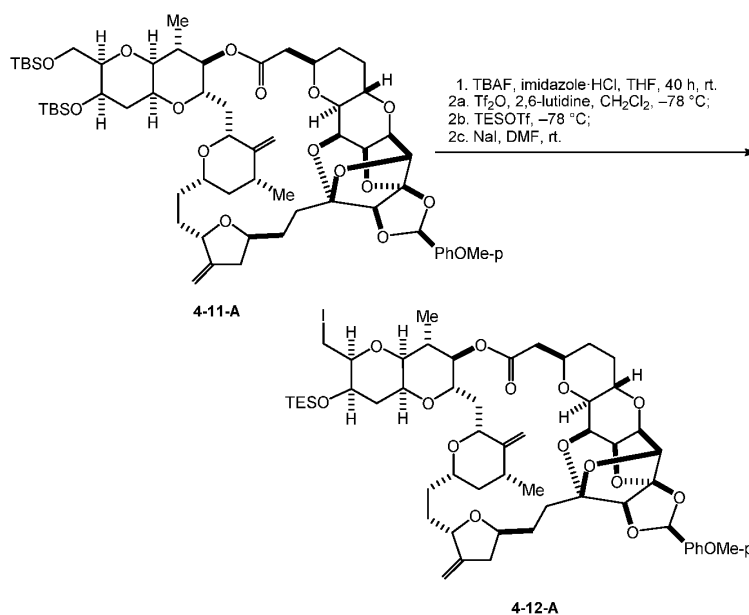
Na₂SO₄, filtered, and concentrated. Without further purification, the crude product was used in the next reaction.

[00764] To a solution of MNBA³ (Shiina's reagent, 0.658 g, 1.91 mmol, 6.0 eq) in hot toluene (200 mL) at 70 °C was added DMAP (0.47 g, 3.83 mmol, 12.0 eq). A solution of carboxylic acid **S10** (estimated as 0.32 mmol, 1.0 eq) and DIPEA (0.66 mL, 3.83 mmol, 12.0 eq) in toluene (20 mL) was added to the reaction mixture via syringe pump over 15 h. After completion of addition, the syringe was washed with toluene (20 mL) and stirred for additional 30 min. The reaction was cooled to room temperature, and concentrated. The residue was dissolved in CH₂Cl₂ and washed with 0.5 M HCl, sat. NaHCO₃ successively. The organic layer was dried over Na₂SO₄, filtered, and concentrated. Flash chromatography (neutral SiO₂, Hexanes/EtOAc= 40/1, 20/1, 5/1 then 4/1 gave white crystalline solid **4-11-A** (211 mg, 0.19 mmol, 54% over 3 steps). The overall stereoselectivity (*dr* = 25:1) was estimated from ¹H NMR spectrum of **4-11-A** thus obtained.

[00765] Under the condition given on page S18, Ni/Cr-Mediated coupling of bis-TBS-**4-8** with **4-9-A** in the presence of Cr-catalyst prepared from (*R*)-**4-G** and (*R*)-**4-H** was carried out. Following the procedure given above, the crude coupling product was converted to the macrolactone acetonide, to determine the overall yield and stereoselectivity; sulfonamide (*R*)-**4-H** gave 58% overall yield from bis-TBS-**4-8** (200-mg scale) and *dr* = 28:1, whereas sulfonamide (*R*)-**4-G** gave 58% overall yield from bis-TBS-**4-8** (200-mg scale) and *dr* = 27:1. **11-A**: [α]_D²⁰ -54.8 (c = 1.0, CDCl₃). mp = 88-89 °C. ¹H NMR (600 MHz, C₆D₆) δ : 7.31 (2H, d, *J* = 8.8 Hz), 6.68 (2H, d, *J* = 8.7 Hz), 6.03 (1H, s), 5.11 (1H, br s), 5.02 (1H, br s), 4.89 (1H, br s), 4.76–4.71 (3H, m), 4.61 (1H, d, *J* = 10.4 Hz), 4.44 (1H, td, *J* = 10.5, 10.0, 4.3 Hz), 4.36 (1H, brs), 4.32 (1H, dt, *J* = 5.3, 0.9 Hz), 3.4 (1H, q, *J* = 3.9 Hz), 3.26 (1H, td, *J* = 6.2, 1.9 Hz), 3.19 (3H, s), 3.09 (1H, dd, *J* = 4.7, 3.1 Hz), 2.72 (1H, dd, *J* = 16.8, 7.7 Hz), 2.62 (1H, ddt, *J* = 15.5, 7.7, 2.4 Hz), 2.48 (1H, dd, *J* = 9.5, 1.8 Hz), 2.35 (1H, dd, *J* = 16.8, 4.4 Hz), 2.32–2.18 (6H, m), 1.97 (1H, dt, *J* = 11.6, 3.9 Hz), 1.88 (1H, ddt, *J* = 14.2, 11.4, 3.1 Hz), 1.77 (1H, tt, *J* = 13.6, 3.7 Hz), 1.70–1.62 (1H, m), 1.49 (1H, dddd, *J* = 11.9, 7.3, 5.6, 2.5 Hz), 1.43 (2H, dt, *J* = 12.5, 2.8 Hz), 1.37 (1H, dt, *J* = 14.6, 4.5 Hz), 1.33–1.19 (3H, m), 1.16 (3H, d, *J* = 7.3 Hz), 1.14–1.0 (2H, M), 0.99 (9H, s), 0.98 (9H, s), 0.14 (3H, s), 0.12 (6H, s), 0.09 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 170.7, 161.0, 152.6, 152.4, 128.4, 128.3, 118.6, 113.7, 108.9, 108.6, 104.6, 103.4, 89.8, 83.4, 79.7, 78.0, 77.8, 77.4, 75.8, 75.6, 75.4, 74.4, 73.9, 73.6, 73.5, 73.4, 69.7, 68.0, 64.13, 64.09, 63.3, 54.4, 43.7, 40.9, 39.2, 39.0, 38.2, 36.0, 35.3, 31.9, 30.9, 30.5, 29.97, 29.9, 27.3, 26.0, 25.9, 25.8, 18.3, 17.8, 16.4, -4.3, -5.0, -5.3, -5.4 ppm.

IR (neat) ν 2952, 2927, 2855, 1732, 1615, 1518, 1251, 1092, 1010, 834, 775, 756 cm^{-1} .

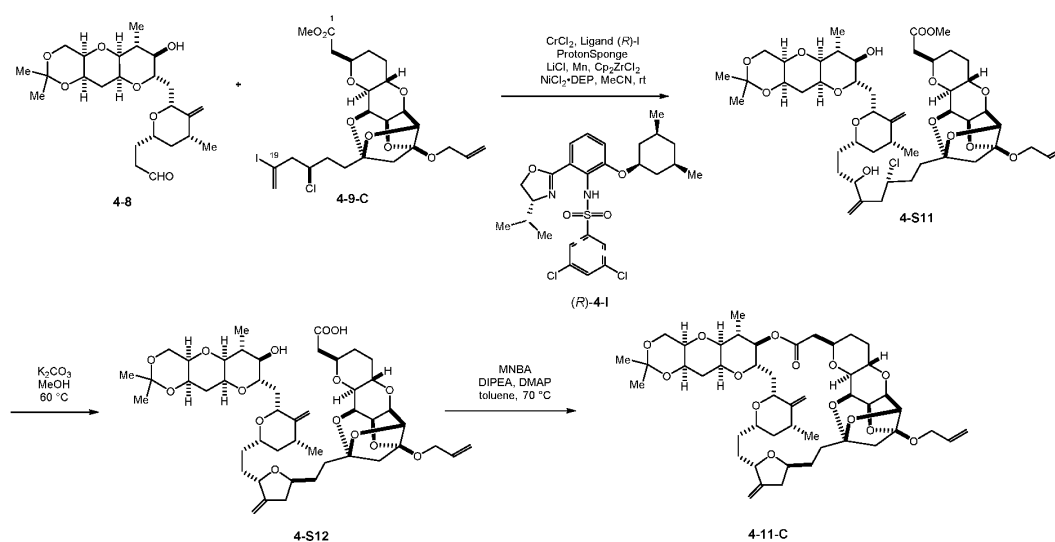
HRMS (ESI) calcd. for $\text{C}_{61}\text{H}_{92}\text{NaO}_{16}\text{Si}_2$ $[\text{M}+\text{Na}]^+$: 1159.7774, found 1159.7816.



[00766] To a solution of **4-11-A** (720 mg, 0.633 mmol) in THF (12.7 mL) was added TBAF solution (0.95 M in THF, buffered with 0.5 eq of imidazole·hydrochloride, 3.2 mL, 3.17 mmol) at room temperature. After stirring for 40 h at room temperature, solvent was removed by a stream of nitrogen gas and redissolved in EtOAc and water. The aqueous layer was extracted with EtOAc four times and the combined organic layer was dried over Na_2SO_4 , filtered, and concentrated. Without further purification, the crude product was used in the next reaction.

[00767] To a stirred solution of crude diol (estimated as 0.633 mmol) in CH_2Cl_2 (6.3 mL) was added 2,6-lutidine (0.733 mL, 6.33 mmol, 10 eq.). The mixture was cooled to $-78\text{ }^\circ\text{C}$, and then added Ti_2O (0.13 mL, 0.760 mmol, 1.2 eq.). After being stirred for 15 min at the same temperature, TESOTf (0.23 mL, 1.01 mmol, 1.6 eq) was added. The reaction was warmed to $0\text{ }^\circ\text{C}$ with ice bath. After being stirred for additional 20 min, to the reaction were added DMF (12.6 mL) and NaI (0.475 g, 3.17 mmol, 5 eq). The resulting mixture was allowed to warm to room temperature and stirred for 2.5 h. Then the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with TBME. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (9%, 20%, 30%, then 40% EtOAc in Hexanes) to give iodide **4-12-A** (595 mg, 0.525 mmol, 83% for 2 steps), which was a mixture of C20-epimer (dr = ~20:1), as a colorless

solid. The C20-epimer was separated by HPLC purification (Column: DuPont Instruments ZORBAL SIL 21.2 mm x 25 cm (880952-101), Solvent: 3% *i*PrOH in Hexanes, Flow rate: 10.0 mL/min., Detection: UV at 254 nm and 220 nm, $t_{R(\text{desired})}$ = 17 min. $t_{R(\text{C20-epimer})}$ = 21 min.). **4-12-A**: white solid, $[\alpha]_D^{20}$ -50.0 (*c* 0.65, CHCl₃). MP: 115-117 °C (recrystallized from H₂O-*i*PrOH). ¹H NMR (600 MHz, C₆D₆) δ : 7.36 (2H, d, *J* = 8.8 Hz), 6.73 (2H, d, *J* = 8.8 Hz), 6.09 (1H, s), 5.17 (1H, s), 5.11 (1H, d, *J* = 1.2 Hz), 4.94 (1H, s), 4.80-4.77 (3H, m), 4.66 (1H, brd, *J* = 11.2 Hz), 4.49 (1H, ddd, *J* = 11.2, 11.2, 4.2 Hz), 4.40 (1H, d, *J* = 1.8 Hz), 4.37 (1H, dd, *J* = 5.4, 1.2 Hz), 4.06 (1H, s), 4.01 (1H, dd, *J* = 6.0, 6.0 Hz), 4.00-3.96 (2H, m), 3.79 (1H, d, *J* = 11.4 Hz), 3.73-3.68 (3H, m), 3.36-3.28 (3H, m), 3.22 (3H, s), 3.20 (1H, ddd, *J* = 6.6, 6.6, 1.8 Hz), 3.04 (1H, dd, *J* = 4.2, 3.0 Hz), 2.77 (1H, dd, *J* = 16.2, 7.8 Hz), 2.66-2.62 (1H, m), 2.51 (1H, dd, *J* = 9.9, 1.5 Hz), 2.39-2.24 (6H, m), 2.19-1.94 (7H, m), 1.85-1.82 (1H, m), 1.76-1.71 (1H, m), 1.55-1.27 (6H, m), 1.18 (3H, d, *J* = 7.2 Hz), 1.08 (1H, dd, *J* = 12.6, 11.4 Hz), 1.03 (9H, t, *J* = 7.9 Hz), 1.01 (3H, d, *J* = 6.5 Hz), 0.70 (6H, q, *J* = 8.2 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 171.2, 161.4, 153.0, 152.6, 119.0, 114.1, 109.3, 109.0, 105.1, 103.7, 90.3, 83.8, 79.8, 78.7, 78.1, 77.9, 76.0, 75.6, 74.7, 74.1, 74.0, 73.8, 73.7, 69.9, 68.4, 64.8, 64.0, 54.8, 44.0, 41.4, 39.4, 39.4, 39.0, 36.4, 36.3, 32.4, 31.1, 30.9, 30.3, 30.2, 27.6, 18.2, 16.6, 7.3($\times 3$), 5.7, 5.3($\times 3$) ppm. IR (film): 2953, 2935, 2875, 1731, 1615, 1518, 1251, 1092, 1011, 907, 831 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₅₅H₇₇IO₁₅SiNa, 1155.3969; found, 1155.3987.



[00768] In a glove box, to a mixture of CrCl₂ (7.0 mg, 10 mol%), sulfonamide ligand (R)-4-I (43.2 mg, 13 mol%), proton scavenger (14.7 mg, 12 mol%) and LiCl (24.1 mg, 0.57 mmol,

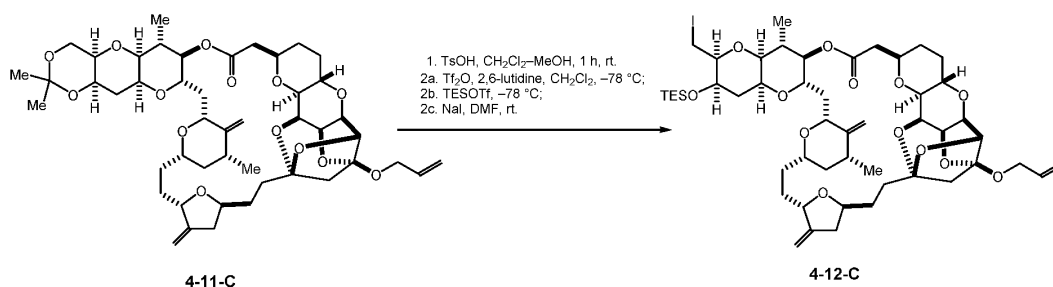
1.0 eq) was added MeCN (1.4 mL, 0.4 M) and stirred for 1 h at r.t. In a separate flask, **4-8** (0.25 g, 0.57 mmol, 1.0 eq), **4-9-C** (0.35 g, 0.63 mmol, 1.1 eq), DTBMP (0.29 g, 1.43 mmol, 2.5 eq), Mn (125 mg, 2.28 mmol, 4.0 eq) and Cp₂ZrCl₂ (0.42 g, 1.43 mmol, 2.5 eq) and the above Cr complex solution was transferred, and mixture was stirred for 1 min, then NiCl₂•DEP (2.2 mg, 1 mol%, 0.01 eq, doped in LiCl) was added. Additional NiCl₂•DEP (2.2 mg each, 1 mol%, 0.01 eq, twice, doped in LiCl) was added after 1 and 2 h respectively. In total the reaction mixture was stirred for 3 h at r.t. The reaction was removed from the glove box and diluted with EtOAc (10 mL). Aqueous potassium serinate (0.5M, 3 mL) and saturated aqueous NaHCO₃ (3 mL) were added. After being stirred for 1 h, the resulting suspension was filtered through a pad of celite. The filtrate was extracted with EtOAc, washed with sat. NaCl, dried over Na₂SO₄, filtered, and concentrated. Without further purification, the crude was used in next reaction.

[00769] To a solution of **4-S11** (estimated as 0.57 mmol, 1.0 eq) in MeOH (10 mL) was added K₂CO₃ (0.79 g, 5.71 mmol, 10.0 eq). The reaction was heated to 60 °C and stirred for 15 h. Then H₂O (0.6 mL) was added. After being stirred for additional 3 h at the same temperature, the mixture was cooled to room temperature, filtered through a pad of Celite, and concentrated. The residue was dissolved in EtOAc, sat. NH₄Cl, and sat. NaCl (adjust pH = ~7 by adding few drops of 1N HCl). The mixture was extracted for five times with EtOAc. The combined organic layers were dried over Na₂SO₄, filtered, and concentrated. Without further purification, the crude product was used in the next reaction.

[00770] To a solution of MNBA (1.18 g, 3.43 mmol, 6.0 eq) in hot toluene (380 mL) at 70 °C was added DMAP (0.84 g, 6.85 mmol, 12.0 eq). A solution of carboxylic acid **4-S12** (estimated as 0.57 mmol, 1.0 eq) and DIPEA (1.2 mL, 6.85 mmol, 12.0 eq) in toluene (27 mL) was added to the reaction mixture via syringe pump over 15 h. After completion of addition, the syringe was washed with toluene (20 mL) and stirred for additional 30 min. The reaction was cooled to room temperature, and concentrated. The residue was dissolved in CH₂Cl₂ and washed with 0.5 M HCl, sat. NaHCO₃ successively. The organic layer was dried over Na₂SO₄, filtered, and concentrated. Flash chromatography (neutral SiO₂, Hexanes/EtOAc= 7/1, 5/1, 3/1, 1/1, 1/2 then 1/3 gave white crystalline solid **4-11-C** (307 mg, 0.36 mmol, 61% over 3 steps). The overall stereoselectivity (*dr* = 17:1) was estimated from ¹H NMR spectrum of **4-11-C** thus obtained. **4-11-C**: [α]_D²⁰ -52.2 (c = 1.0, CDCl₃). mp = 157-158 °C. ¹H NMR (600 MHz, C₆D₆) δ: 5.71 (1H, ddt, *J* = 17.3, 10.6, 5.3 Hz), 5.18 (1H, brs), 5.11 (1H, dq, *J* = 17.3, 1.7 Hz), 5.05 (1H, q, *J* = 2.2 Hz), 5.02 – 4.91 (2H, m), 4.90 (1H, s), 4.77 (1H, dd, *J* = 7.7, 5.0 Hz), 4.72 (1H, d, *J* = 1.8 Hz), 4.62 (1H, dt, *J* = 10.3, 2.4 Hz), 4.54 –

4.47 (1H, m), 4.35 (1H, dd, $J = 4.2, 2.0$ Hz), 4.30 – 4.26 (1H, m), 4.11 – 4.00 (2H, m), 3.94 (1H, m), 3.83 – 3.73 (3H, m), 3.72 – 3.62 (m, 3H), 3.53 – 3.45 (2H, m), 3.00 (1H, dd, $J = 5.1, 4.0$ Hz), 2.80 (1H, dd, $J = 16.5, 7.2$ Hz), 2.73 – 2.63 (2H, m), 2.53 (1H, dd, $J = 9.6, 2.1$ Hz), 2.46 – 2.37 (2H, m), 2.29 (2H, dt, $J = 19.4, 7.2, 4.9, 2.1$ Hz), 2.25 – 2.09 (4H, m), 2.09 – 1.98 (4H, m), 1.92 – 1.80 (2H, m), 1.75 – 1.62 (1H, m), 1.56 – 1.40 (4H, m), 1.40 – 1.23 (3H, m), 1.19 (5H, d, $J = 6.9$ Hz), 1.02 (1H, td, $J = 12.4, 11.0$ Hz), 0.93 (2H, d, $J = 6.4$ Hz) ppm. ^{13}C NMR(125 MHz, C_6D_6) δ : 170.9, 152.5, 152.3, 134.3, 116.3, 115.9, 108.9, 104.6, 103.4, 98.4, 82.9, 77.9, 77.6, 76.9, 75.9, 75.7, 74.7, 74.7, 73.9, 73.7, 73.6, 70.0, 69.5, 68.1, 65.6, 63.7, 63.5, 62.1, 51.3, 43.4, 41.1, 39.0, 38.9, 38.4, 35.9, 35.6, 32.2, 31.9, 30.7, 30.7, 30.1, 28.5, 28.1, 19.8, 17.8, 16.1 ppm. IR (neat) ν 2925, 2867, 1729, 1373, 1311, 1186, 1118, 1073, 995, 911, 832, 756 cm^{-1} . HRMS (ESI) calcd. for $\text{C}_{47}\text{H}_{66}\text{NaO}_{14}$ $[\text{M}+\text{Na}]^+$: 877.4345, found 821.4353.

[00771] Under the condition given on page S18, Ni/Cr-Mediated coupling of **4-8** with **4-9-C** in the presence of Cr-catalyst prepared from (*R*)-**4-G** and (*R*)-**4-H** was carried out. Following the procedure given above, the crude coupling product was converted to the macrolactone acetonide, to determine the overall yield and stereoselectivity; sulfonamide (*R*)-**4-H** gave 63% overall yield from **8** (250-mg scale) and $dr = 20:1$, whereas sulfonamide (*R*)-**4-G** gave 63% overall yield from **4-8** (250-mg scale) and $dr = 21:1$.



[00772] To a stirred solution of **4-11-C** (700 g, 0.819 mmol, 1 eq., $dr = \sim 20:1$) in CH_2Cl_2 (8.2 mL) and MeOH (8.2 mL) was added *p*-TsOH· H_2O (3.1 mg, 0.016 mmol, 2 mol%) at room temperature. After being stirred for 1 h, the reaction was quenched with Et_3N (0.1 mL) and concentrated under reduced pressure. The residue was dissolved in EtOAc and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude diol, which was used in the next reaction without further purification.

[00773] To a stirred solution of crude diol (estimated as 0.819 mmol) in CH_2Cl_2 (8.2 mL) was added 2,6-lutidine (0.57 mL, 4.91 mmol, 6 eq.). The mixture was cooled to -78°C , and then added Tf_2O (0.18 mL, 1.1 mmol, 1.2 eq.). After being stirred for 15 min at the same temperature, TESOTf (0.30 mL, 1.31 mmol, 1.5 eq) was added. The reaction was warmed to 0°C with ice bath. After being stirred for additional 20 min, to the reaction were added DMF (16.4 mL) and NaI (650 g, 4.43 mmol, 5 eq). The resulting mixture was allowed to warm to room temperature and stirred for 6 h. Then the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with TBME. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on neutral silica gel (9%, 13%, 20%, then 30% EtOAc in Hexanes) to give iodide **4-12-C** (689 mg, 0.663 mmol, 81% for 2 steps), which was a mixture of C20-epimer (dr = ~20:1), as a colorless solid. The C20-epimer was separated by HPLC purification (Column: DuPont Instruments ZORBAL SIL 21.2 mm x 25 cm (880952-101), Solvent: 2% *i*PrOH in Hexanes, Flow rate: 10.0 mL/min., Detection: UV at 254 nm and 220 nm, $t_{\text{R(desired)}}$ = 22.8 min. $t_{\text{R(C20-epimer)}}$ = 28.0 min.). **4-12-C**: colorless solid, $[\alpha]_{\text{D}}^{20}$ -46.8 (*c* 1.0, CHCl_3). MP: $84-86^\circ\text{C}$ (recrystallized from H_2O -*i*PrOH). ^1H NMR (600 MHz, C_6D_6) δ : 5.75 (1H, dddd, J = 17.3, 10.5, 5.1, 5.1 Hz), 5.15 (1H, ddd, J = 17.4, 1.8, 1.8 Hz), 5.14 (1H, s), 5.11 (1H, s), 4.97 (1H, ddd, J = 10.2, 1.8, 1.8 Hz), 4.90 (1H, s), 4.77-4.72 (3H, m), 4.62 (1H, d, J = 11.2 Hz), 4.49 (1H, ddd, J = 10.2, 10.2, 4.4 Hz), 4.35 (1H, ddd, J = 1.8, 1.8, 1.8 Hz), 4.31 (1H, d, J = 4.8 Hz), 4.11 (1H, dd, J = 6.6, 4.8 Hz), 4.0-3.97 (1H, m), 3.83-3.80 (1H, m), 3.74 (1H, d, J = 11.4 Hz), 3.72 (1H, ddd, J = 5.4, 1.8, 1.8 Hz), 3.70-3.67 (2H, m), 3.32-3.27 (3H, m), 3.19 (1H, ddd, J = 6.6, 6.6, 1.8 Hz), 3.03 (1H, dd, J = 4.8, 3.6 Hz), 2.74 (1H, dd, J = 17.4, 7.8 Hz), 2.71-2.67 (1H, m), 2.56 (1H, dd, J = 9.6, 1.8 Hz), 2.33 (1H, dd, J = 16.8, 4.2 Hz), 2.29-2.20 (5H, m), 2.23-1.97 (8H, m), 1.93-1.89 (1H, m), 1.72-1.66 (1H, m), 1.53-1.28 (8H, m), 1.14 (3H, d, J = 7.2 Hz), 1.05 (1H, dd, J = 12.6, 10.8 Hz), 1.00 (9H, t, J = 8.4 Hz), 0.98 (3H, d, J = 7.2 Hz), 0.66 (6H, q, J = 7.8 Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 171.4, 153.0, 152.6, 134.6, 128.3, 116.6, 116.2, 109.3, 105.0, 103.7, 83.3, 79.8, 78.6, 77.9, 77.7, 76.0, 75.6, 75.1, 75.0, 74.7, 74.1, 73.9, 70.0, 68.4, 65.9, 64.9, 63.9, 51.7, 43.9, 41.3, 39.4, 39.3, 38.9, 36.3, 36.2, 35.9, 32.5, 31.1, 30.7, 30.5, 28.5, 18.1, 16.6, 7.3($\times 3$), 5.7, 5.3($\times 3$) ppm. IR (film): 2952, 2948, 2875, 1733, 1457, 1338, 1218, 1138, 1073, 831 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{50}\text{H}_{75}\text{IO}_{13}\text{SiNa}$, 1061.3914; found, 1061.3919.

Synthesis of Left Halves

[00774] One of the central questions was how to construct the [6,6]-spiroketal in a stereoselective manner. Based on precedents, the oxy-Michael process of **E** → **F** under the basic conditions was used (*Figure 15*). The first precedent was found in the synthetic work on polyether antibiotic (-)-A23187 (calcimycin) (see, *e.g.*, Negri, D. P.; Kishi, Y. *Tetrahedron Lett.*, **1987**, 28, 1063); on treatment with a catalytic amount of NaOMe, **G** was transferred exclusively to **H**, the relative stereochemistry of which corresponded to that of **F**.

[00775] The second precedent was found in halichondrin syntheses. As the model study for the original enone route, the oxy-Michael reaction on **I** was studied; on treatment with DBU and LiCl in CH₂Cl₂, **I** gave selectively **J**, which isomerized to **K** on treatment with DBU and LiCl in MeCN or DBU and LiBr in CH₂Cl₂. Alternatively, **K** was obtained directly from **I** on treatment with Hunig base, LiBr in CH₂Cl₂ MeCN. These results indicated that **J** was as the kinetically controlled product, whereas **K** was as the thermodynamically controlled product. The stereochemistry of **J** and **K** was reduced from NOE experiments; in particular, a strong cross peak was observed between C39-H and C48-H of **J-a**, whereas a strong cross peak was detected between C40-H and C48-H of **K-a**, cf. red double-headed arrows.

[00776] A primitive estimation was made on the relative stability of **J** and **K**, on the basis of the literature-known data on A-values of cyclohexane and tetrahydropyran, destabilization energy due to 1,3-diaxial interaction, and destabilization-energy due to boat-conformation. The preferred conformation of **K** was assumed to be **K-a**, whereas that of **J** to be either **J-a**, **J-b**, or half-boat conformer (not shown). For the simplicity of analysis, conformers **K-a**, **J-a**, and **J-b** were used. Around the structure indicated by a dotted box, all of them had the same structural arrangement, including the double stereoelectronic stabilization. Therefore, this structural moiety was excluded from the analysis. A-value for Me-group at 2-position of tetrahydropyran was reported to be 2.86 kcal/mol, which was roughly 1.6 times of A-value for Me-group of cyclohexane (see, *e.g.*, Eliel, E. L.; Hargrave, K. D.; Pietrusiewicz, K. M.; Manoharan, M. *J. Am. Chem. Soc.* **1982**, 104, 3635). 1,3-Diaxial interaction of CH₂R/OMe on cyclohexane was known around 1.9 kcal/mol, which would correspond to ~3 kcal/mol (1.6 x 1.9 kcal/mol) on tetrahydropyran (see, *e.g.*, Corey, E. J.; Feiner, N. F. *J. Org. Chem.* **1980**, **45**, 765 and references cited therein). In addition, a boat conformer of cyclohexane and tetrahydropyran was estimated to be destabilized by 5.5 and 3.9 kcal/mol, respectively, relative to the corresponding chair conformer. Thus, we would assume the destabilization due to the boat conformation in **J-b** to be in the range of 3.9~5.5 kcal/mol (see, *e.g.*, Eliel, E. L.;

Allinger, N. L. Angyal, S. J.; Morrison, G. A. *Conformational Analysis*; John Wiley: New York, 1965, p 244). Counting these factors, **K** was roughly estimated to be energetically favored over **J** by 3~4 kcal/mol.

[00777] To test the feasibility of the synthetic plan proposed above, studied first was the synthesis of 2-pyridyl thioester **8** for two reasons. First, γ -lactone **3** was readily available from **2**, which was the starting material for the synthesis of C27-C37 building block and available from D-galactal (**1**) in a large quantity (see, *e.g.*, Chen, C.-L.; Namba, K.; Kishi, Y. *Org. Lett.* **2009**, *11*, 409). Second, 2-pyridyl thioester **8** should be a good model substrate to study the following Zr/Ni-mediated one-pot ketone synthesis.

[00778] *Figure 16* summarizes the synthesis of **8** from γ -lactone **3**. Thus, γ -lactone **3** was straightforwardly transformed to δ -lactone **4** in an excellent overall yield. The coupling of **4** and **5-5** was achieved by slow addition of *t*-BuLi into a mixture of the two substrates in THF. The unsaturated benzyl ester group was incorporated on **6** in 3 steps, *i.e.*, (1) dihydroxylation of the terminal olefin, (2) oxidative cleavage, and (3) Horner-Emmons reaction, to furnish **7** in an excellent overall yield. Alternatively, the transformation of **6** into **7** was realized in one step with metathesis (see, *e.g.*, Trnka, T. M.; Grubbs, R. H. *Acc. Chem. Res.* **2001**, *34*, 18; Schrock, R. R. *Adv. Syn. Catal.* **2007**, *349*, 41; Hoveyda, A. H.; Zhugralin, A. R. *Nature* **2007**, *450*, 243; Araki, M.; Sakata, S.; Takei, H.; Mukaiyama, T. *Bull. Chem. Soc. Jpn.* **1974**, *47*, 1777). Because of the cost-effectiveness, the former three-step procedure was chosen for preparative purpose.

[00779] The behavior of **7** in oxy-Michael process was similar to the case of **I** into **K**. On treatment with DBU and LiBr in MeCN at room temperature, **7** initially gave a 2:1 mixture of kinetically- and thermodynamically-controlled products, which equilibrated almost exclusively to the thermodynamically controlled product at room temperature for 12 hours. In this transformation, the benzyl ester was partially hydrolyzed, but it was found that the hydrolysis of benzyl ester was completely suppressed by addition of BnOAc. Preparatively, on treatment with DBU (5 equiv)/LiBr (10 equiv)/BnOAc (10 equiv.), **7** gave the desired, thermodynamically controlled product with in an excellent overall yield. The stereochemistry of oxy-Michael product was confirmed by NOE experiments; a strong cross peak was observed between C40-H and C48-H. Finally, the necessary functional group adjustment was made in four steps, *i.e.*, (1) deprotection of C41 MPM group, (2) protection of the resultant alcohol with TES, (3) debenzylation, and (4) thioester formation. 2-Pyridyl thioester **8** thus

obtained was fully characterized and was found stable to store at -20 °C in dark. The overall yield of **8** from **3** was >50% in a 5-g scale preparation.

[00780] *Figure 17* summarizes the synthesis of left half in the halichondrin series. This synthesis was designed on the basis of the knowledge gained in the synthesis of C38-C52 building block discussed in the preceding section, coupled with the original synthesis of the left half of halichondrin B. Epoxy γ -lactone **9** was synthesized from L-gulono- γ -lactone by the method reported in 1992 (see, *e.g.*, Buszek, K. R.; Fang, F. G.; Forsyth, C. J.; Jung, S. H.; Kishi, Y.; Scola, P. M.; Yoon, S. K. *Tetrahedron Lett.* **1992**, 33, 1553), whereas *cis*-vinyl iodide **5-10** was prepared from commercially available (*S*)-glycidol in 4 steps. Then, **9** and **10** were coupled with use of a standard Cu-mediated chemistry. With use of 1.8 equivalent of **10**, γ -lactone **11** was isolated in 81% yield. The homoallylic alcohol in **11** was epoxidized with Sharpless VO(TMHD)₂-mediated method, followed by TES-protection of the C48 alcohol, to give epoxy- γ -lactone **12** in 85% overall yield from **11**, with >50:1 stereoselectivity (see, *e.g.*, Sharpless, K. B.; Michaelson, R. C. *J. Am. Chem. Soc.* **1973**, 95, 6136). The next coupling of **12** with **5** was best achieved by slow addition of *t*-BuLi into a mixture of the two substrates in THF, to give **13** in 85% yield, which existed as a mixture of ketol (major) and keto/alcohol (minor). Among acids tested, (PhO)₂P(=O)OH was found to be most effective in facilitating an intramolecular epoxide-ring opening by the masked alcohol at C47. With minor modifications on the transformation outlined in *Figure 16*, **14** was transformed to the left half **17**. Thus, the terminal olefin of **14** was converted to unsaturated benzyl ester **16**. In this series, it was necessary to unmask the protecting group of the C48 alcohol; on treatment with (PhO)₂P(=O)OH, two TES-groups in **14** were selectively removed, to furnish **15**.

[00781] Unsaturated benzyl ester **15** was subjected to oxy-Michael reaction under the condition optimized with **7** (*Figure 16*), to furnish the desired product **16** in 86% yield. The stereochemical behavior of **16** in the oxy-Michael reaction was found to be very similar to that of **7**, including the overall stereoselectivity and the equilibration of the kinetically-controlled product into the thermodynamically-controlled product. In this series, however, one-minor by-product was isolated in ~8% yield. The spectroscopic analysis (HRMS, ¹H- and ¹³C-NMR, IR) suggested this minor product to be **18**, which likely arose via the carbanion-attack to the unsaturated benzyl ester, cf., the red arrow in **L**. The structure information of **18** immediately suggested a possibility of eliminating or suppressing the by-product formation with use of organocatalysts recently reported by Asano, Matsubara and coworkers (see, *e.g.*,

Yoneda, N.; Fukata, Y.; Asano, K.; Matsubara, S. *Angew. Chem. Int. Ed.* **2015**, *54*, 15497). Indeed, an addition of non-chiral thiourea **M** had an expected effect, to give a 2:1 mixture of two diastereomers, free from **18**. In principle, a chiral thiourea should enforce preferentially to form one of the two diastereomers. In practice, however, the asymmetric induction was only modest with chiral organocatalysts. Thus, the 2:1 mixture was subjected to the equilibrating condition, to furnish almost exclusively **16** in 93% overall yield from **15**. Obviously, the equilibration took place via the retro-oxy-Michael/oxy-Michael process. Curiously, however, there was no detectable amount of **18** formed in the equilibration step. Structurally, compared with **7**, **15** had one additional free-hydroxyl group at C51. That additional free-hydroxyl group might affect the overall outcome in the oxy-Michael reaction, because it was carried out under basic conditions. However, it was demonstrated that the substrate with the C51-alcohol protected with TBS did exhibit virtually identical reactivity profiles as **15** did. Finally, the left-half **17** in the halichondrin series was obtained uneventfully from **16** in 82% overall yield.

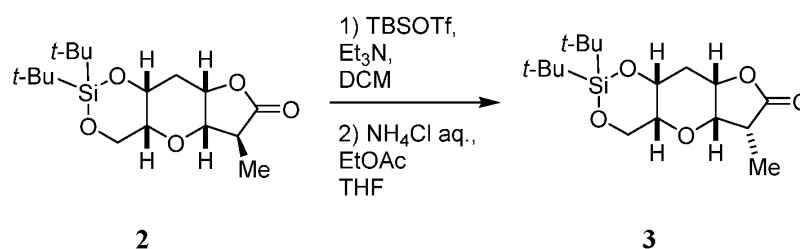
[00782] *Figure 18* summarizes the synthesis of left half in the homohalichondrin series. Once again, this synthesis was based on the knowledge gained through the previous work (see, e.g., Fang, F. G.; Kishi, Y.; Matelich, M. C.; Scola, P. M. *Tetrahedron Lett.* **1992**, *33*, 1557). The synthesis began with γ -lactone **3**, which was transformed to **21** via **20** by standard synthetic operations. Sharpless asymmetric epoxidation (see, e.g., Katsuki, T.; Sharpless, K. B. *J. Am. Chem. Soc.* **1980**, *102*, 5974) was used stereoselectively to install the C53-C55 moiety. Namely, asymmetric epoxidation of the allylic alcohol of **21**, followed by acid treatment, gave triol **22** in an excellent overall yield. Based on the absolute stereochemistry of the epoxide introduced by Sharpless asymmetric epoxidation, the stereochemistry of **22** was assigned as indicated, which was confirmed by X-ray analysis of its C55 mono-3,5-dinitrobenzoate. This experiment proved the C53/C54-stereochemistry previously proposed for the homohalichondrins from NMR analysis. Triol **22** was transferred to δ -lactone **23** and then to unsaturated benzyl ester **24** in a good overall yield. The overall behavior of **24** in the intramolecular oxy-Michael to construct the [6,6]-spiroketal was similar to that observed on **15**. Lastly, 2-pyridyl thioester **26** was secured from **25** as before.

[00783] *Figure 19* summarizes the synthesis of left half in the norhalichondrin series. With two modifications, this synthesis was carried out as the previous cases. First, the C53 terminal in the norhalichondrin series was a carboxylic acid, which was introduced via oxidation of the primary alcohol selectively prepared from **29**. Second, TES-group was

chosen to protect the alcohol at C50. Notably, the behavior of unsaturated benzyl ester **28** in the oxy-Michael transformation was found to be virtually identical with that of **7**. 2-Thiopyridine ester **32** thus synthesized was isolated via neutral silica gel chromatography and fully characterized.

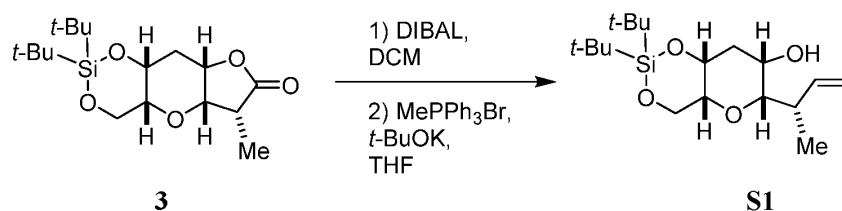
Experimental Procedures for Synthesis of Left Halves

Halichondrin Analogs

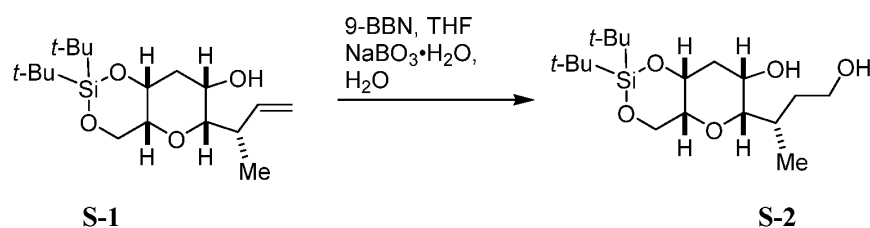


[00784] To a stirred solution of lactone **2** (10.0 g, 29.2 mmol, 1 eq.) and Et₃N (20.0 mL, 143 mmol, 5.0 eq.) in CH₂Cl₂ (100 mL) was added TBSOTf (17.0 mL, 74.0 mmol, 2.5 eq.) at 0 °C. After being stirred for 3 h at room temperature, the reaction was quenched with sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with CH₂Cl₂. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was used in the next reaction without further purification.

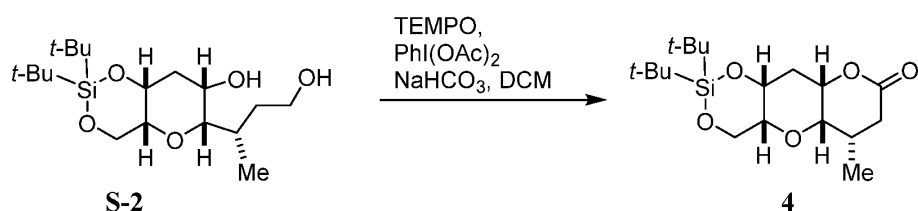
The crude silyl enol ether (calculated as 29.2 mmol, 1 eq.) was dissolved in a mixture of EtOAc (200 mL), THF (30 mL), and sat. NH₄Cl aq. (300 mL). After being stirred for 3 h at 50 °C, the mixture was cooled to room temperature. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The crude lactone was recrystallized from CH₂Cl₂/Hexanes to give lactone **3** (9.68 g from 1st recrystallization and 300 mg from 2nd recrystallization: total 9.98 g, 29.1 mmol, 100% for 2 steps) as a colorless crystal. The spectroscopic data obtained are consistent with those previously reported in the literature.



[00785] To a stirred solution of **3** (4.3 g, 12.6 mmol, 1 eq.) in CH_2Cl_2 (63 mL) at -78°C was added DIBAL solution (16.4 mL of 1.0 M in hexanes, 16.4 mmol, 1.3 eq.). After being stirred for 40 min, the reaction was quenched with MeOH at the same temperature. Then 10% Rochelle's salt aq. was added. The mixture was stirred for 2 h at room temperature to give a clear biphasic mixture. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude lactol, which was used in the next reaction without further purification. To a suspension of $\text{Ph}_3\text{PCH}_3\text{Br}$ (18.0 g, 50.4 mmol, 4 eq.) in THF (50 mL) was added $t\text{-BuOK}$ (4.24 g, 37.8 mmol, 3 eq.) at 0°C . A solution of the crude lactol (calculated as 12.6 mmol) in THF (13 mL) was added into the reaction mixture, and then the ice bath was removed. After being stirred for 1.5 h at room temperature, the reaction was quenched with sat. NH_4Cl aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was dissolved in hexanes-EtOAc (10:1) and passed through a pad of silica gel (9% EtOAc in Hexanes) to give **S-1** (4.15 g, 12.1 mmol, 96% for 2 steps) as a colorless solid. **S-1**: $[\alpha]_D^{20} -36.8$ (c 1.14, CHCl_3). MP: $90\text{--}93^\circ\text{C}$ (recrystallized from Hexanes). ^1H NMR (600 MHz, C_6D_6) δ : 6.19 (1H, ddd, $J = 17.3, 10.6, 6.7$ Hz), 5.20–5.11 (2H, m), 4.13 (1H, d, $J = 12.3$ Hz), 3.93 (1H, dd, $J = 2.9, 2.9$ Hz), 3.84 (1H, dd, $J = 12.3, 2.9$ Hz), 3.71 (1H, d, $J = 10.6$ Hz), 3.58 (1H, m), 3.00 (1H, m), 2.74 (1H, d, $J = 9.4$ Hz), 2.64 (1H, d, $J = 2.9$ Hz), 2.18 (1H, ddd, $J = 14.7, 2.9, 2.9$ Hz), 1.18 (9H, s), 1.15 (1H, ddd, $J = 14.7, 3.2, 3.2$ Hz), 1.08 (3H, d, $J = 6.5$ Hz), 1.05 (9H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 142.6, 113.5, 85.5, 76.5, 69.7, 68.6, 64.2, 38.6, 36.9, 27.9, 27.5, 23.3, 20.6, 15.0 ppm. FTIR (film): 3503, 2960, 2933, 2858, 1474, 1132, 1091, 1022, 950, 908, 844, 825, 766, 651 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{18}\text{H}_{34}\text{O}_4\text{SiNa}$, 365.2119; found, 365.2116.

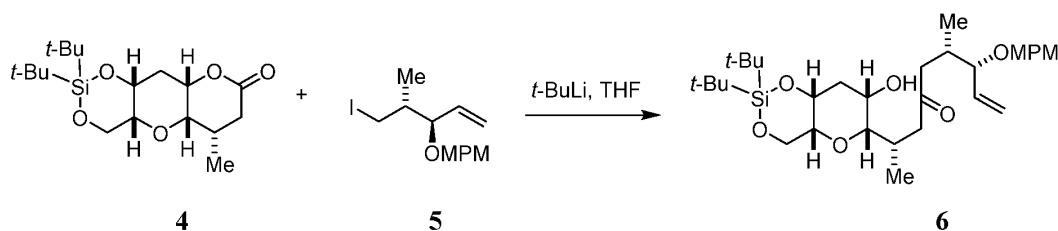


[00786] **S-1** (4.66 g, 13.6 mmol, 1 eq.) was dissolved in 9-BBN solution (68 mL of 0.5 M in THF, 34 mmol, 2.5 eq.) at room temperature. After being stirred for 1.5 h at the same temperature, H_2O (68 mL) and $\text{NaBO}_3 \cdot \text{H}_2\text{O}$ (20.4 g, 204 mmol, 15 eq.) were added. After being stirred for 2 h at the same temperature, the reaction was filtered through a filter paper. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (9%, 25%, then 50% EtOAc in Hexanes) to give **S-2** (4.94 g, 13.7 mmol, quantitative) as a colorless amorphous solid. **S-2**: $[\alpha]_D^{20}$ -7.6 (*c* 1.01, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ 4.33 (1H, m), 4.16 (1H, dd, $J = 12.3, 2.4$ Hz), 4.11 (1H, m), 3.67 (1H, brs), 3.63 (1H, dt, $J = 10.7, 6.4, 6.4$ Hz), 3.59-3.45 (2H, m), 3.27 (1H, m), 2.95 (1H, d, $J = 9.4$ Hz), 2.26 (1H, dt, $J = 14.7, 2.9$ Hz), 2.02 (1H, m), 1.93 (1H, br), 1.70 (1H, dddd, $J = 13.6, 6.6, 6.6, 6.6$ Hz), 1.65 (1H, dt, $J = 14.7, 2.9, 2.9$ Hz), 1.51 (1H, m), 0.95 (9H, s), 0.94 (9H, s), 0.86 (3H, d, $J = 7.0$ Hz). ^{13}C NMR (125 MHz, C_6D_6) δ : 86.3, 76.6, 69.7, 68.5, 64.3, 61.1, 38.3, 36.9, 32.2, 27.9, 27.4, 23.3, 20.5, 16.6 ppm. FTIR (film): 3525, 2933, 2859, 1473, 1145, 1091, 1053, 1023, 954, 869, 839, 826, 764, 651, 447 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{18}\text{H}_{37}\text{O}_5\text{Si}$, 361.2405; found, 361.2415.



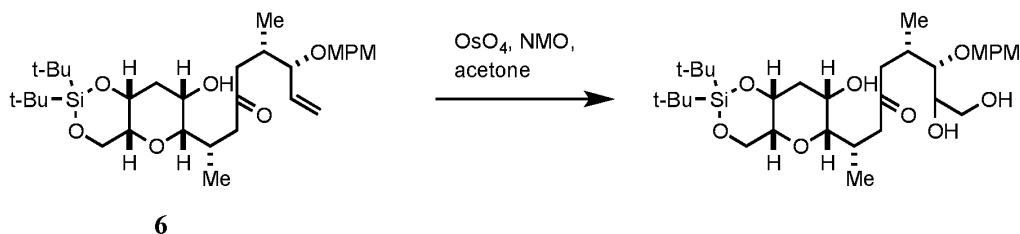
[00787] To a stirred solution of **S-2** (calculated as 13.6 mmol) in CH_2Cl_2 (136 mL) was added NaHCO_3 (11.4 g, 136 mmol, 10 eq.). The mixture was cooled to 4 $^\circ\text{C}$, then PhI(OAc)_2 (13.1 g, 40.8 mmol, 3 eq.) and TEMPO (213 mg, 1.36 mmol, 10 mol%) were added. After being stirred for 15 h at the same temperature, the reaction was quenched with 10% $\text{Na}_2\text{S}_2\text{O}_3$ aq. and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated

under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 9%, 17%, then 33% EtOAc in Hexanes) to give **4** (4.68 g, 13.1 mmol, 97% in 2 steps) as a colorless solid. **4**: $[\alpha]_D^{20}$ -12.7 (*c* 2.29, CHCl₃). MP: 147-150 °C (recrystallized from Hexanes-EtOAc). ¹H NMR (600 MHz, C₆D₆) δ 4.11 (1H, dd, *J* = 12.9, 1.2 Hz), 3.96 (1H, dd, *J* = 12.6, 3.2 Hz), 3.89 (1H, m), 3.43 (1H, m), 2.55 (1H, m), 2.53 (1H, m), 2.43 (1H, dd, *J* = 17.0, 13.5 Hz), 2.28-2.20 (2H, m), 1.29 (1H, m), 1.21 (9H, s), 1.13-1.07 (10H, m), 0.76 (3H, d, *J* = 6.5 Hz). ¹³C NMR (150 MHz, C₆D₆) δ : 169.0, 77.1, 75.0, 73.3, 68.0, 66.5, 35.9, 33.9, 31.3, 28.1, 27.2, 23.4, 20.4, 16.7 ppm. FTIR (film): 2933, 2857, 1727, 1474, 1364, 1180, 1136, 1109, 1045, 976, 829, 770, 645, 443 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₁₈H₃₂NaO₅Si, 379.1911; found, 379.1912.



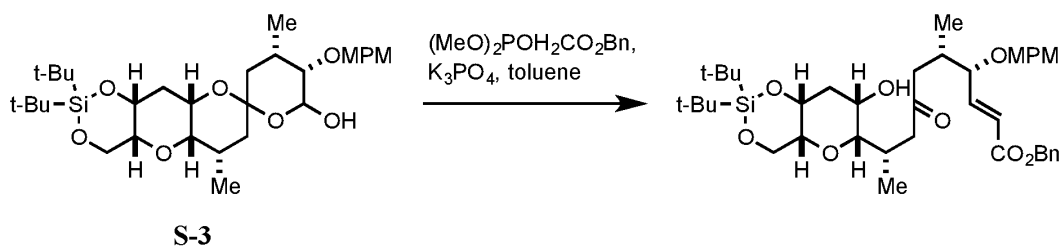
[00788] A solution of **4** (1.0 g, 2.80 mmol) and **5** (1.36 g, 3.93 mmol, 1.4 eq.) in THF was degassed by bubbling with Ar. To the solution was added *t*-BuLi solution (4.23 mL of 1.7 M in pentane, 7.29 mmol, 2.6 eq.) dropwise at -78 °C over 10 min. After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NH₄Cl aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (10% then 30% EtOAc in Hexanes) to give **6** (1.46 g, 2.53 mmol, 90%) as colorless oil. **6** was obtained as an equilibrium mixture of ketone form and ketal form (ca. 4:1 ratio in C₆D₆). Spectral data only for major ketone form are shown here. **6**: $[\alpha]_D^{20}$ -17.4 (*c* 1.44, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.26 (2H, d, *J* = 8.7 Hz), 6.83 (2H, d, *J* = 8.7 Hz), 5.63 (1H, ddd, *J* = 17.1, 10.8, 7.8 Hz), 5.14-5.09 (2H, m), 4.53 (1H, d, *J* = 11.4 Hz), 4.21 (1H, d, *J* = 11.4 Hz), 4.10 (1H, d, *J* = 12.6 Hz), 3.93 (1H, dd, *J* = 2.7, 2.7 Hz), 3.85 (1H, dd, *J* = 12.6, 3.0 Hz), 3.64 (1H, d, *J* = 10.2 Hz), 3.57 (1H, d, *J* = 10.8 Hz), 3.48 (1H, dd, *J* = 7.2, 7.2 Hz), 3.34 (3H, s), 2.81-2.68 (4H, m), 2.64 (1H, s), 2.50-2.46 (1H, m), 2.28 (1H, dd, *J* = 15.3, 7.5 Hz), 2.24 (1H, dd, *J* = 16.5, 8.7 Hz), 2.15 (1H, ddd, *J* = 14.4, 2.7, 2.7 Hz), 1.16 (9H, s), 1.16-1.13 (1H, m), 1.04 (9H, s), 1.02 (3H, d, *J* = 6.6 Hz), 0.97 (3H, d, *J* = 7.2 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ :

209.1, 159.6, 137.8, 131.3, 129.5, 118.1, 114.0, 84.9, 84.2, 76.5, 70.2, 69.7, 68.5, 64.1, 54.8, 47.3, 46.2, 37.0, 34.1, 31.4, 27.9, 27.5, 23.3, 20.5, 16.5, 16.1 ppm. FTIR (film): 3527, 2933, 2858, 1708, 1613, 1513, 1473, 1247, 1146, 1091, 1022, 956, 652, 447 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{32}\text{H}_{52}\text{NaO}_7\text{Si}$, 599.3375; found, 599.3376.



[00789] To a stirred solution of **6** (1.0 g, 1.73 mmol, 1 eq.) in acetone (17.4 mL) were added NMMO (405 mg, 3.46 mmol, 2 eq.) and OsO_4 solution (8.8 mL of 0.02 M in H_2O , 0.173 mmol, 10 mol%) at room temperature. After being stirred for 21 h at the same temperature, the mixture was diluted with water. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was passed through a pad of silica gel (EtOAc) and concentrated under reduced pressure. The obtained crude material was used in the next reaction without further purification. To a stirred solution of diol (calculated as 1.73 mmol) in CH_2Cl_2 (17.3 mL) was added K_2CO_3 (717 mg, 5.19 mmol, 3 eq.) and $\text{Pb}(\text{OAc})_4$ (920 mg, 2.08 mmol, 1.2 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was passed through a pad of silica gel (EtOAc). The filtrate was concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 9%, 17%, then 25% EtOAc in Hexanes) to give **S-3** (826 mg, 1.43 mmol, 83% for 2 steps) as colorless oil. **S-3** was obtained as an equilibrium mixture of hemiacetal (ca. 4:1 ratio in C_6D_6). **S-3**: $[\alpha]_D^{20}$ -39.3 (c 1.01, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.19-7.16 (2H, m), 6.78-6.75 (2H, m), 5.41 (0.2H, dd, 11.1, 1.5 Hz), 4.93 (0.8H, dd, J = 13.2, 1.8 Hz), 4.54 (0.8H, d, J = 10.8 Hz), 4.44 (0.2H, d, J = 11.4 Hz), 4.31 (0.8H, d, J = 10.8 Hz), 4.26 (0.2H, d, J = 11.4 Hz), 4.21-4.17 (1.2H, m), 3.99 (0.8H, dd, J = 12.3, 2.7 Hz), 3.98 (0.2H, dd, J = 12.6, 3.6 Hz), 3.90-3.89 (0.8H, m), 3.87-3.86 (0.2H, m), 3.69 (0.2H, d, J = 10.8 Hz), 3.62 (0.8H, d, J = 4.2 Hz), 3.36 (0.8H, d, J = 12.6 Hz), 3.62 (0.8H, d, 4.2 Hz), 3.36 (0.8H, d, J = 12.6 Hz), 3.29 (0.6H, s), 3.28 (2.4H, s), 3.19 (0.8H, s), 2.81 (0.2H, d, J = 3.0 Hz), 2.77 (0.8H, d, J = 2.4 Hz), 2.63 (0.8H, dd, J = 1.5, 1.5 Hz), 2.61 (0.2H, dd, J = 1.8, 1.8 Hz), 2.47-2.42 (0.2H, m), 2.34-2.25 (1.8H, m), 2.09 (0.8H, ddd, J = 15.0, 1.8, 1.8 Hz), 2.00 (0.2H, ddd, J = 15.0, 1.8, 1.8 Hz), 1.91-1.88 (0.4H, m), 1.74-1.61 (2.8H, m), 1.53-1.49

(1H, m), 1.42 (0.2H, ddd, $J = 15.6, 4.2, 4.2$ Hz), 1.36 (0.8H, ddd, $J = 15.0, 4.5, 4.5$ Hz), 1.29 (7.2H, s), 1.27 (1.8H, s), 1.14 (7.2H, s), 1.13 (1.8H, s), 1.07 (0.6H, d, $J = 6.6$ Hz), 1.03 (2.4H, d, $J = 7.2$ Hz), 1.01 (2.4H, d, $J = 7.2$ Hz), 1.00 (0.6H, d, $J = 6.6$ Hz) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 159.9, 159.7, 131.2, 129.6, 129.5, 128.5, 114.3, 114.0, 99.0, 98.4, 93.9, 91.2, 80.2, 78.6, 77.5, 77.4, 76.0, 72.2, 68.6, 68.5, 67.2, 67.0, 64.7, 64.0, 54.7, 38.5, 37.5, 37.4, 36.9, 36.4, 36.3, 29.5, 29.0, 27.8, 27.6, 23.41, 23.38, 23.1, 20.9, 17.9, 17.5, 17.2 ppm. FTIR (film): 3512, 2931, 2856, 1612, 1514, 1474, 1246, 1195, 1127, 1094, 1043, 1014, 966, 942, 828, 769, 735, 651, 441 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{31}\text{H}_{50}\text{NaO}_8\text{Si}$, 601.3167; found, 601.3168.



[00790] To a stirred solution of **S-3** (208 mg, 0.359 mmol, 1 eq.) in toluene (3.6 mL) were added benzyl dimethylphosphonoacetate (0.30 mL, 1.44 mmol, 4 eq.) and K_3PO_4 (232 mg, 1.08 mmol, 3 eq.) at room temperature. After being stirred for 23 h at the same temperature, the reaction mixture was passed through a pad of silica gel (50% EtOAc in Hexanes) to give crude unsaturated ester (**7**) (as a ~3:1 mixture of *E/Z* isomers). The crude material was used in the next reaction without further purification.

[00791] To a stirred solution of the crude **7** (calculated as 0.359 mmol, 1 eq.), BnOAc (541 mg, 3.6 mmol, 10 eq.), and LiBr (313 mg, 3.6 mmol, 10 eq.) in MeCN (3.6 mL) was added DBU (0.269 μL , 1.80 mmol, 5 eq.) at room temperature. After being stirred for 12 h at the same temperature, the reaction was quenched with sat. NH_4Cl aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude spiro acetal, which was used in the next reaction without further purification. To a stirred solution of the crude spiro acetal (calculated as 0.359 mmol) in CH_2Cl_2 (3.6 mL) and phosphate buffer (pH=7, 0.7 mL) was added DDQ (163 mg, 0.72 mmol, 2 eq.). After being stirred for 40 min, the reaction was quenched with 10% $\text{Na}_2\text{S}_2\text{O}_3$ aq. and sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced

pressure. The residue was purified by flash column chromatography on silica gel (0%, 9%, 13%, the 17% EtOAc in Hexanes) to give alcohol **S-4** (160 mg, 0.271 mmol, 75% for 3 steps) as colorless oil. **S-4**: $[\alpha]_D^{20}$ -51.3 (*c* 1.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.24-7.01 (5H, m), 5.07 (1H, d, *J* = 12.3 Hz), 4.98 (1H, d, *J* = 12.3 Hz), 4.30 (1H, m), 4.22 (1H, d, *J* = 12.9 Hz), 4.16 (1H, dd, *J* = 10.0, 2.9 Hz), 4.00 (1H, dd, *J* = 12.6, 2.6 Hz), 3.93 (1H, m), 3.87 (1H, m), 2.99 (1H, d, *J* = 8.2 Hz), 2.92-2.84 (2H, m), 2.71 (1H, brs), 2.32 (1H, dd, *J* = 16.1, 3.2 Hz), 2.28 (1H, d, *J* = 15.3 Hz), 2.18 (1H, m), 2.11 (1H, m), 1.74 (1H, dd, *J* = 13.5, 4.1 Hz), 1.66 (1H, dd, *J* = 12.9, 12.9 Hz), 1.55 (1H, ddd, *J* = 15.4, 4.5, 4.5 Hz), 1.36-1.20 (11H, m), 1.13 (9H, s), 1.05 (3H, d, *J* = 7.0 Hz), 0.92 (3H, d, *J* = 6.5 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 171.6, 136.7, 128.7, 128.5, 126.9, 92.3, 77.52, 77.48, 70.5, 69.4, 68.6, 67.2, 66.1, 63.7, 37.42, 37.38, 36.5, 36.4, 29.8, 29.1, 27.8, 27.6, 23.4, 20.9, 17.6, 17.4 ppm. IR (film): 3473, 2957, 2931, 2857, 1736, 1131, 1017, 974 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₃₂H₅₀NaO₈Si, 613.3167; found, 613.3169.



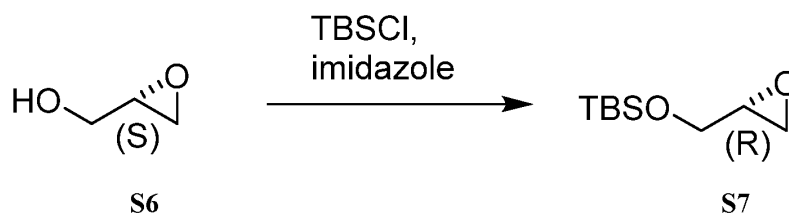
[00792] To a stirred solution of **S-4** (141 mg, 0.239 mmol, 1 eq.) in CH₂Cl₂ (2.4 mL, 0.1 M) were added imidazole (65 mg, 0.96 mmol, 4 eq.) and TESCl (80.2 μ L, 0.48 mmol, 2 eq.) at room temperature. After being stirred for 16 h at the same temperature, the reaction was quenched with MeOH. The mixture was concentrated and passed through a pad of silica gel (25% EtOAc in Hexanes), and concentrated under reduced pressure to give a crude TES ether, which was used in the next reaction without further purification.

[00793] To a stirred solution of the crude TES ether (calculated as 0.239 mmol) in EtOAc (3.5 mL) was added wet 10% Pd/C (15 mg). The reaction was stirred under 1 atmosphere of hydrogen for 45 min at room temperature. The mixture was degassed and filled with N₂, passed through a pad of silica gel (EtOAc), and concentrated under reduced pressure to give a crude acid, which was used in the next reaction without further purification.

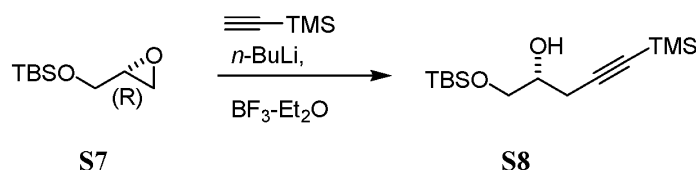
[00794] To a stirred solution of the crude acid (calculated as 0.239 mmol) in CH₂Cl₂ (2.4 mL, 0.1 M) were added PPh₃ (94 mg, 0.389 mmol, 1.2 eq.) and (PyS)₂ (73.7 mg, 0.335 mmol, 1.4 eq.) at room temperature. After being stirred for 17 h at the same temperature, the reaction mixture was concentrated under reduced pressure. The residue was purified by flash column

chromatography on neutral silica gel (0%, 9%, then 17% EtOAc in Hexanes) to give thioester **8** (162 mg, 0.229 mmol, 96% for 3 steps), which contained 2% of (PyS)₂. The disulfide impurity can be removed by additional column chromatography (0%, 1%, 2%, 3%, then 9% EtOAc in CH₂Cl₂) or HPLC (Column: DuPont Instruments ZORBAL SIL 21.2 mm x 25 cm (880952-101), Solvent: 3% *i*PrOH in Hexanes, Flow rate: 10.0 mL/min, Detection: UV at 254 nm and 220 nm, *t_R* = 15 min. The product was obtained as pale yellow foam. **8**: $[\alpha]_D^{20}$ -74.3 (*c* 1.0, CHCl₃). ¹H NMR (500 MHz, C₆D₆) δ : 8.33 (1H, m), 7.52 (1H, m), 6.94 (1H, m), 6.47 (1H, m), 4.25-4.16 (2H, m), 3.99 (1H, dd, *J* = 12.4, 2.7 Hz), 3.97-3.91 (2H, m), 3.27 (1H, m), 3.19 (1H, brs), 2.91 (1H, m), 2.72 (1H, brs), 2.59 (1H, dd, *J* = 14.9, 2.2 Hz), 2.34 (2H, d, *J* = 14.6 Hz), 2.24 (1H, m), 1.77-1.60 (4H, m), 1.52 (1H, m), 1.31 (9H, s), 1.13 (9H, s), 1.04-0.96 (12H, m), 0.93 (3H, m), 0.59 (6H, q, *J* = 7.8 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 194.7, 152.7, 150.6, 136.5, 128.5, 123.1, 97.4, 77.7, 77.5, 72.5, 70.0, 68.6, 67.3, 63.9, 47.8, 37.5, 36.7, 36.5, 30.5, 29.1, 27.8, 27.7, 23.4, 21.0, 18.5, 17.3, 7.4, 5.9 ppm. IR (film): 2955, 2931, 2874, 2857, 1708, 1132, 1035, 974 cm⁻¹. HRMS (ESI) *m/z*: [M+H]⁺ calcd for C₃₆H₆₂O₇SSi₂, 708.3780; found, 708.3779.

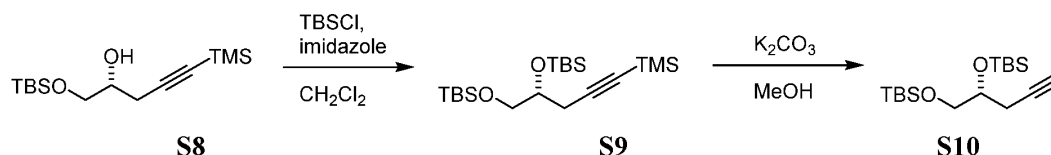
Halichondrin Left Halves



[00795] To a solution of (*S*)-glycidol **S6** (6.70 g, 90.0 mmol; AK Scientific) in dichloromethane (250 mL) was added imidazole (7.40 g, 108 mmol) and TBSCl (14.9 g, 99.0 mmol) at 0 °C. The reaction was stirred for 6 hr at room temperature before quenched with water (200 mL), and the resulting two layers were separated. The organic layer was dried over anhydrous Na₂SO₄, and filtered, concentrated under reduced pressure to give a crude oil. Purification over SiO₂ (hexanes/ethyl acetate = 50/1) gave **S7** as a clear oil (16.2 g, 95%). **S7**: $[\alpha]_D^{22}$ = -2.2 (*c* 1.0, CHCl₃); ¹H NMR (500 MHz, CDCl₃) δ : 3.85 (1H, dd, *J* = 9.6, 3.2 Hz), 3.66 (1H, dd, *J* = 11.9, 5.0 Hz), 3.09 (1H, ddd, *J* = 7.5, 4.4, 3.0 Hz), 2.77 (1H, dd, *J* = 5.1, 4.5 Hz), 2.64 (1H, dd, *J* = 5.2, 2.6 Hz), 0.9 (9H, s), 0.09 (3H, s), 0.08 (3H, s) ppm. ¹³C NMR (125 MHz, CDCl₃) δ : 63.7, 52.4, 44.4, 25.8, 18.3, -5.3, -5.4 ppm. HRMS (ESI) *m/z*: [M+H]⁺ calcd for C₉H₂₁O₂Si 189.1305; found 189.1299.



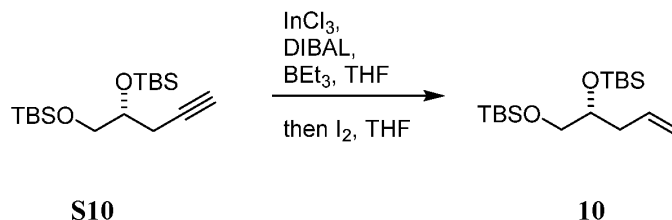
[00796] A solution of trimethylsilyl acetylene (23.8 mL, 167 mmol; Oakwood) in dry THF (76.0 mL) was cooled to -78°C under Ar atmosphere and treated with *n*-BuLi (2.50 M in hexane, 61.3 mL, 159 mmol). After 30 min, $\text{BF}_3 \cdot \text{OEt}_2$ (18.9 mL, 159 mmol) was added dropwise followed by slow addition of a solution of epoxide **S7** (15.0 g, 79.7 mmol) in THF (30.0 mL) at -78°C . The reaction mixture was stirred for 1 h at the same temperature before addition of *sat'd*- NaHCO_3 (100 mL) and diethyl ether (300 mL). The resulting biphasic solution was warmed up to rt, and the layers were separated. The organic layer was dried over anhydrous Na_2SO_4 , filtered, and concentrated under reduced pressure to give crude oil. Purification over SiO_2 (hexanes/ethyl acetate = 10/1) gave **S8** as a clear oil (20.5 g, 90%). **S8**: $[\alpha]_D^{22} = -15.6$ (*c* 1.0, CHCl_3); ^1H NMR (500 MHz, CDCl_3): δ 3.82-3.78 (1H, m), 3.73 (1H, dd, $J = 10.0, 4.5$ Hz), 3.67 (1H, dd, $J = 9.5, 5.5$ Hz), 2.52-2.44 (2H, m), 0.92 (9H, s), 0.16 (9H, s), 0.10 (6H, s) ppm. ^{13}C NMR (125 MHz, CDCl_3): δ 102.8, 87.0, 70.1, 65.4, 25.9, 24.5, 18.3, 0.02, -5.4, -5.4 ppm. HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{14}\text{H}_{30}\text{NaO}_2\text{Si}_2$ 309.1677; found 309.1677.



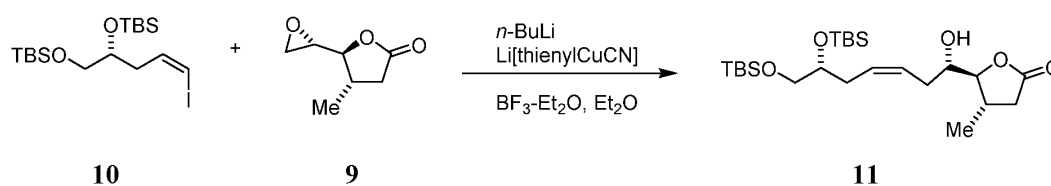
[00797] To a solution of alcohol **S8** (20.5 g, 71.5 mmol) and imidazole (7.3 g, 107 mmol) in dichloromethane (300 mL) was added TBSCl (14.0 g, 92.9 mmol) at 0°C . The reaction was stirred for 6 hr at rt before addition of water (100 mL). The mixture was extracted with hexanes (200 mL), and the organic layer was dried over anhydrous Na_2SO_4 , filtered, and concentrated under reduced pressure to give the crude product **S9** which was used in next step without further purification.

[00798] To a solution of **S9** in methanol (240 mL) was treated with potassium carbonate (11.9 g, 85.8 mmol) at rt. After 6 h, the reaction was diluted with hexanes (200 mL) and filtered through a pad of Celite[®]. Concentration under reduced pressure gave a crude oil, which was subject to purification over SiO_2 (hexanes/ethyl acetate = 50/1) to give **S10** as a clear oil (21.2 g, 90% for 2 steps). **S10**: $[\alpha]_D^{22} = +6.0$ (*c* 1.0, CHCl_3). ^1H NMR (500 MHz, CDCl_3): δ : 3.81 (1H, quint, $J = 5.5$ Hz), 3.53-3.52 (2H, m), 2.46 (1H, ddd, $J = 14.0, 5.5, 2.5$

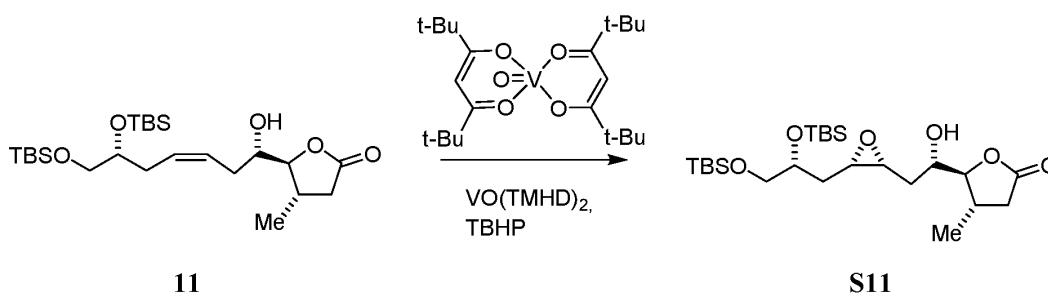
(Hz), 2.29 (1H, ddd, $J = 14.0, 6.0, 2.5$ Hz), 1.94 (1H, t, $J = 2.5$), 0.89 (18H, s), 0.12 (3H, s), 0.08 (3H, s), 0.06 (3H, s), 0.05 (3H, s). ^{13}C NMR (125 MHz, CDCl_3) δ : 81.7, 71.8, 69.6, 66.4, 25.9, 25.8, 24.3, 18.3, 18.1, -4.5, -4.7, -5.4, -5.4 ppm. FTIR (film): 2955, 2929, 2886, 2857, 2124, 1472, 1361, 1253, 1116, 1078, 832, 773, 637 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{17}\text{H}_{37}\text{O}_2\text{Si}_2$ 329.2327; found 329.2328.



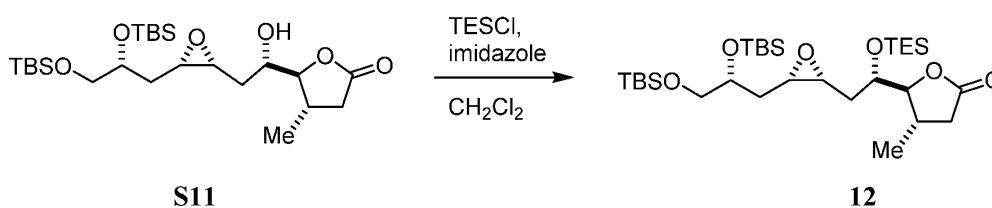
[00799] Anhydrous indium trichloride (16.4 g, 74.0 mmol; Alfa Aesar 99.99%) was placed in flask and heated with hot gun *in vacuo* for 3 min.³ The indium salt was dissolved with THF (220 mL) at 0 °C under an argon atmosphere. The solution turned to a white suspension upon cooling to -78 °C. DIBAL (1.0 M in hexane, 71.3 mL, 71.3 mmol) was then added dropwise to suspension at -78 °C. The mixture was stirred for 2.5 hr to prepare dichloroindium hydride. **S10** (18.1 g, 54.8 mmol) and triethylborane (1.0 M hexane solution, 11.0 mL, 11.0 mmol; Aldrich) were added to the reaction mixture and the resulting mixture was stirred for 4.5 hr at -78 °C. Iodine (41.8 g, 164 mmol) in THF (80.0 mL) was added to the reaction mixture. After being stirred for 20 min at -78 °C, the reaction was poured into saturated NaHCO_3 solution. $\text{Na}_2\text{S}_2\text{O}_3$ solution was added to consume excess iodine. Then saturated potassium sodium tartrate solution was added and stirring vigorously for 1 hr. The mixture was extracted with hexane and ethyl acetate (4:1, 1000 mL) twice. The organic layer was dried over anhydrous Na_2SO_4 , filtered, and concentrated under reduced pressure to give crude oil. Purification over SiO_2 (hexanes/ethyl acetate = 100/1) gave **10** as a clear oil (21.3 g, 85%, $Z/E > 99:1$)². **10**: colorless oil. $[\alpha]_D^{22} = +3.0$ (c 0.9, CHCl_3). ^1H NMR (500 MHz, CDCl_3) δ : 6.29-6.24 (2H, m), 3.77 (1H, quint, $J = 5.4$ Hz), 3.51 (1H, dd, $J = 10.2, 6.6$ Hz), 3.40 (1H, dd, $J = 10.2, 6.6$ Hz), 0.88 (9H, s), 0.86 (9H, s), 0.05 (3H, s), 0.04 (3H, s), 0.04 (3H, s), 0.03 (3H, s) ppm. ^{13}C NMR (125 MHz, CDCl_3) δ : 138.0, 83.7, 71.9, 67.0, 39.8, 26.0, 25.9, 18.4, 18.1, -4.4, -4.7, -5.3, -5.3 ppm. FTIP (film): 2954, 2928, 2885, 2857, 1610, 1471, 1389, 1306, 1113, 1081, 831, 772 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{17}\text{H}_{38}\text{IO}_2\text{Si}_2$ 457.1450; found 457.1455.



[00800] To a solution of vinyl iodide **10** (5.79 g, 12.7 mmol, 1.8 equiv.) in anhydrous Et₂O (64 mL) was added *n*-BuLi (2.52 M in hexane, 4.8 mL, 1.75 equiv.; Aldrich) dropwise at -78°C under Ar atmosphere, and the clear mixture was stirred for 1 h at the same temperature. Li 2-thienylcyanocuprate (0.25 M in THF, 53.4 mL, 2.0 equiv.; Aldrich) was slowly added over 10 min at -78°C , which was stirred for 30 min before addition of BF₃·Et₂O (0.87 mL, 7.04 mmol, 1.6 equiv.). After 20 min, a solution of epoxide **9** (1000 mg, 7.04 mmol, 1.0 equiv.) in anhydrous Et₂O (5.0 mL) was added dropwise at -78°C , and the yellow reaction mixture was stirred for 1 hr at the same temperature. The reaction was quenched by slow addition of a mixture of sat. NH₄Cl aq. (90 mL) and 30% aq. NH₄OH (10 mL) at 0°C and stirred 2 hr at rt. The biphasic mixture was diluted with Et₂O (200 mL), and layers were separated, and organic layer was dried over Na₂SO₄ and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (25% EtOAc in Hexanes) to afford **11** as a oil (2.69 g, 81%). **11**: $[\alpha]_{\text{D}}^{22} = +13.0$ (c 1.0, CDCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 5.74 (1H, dd, *J* = 10.8, 8.4 Hz), 5.47 (1H, dd, *J* = 10.8, 7.8 Hz), 3.83–3.79 (m, 1H), 3.63 (1H, dd, *J* = 9.9, 5.1 Hz), 3.56 (1H, dd, *J* = 10.2, 6.6 Hz), 3.45 (1H, dd, *J* = 6.0, 2.4 Hz), 3.25–3.24 (1H, m), 2.50–2.35 (3H, m), 2.37 (1H, dd, *J* = 16.8, 9.0 Hz), 2.23–2.14 (2H, m), 1.95 (1H, d, *J* = 6.0 Hz), 1.59 (1H, dd, *J* = 17.1, 8.1 Hz), 1.00 (9H, s), 0.99 (9H, s), 0.53 (3H, d, *J* = 7.2 Hz), 0.13 (3H, s), 0.12 (3H, s), 0.10 (3H, s), 0.09 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 175.5, 129.6, 127.1, 87.8, 73.3, 70.9, 67.3, 36.8, 32.8, 32.5, 31.3, 26.1 (×3), 26.2 (×3), 18.6, 18.4, 18.1, -4.2, -4.4, -5.1, -5.2 ppm. FTIP (film): 3450, 2958, 2930, 2858, 1778, 1472, 1428, 1389, 1361, 1252, 1113, 835, 776, 738, 703 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₄H₄₈O₅NaSi₂⁺ 495.2932; found 495.2940.

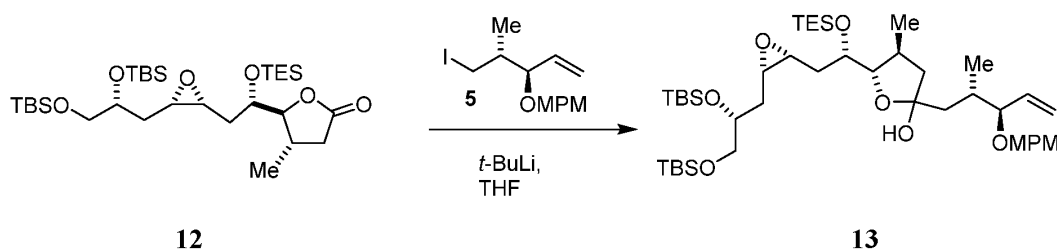


[00801] To a solution of alkene **11** (490 mg, 1.04 mmol, 1.0 equiv.) in toluene (10 mL) was added VO(TMHD)₂ (23 mg, 5 mol%) and *tert*-butylhydrogenperoxide (TBHP) (5.5 M in decane, 380 μ L, 2.0 equiv.; Aldrich) to form a reddish solution. The reaction mixture was stirred for 5.5 hr at rt before quenched with sat. Na₂S₂O₃/NaHCO₃ solution (v:v = 1:1). The resulting biphasic mixture was diluted with Et₂O (10 mL), and the layers were separated. The organic layer was dried over anhydrous Na₂SO₄, concentrated, and purified by flash column chromatography on silica gel (33% EtOAc in Hexanes) to afford **S11** as a colorless oil (450 g, 89 %, *dr* > 50:1). **S11**: [α]_D²² = + 26.0 (c 1.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 3.92 (1H, quint, *J* = 5.4 Hz), 3.70 (1H, dd, *J* = 10.2, 5.1 Hz), 3.66-3.62 (2H, m), 3.76 (1H, dd, *J* = 9.9, 6.3 Hz), 3.14-3.11 (2H, m), 2.97 (1H, dt, *J* = 14.4, 4.2 Hz), 2.47 (1H, dd, *J* = 18.0, 9.6 Hz), 2.31-2.24 (1H, m), 1.84 (1H, dt, *J* = 14.4, 4.8 Hz), 1.77-1.62 (4H, m), 0.97 (9H, s), 0.96 (9H, s), 0.67 (3H, d, *J* = 6.6 Hz), 0.12 (3H, s), 0.11 (3H, s), 0.08 (3H, s), 0.07 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 176.1, 88.3, 72.2, 70.5, 67.5, 53.7, 53.4, 36.8, 33.2, 31.9, 31.2, 26.2 ($\times$ 3), 26.1 ($\times$ 3), 18.6, 18.5, 18.4, -4.2, -4.6, -5.2, -5.2 ppm. FTIP (film): 3450, 2955, 2929, 2857, 1779, 1472, 1463, 1388, 1361, 1253, 1115, 835, 776, 738 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₄H₄₈O₆NaSi₂⁺ 511.2882; found 511.2877.

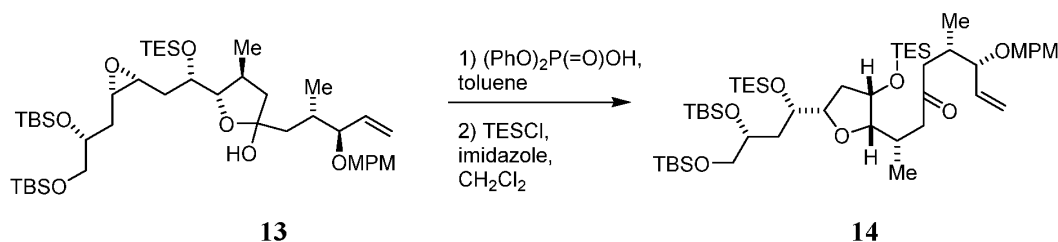


[00802] To a solution of **S11** (450 mg, 0.92 mmol, 1.0 equiv.) and imidazole (251 mg) in dichloromethane (9.2 mL) was added TESCl (308 μ L) at 0 °C. The reaction was stirred for 1 hr at 0 °C before quenched with sat. NaHCO₃. The biphasic mixture was diluted with dichloromethane, and the layers were separated. The organic layer was dried over anhydrous Na₂SO₄, filtered, and concentrated under reduced pressure to give a crude oil. Purification over SiO₂ (9% EtOAc in Hexanes) afforded **12** as a clear oil (528 mg, 95%). **12**: [α]_D²² = +

26.9 (c 1.0, CHCl₃). ¹H NMR (500 MHz, C₆D₆) δ: 3.97 (1H, quint, *J* = 5.4 Hz), 3.94 (1H, dd, *J* = 5.0, 3.5 Hz), 3.89-3.85 (1H, m), 3.76 (1H, dd, *J* = 10.0, 5.5 Hz), 3.65 (1H, dd, *J* = 10.5, 6.0 Hz), 3.15 (1H, dt, *J* = 7.5, 4.0 Hz), 2.92 (1H, dt, *J* = 9.0, 4.0 Hz), 2.40 (1H, dd, *J* = 17.2, 9.2 Hz), 2.13-2.09 (1H, m), 2.04 (1H, ddd, *J* = 14.0, 7.5, 3.5 Hz), 1.95 (1H, ddd, *J* = 14.5, 5.5, 4.5 Hz), 1.71-1.64 (2H, m), 1.59 (1H, ddd, *J* = 13.5, 8.0, 4.5 Hz), 1.00 (9H, s), 0.98 (9H, s), 0.97 (9H, t, *J* = 8.5 Hz), 0.68 (3H, d, *J* = 6.5 Hz), 0.58 (6H, q, *J* = 8.0 Hz), 0.14 (3H, s), 0.13 (3H, s), 0.11 (3H, s), 0.09 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ: 175.3, 86.9, 72.2, 71.7, 67.7, 53.7, 52.4, 36.8, 33.5, 32.4, 31.2, 26.2 (×3), 26.1 (×3), 19.2, 18.6, 18.3, 7.1, 7.0, 6.3, 5.3, -4.2, -4.7, -5.2, -5.2 ppm. FTIP (film): 2955, 2929, 2878, 2857, 1781, 1472, 1463, 1388, 1361, 1250, 1097, 1005, 832, 774, 735 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₃₀H₆₂O₆NaSi₃⁺ 625.3746; found 625.3748.

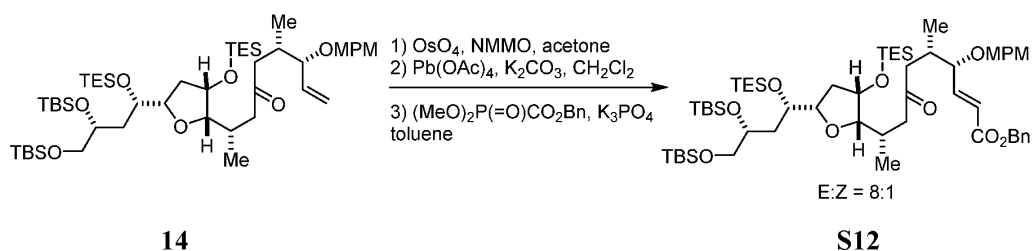


[00803] A solution of **12** (193 mg, 0.32 mmol) and **5** (155 mg, 0.45 mmol, 1.4 eq.) in THF was degassed by bubbling with Ar. To the solution was added *t*-BuLi solution (0.49 mL of 1.7 M in pentane, 14.6 mmol, 2.6 eq.) dropwise at -78 °C over 10 min. After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NH₄Cl aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with brine, dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (10% then 30% EtOAc in Hexanes) to give **13** (225 mg, 0.27 mmol, 85%) as colorless oil. **13** was obtained as an equilibrium mixture of ketone form and ketal form (ca. 1:1 ratio).



[00804] To a solution of **13** (500 mg, 0.607 mmol) in 10.1 mL of toluene, (PhO)₂P(=O)OH (15.2 mg in 2 mL toluene) was added at 0 °C. The resulting mixture was stirred at room temperature for 12 hr before it was quenched by 5 mL of saturated aqueous NaHCO₃. The

organic layer was removed, and the aqueous layer was extracted with ethyl acetate (30 mL) three times. The combined organic layers were dried over Na_2SO_4 , filtered and concentrated. The resulting oil was then dissolved in 6 mL of dichloromethane before the addition of imidazole (413 mg, 6.07 mmol) and TESCl (509 μL , 3.04 mmol) at 0 °C. The resulting mixture was stirred at room temperature for 2 hr before it was quenched by 5 mL of saturated aqueous NH_4Cl . The organic layer was removed, and the aqueous layer was extracted with ethyl acetate (30 mL) three times. The combined organic layers were dried over MgSO_4 , filtered and concentrated. The residue was purified by flash column chromatography on silica gel (9% EtOAc in Hexanes) afforded **14** as a clear oil (484 mg, 85% for 2 steps). **14**: $[\alpha]_D^{22} = -5.8$ (c 1.1, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.24 (1H, d, $J = 8.4$ Hz), 6.82 (1H, d, $J = 9.0$ Hz), 5.63 (1H, ddd, $J = 17.4, 10.8, 7.8$ Hz), 5.13 (1H, dd, $J = 8.4, 3.0$ Hz), 5.10 (1H, d, $J = 10.2$ Hz), 4.54 (1H, d, $J = 10.8$ Hz), 4.27-4.24 (1H, m), 4.21 (1H, d, $J = 7.6$ Hz), 4.06-4.03 (1H, m), 4.00-3.98 (1H, m), 3.83 (1H, dd, $J = 10.5, 3.3$ Hz), 3.76 (1H, q, $J = 4.8$ Hz), 3.72 (1H, dd, $J = 10.2, 6.0$ Hz), 3.49 (1H, t, $J = 6.9$ Hz), 3.33 (3H, s), 3.12 (1H, dd, $J = 8.4, 4.2$ Hz), 3.04 (1H, dd, $J = 16.8, 3.0$ Hz), 2.77-2.71 (2H, m), 2.59-2.53 (1H, m), 2.36 (1H, dd, $J = 16.2, 10.2$ Hz), 2.31 (1H, dd, $J = 16.8, 9.0$ Hz), 2.00-1.91 (2H, m), 1.76 (1H, ddd, $J = 12.6, 7.8, 3.6$ Hz), 1.59 (1H, ddd, $J = 12.6, 7.2, 2.4$ Hz), 1.14 (9H, t, $J = 7.8$ Hz), 1.08 (9H, s), 1.03 (9H, s), 1.01 (3H, d, $J = 5.4$ Hz), 0.99 (1H, t, $J = 8.4$ Hz), 0.82 (3H, q, $J = 7.8$ Hz), 0.81 (3H, q, $J = 7.8$ Hz), 0.57 (6H, q, $J = 7.8$ Hz), 0.28 (3H, s), 0.28 (3H, s), 0.15 (6H, s) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 208.4, 159.7, 137.8, 131.4, 129.5, 118.1, 114.0, 87.4, 84.1, 81.3, 72.7, 72.2, 71.3, 70.3, 68.2, 54.8, 47.6, 46.4, 39.1, 39.0, 33.9, 28.9, 26.3 ($\times 6$), 18.7, 18.5, 17.4, 16.6, 7.9, 7.2, 5.8, 5.3, -3.9, -4.2, -5.0, -5.1 ppm. FTIP (film): 2956, 2932, 2880, 2855, 1713, 1615, 1514, 1463, 1381, 1361, 1249, 1079, 1039, 944, 833, 775, 736 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{50}\text{H}_{96}\text{O}_8\text{NaSi}_4^+$ 959.6074; found 959.6070.

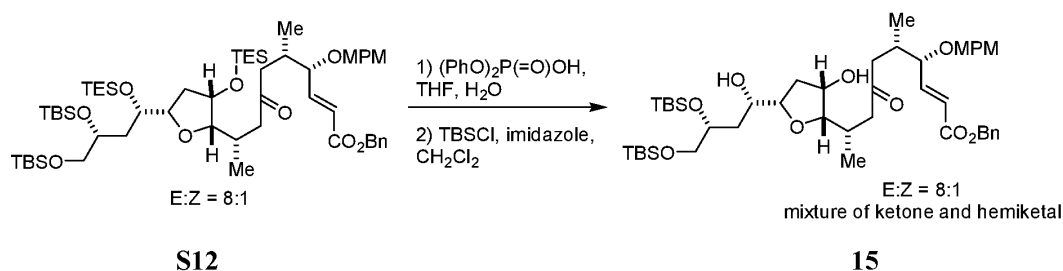


[00805] To a solution of **14** (1.80 g, 1.92 mmol, 1.0 equiv.) in acetone (19.2 mL) was added NMMO (449 mg, 3.84 mmol, 2.0 equiv.) and aqueous solution of OsO_4 (4.88 mL, 0.096 mmol, 5 mg/mL H_2O , 5 mol%). The reaction mixture was stirred for 15 hr, and then

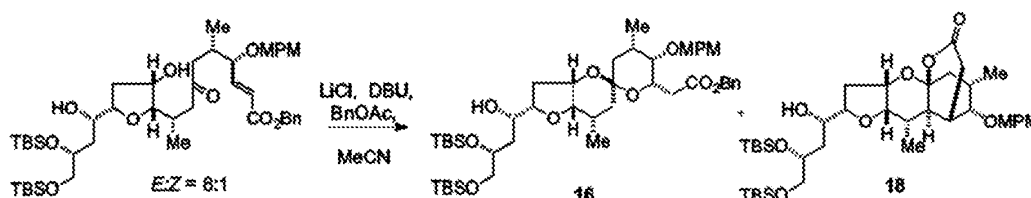
quenched with Na₂SO₃ aq.. The mixture was extracted twice with EtOAc, dried over Na₂SO₄, filtered, and concentrated. The residue was passed through a pad of SiO₂ with EtOAc and concentrated. Without further purification the crude material was used in the next reaction.

[00806] To a solution of diol (estimated as 1.92 mmol, 1.0 equiv.) in DCM (19.2 mL) was added K₂CO₃ (2.66 g, 19.2 mmol, 10 equiv.) and Pb(OAc)₄ (1.36 g, 3.07 mmol, 1.6 equiv.). After being stirred for 1 hr, the reaction mixture was passed through SiO₂ pad with EtOAc and concentrated. Without further purification the crude material was used in the next reaction.

[00807] To a solution of aldehyde (estimated as 1.92 mmol, 1.0 equiv.) in toluene (19.2 mL) at room temperature was added (MeO)₂P(=O)CH₂CO₂Bn (2.0 mL, 9.6 mmol, 5.0 equiv.) and K₃PO₄ (4.08 g, 19.2 mmol, 10 equiv.). After being stirred for additional 12 h, the reaction mixture was passed through a pad of SiO₂ with EtOAc/hexanes (1/1), and concentrated to give **S12** (1.65 g, 1.51 mmol, 82% for 3 steps, ~8:1 mixture of *E/Z* isomers). **S12**: $[\alpha]_D^{22} = -7.0$ (c 0.9, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.24 (2H, d, *J* = 7.2 Hz), 7.15 (2H, d, *J* = 8.4 Hz), 7.10 (2H, dd, *J* = 7.2, 7.2 Hz), 7.05 (1H, t, *J* = 7.2 Hz), 7.00 (1H, dd, *J* = 15.5, 6.3 Hz), 6.59 (2H, d, *J* = 8.4 Hz), 6.13 (1H, d, *J* = 15.5 Hz), 5.10 (1H, d, *J* = 12.0 Hz), 5.06 (1H, d, *J* = 12.0 Hz), 4.30 (1H, d, *J* = 12.0 Hz), 4.27-4.24 (1H, m), 4.03 (1H, d, *J* = 12.0 Hz), 4.00-3.97 (1H, m), 3.97-3.94 (1H, m), 3.78 (1H, dd, *J* = 10.6, 3.8 Hz), 3.71 (1H, q, *J* = 7.8 Hz), 3.65 (1H, dd, *J* = 16.8, 6.6 Hz), 3.55 (1H, t, *J* = 6.9 Hz), 3.27 (3H, s), 3.05 (1H, dd, *J* = 12.6, 4.2 Hz), 2.93 (1H, dd, *J* = 16.8, 3.0 Hz), 2.66-2.61 (1H, m), 2.50-2.44 (2H, m), 2.21 (1H, dd, *J* = 8.4, 8.4 Hz), 2.19 (1H, dd, *J* = 8.4, 8.4 Hz), 2.00-1.88 (2H, m), 1.70 (1H, ddd, *J* = 12.6, 7.8, 3.6 Hz), 1.55 (1H, ddd, *J* = 12.6, 7.2, 2.4 Hz), 1.08 (9H, t, *J* = 7.8 Hz), 1.03 (9H, s), 0.98 (9H, s), 0.96 (3H, d, *J* = 5.4 Hz), 0.93 (1H, t, *J* = 8.4 Hz), 0.88 (3H, q, *J* = 7.8 Hz), 0.76 (3H, q, *J* = 7.8 Hz), 0.52 (6H, q, *J* = 7.8 Hz), 0.23 (6H, s), 0.20 (6H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 207.8, 165.5, 159.8, 147.7, 136.6, 130.6, 129.6, 128.69, 128.66, 128.3, 123.3, 114.0, 87.3, 81.6, 81.2, 72.6, 72.1, 71.2, 68.2, 66.4, 54.7, 47.6, 45.7, 39.0, 38.96, 33.5, 28.8, 26.2 ($\times 6$), 18.6, 18.4, 17.2, 16.5, 7.4, 7.2, 5.8, 5.2, -4.0, -4.3, -5.1, -5.2 ppm. FTIP (film): 2954, 2929, 2876, 2856, 1720, 1655, 1612, 1514, 1462, 1381, 1301, 1249, 1158, 1079, 1005, 835, 776, 740 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₅₀H₉₆O₈NaSi₄⁺ 959.6074; found 959.6070.

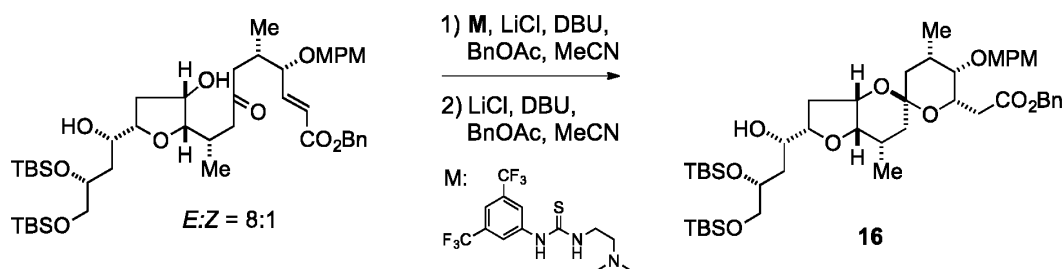


[00808] To a THF-H₂O (4:1, 0.05 M) solution of **S12** (1.65 g, 1.51 mmol, 1 eq.), (PhO)₂P(=O)OH (113 mg, 0.45 mmol, 0.3 eq.) was added at 0 °C. The resulting mixture was stirred at room temperature for 24 hr before it was quenched by 5 mL of saturated aqueous NaHCO₃. The organic layer was removed, and the aqueous layer was extracted with ethyl acetate (30 mL) three times. The combined organic layers were dried over Na₂SO₄, filtered and concentrated. The resulting oil was then dissolved in 6 mL of dichloromethane before the addition of imidazole (414 mg, 7.55 mmol, 5 eq.) and TBSCl (274 mg, 1.81 mmol, 1.2 eq.) at 0 °C. The resulting mixture was stirred at room temperature for 1 hour before it was quenched by 5 mL of saturated aqueous NH₄Cl. The organic layer was removed, and the aqueous layer was extracted with ethyl acetate (30 mL) three times. The combined organic layers were dried over MgSO₄, filtered and concentrated. Purification over SiO₂ (30% EtOAc in Hexanes) afforded **15** as a clear oil (1.02 g, 1.21 mmol, 80% for 2 steps). **15** was obtained as an equilibrium mixture of ketone form and ketal form (ca. 1:1 ratio in C₆D₆) along with ~8:1 mixture of *E/Z* isomers.



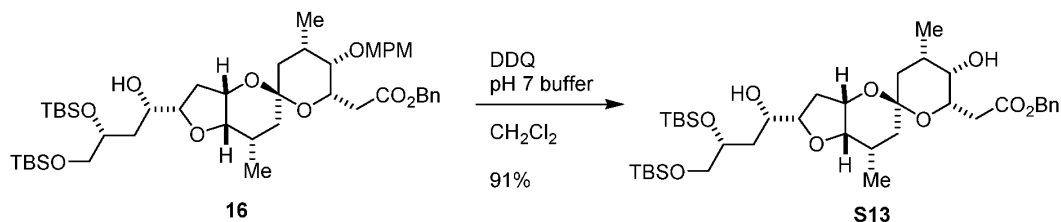
[00809] To a solution of **15** (600 mg, 0.711 mmol, 1 eq.), BnOAc (122 μ L, 0.711 mmol, 1 eq.), and LiCl (302 mg, 7.11 mmol, 10 eq.) in MeCN (14.2 mL, 0.05 M) was added DBU (2.13 mL, 14.2 mmol, 20 eq.). After being stirred for 12 hr at room temperature, the reaction mixture was loaded to a SiO₂ pad and washed by EtOAc. The mixture was concentrated and purified over SiO₂ (25% EtOAc in Hexanes) to afford **16** (516 mg, 0.612 mmol, 86%, *dr* = 22:1) along with byproduct **18** (48 mg, 0.057 mmol, 8%). **16**: [α]_D²² = -29.3 (c 1.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.23 (2H, d, *J* = 7.1 Hz), 7.19 (2H, d, *J* = 8.4 Hz), 7.11 (2H, dd, *J* = 7.5, 7.1 Hz), 7.06 (1H, dd, *J* = 7.5, 7.5 Hz), 6.76 (2H, d, *J* = 8.4 Hz), 5.09 (1H, d, *J* = 12.0 Hz), 5.00 (1H, d, *J* = 12.0 Hz), 4.27-4.21 (2H, m), 4.15 (2H, d, *J* = 7.6 Hz), 3.95-3.77 (4H,

m), 3.36 (1H, dd, $J = 6.3, 3.6$ Hz), 3.26 (1H, dd, $J = 10.2, 2.4$ Hz), 3.25 (3H, s), 2.89 (1H, dd, $J = 15.0, 10.2$ Hz), 2.77 (1H, s), 2.19 (1H, dd, $J = 16.2, 3.0$ Hz), 2.18-2.09 (1H, m), 2.08-2.02 (1H, m), 1.93-1.84 (2H, m), 1.57 (1H, t, $J = 12.9$ Hz), 1.49 (1H, t, $J = 12.9$ Hz), 1.38 (1H, dd, $J = 12.0, 3.0$ Hz), 1.31 (1H, dd, $J = 12.8, 3.0$ Hz), 1.14 (9H, t, $J = 7.8$ Hz), 1.08 (9H, s), 1.03 (9H, s), 1.01 (3H, d, $J = 5.4$ Hz), 0.88 (1H, t, $J = 8.4$ Hz), 0.82 (3H, q, $J = 7.8$ Hz), 0.81 (3H, q, $J = 7.8$ Hz), 0.57 (6H, q, $J = 7.8$ Hz), 0.28 (3H, s), 0.28 (3H, s), 0.15 (6H, s) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 171.5, 159.8, 136.6, 131.0, 129.7, 128.8, 128.7, 128.4, 114.0, 97.4, 80.5, 79.8, 78.1, 75.4, 72.2, 71.9, 70.7, 70.2, 67.9, 66.2, 54.7, 39.0, 37.9, 37.5, 37.0, 35.7, 30.5, 26.3 ($\times 6$), 18.7, 18.5, 18.2, 18.0, 7.2, 4.9, -4.0, -4.4, -5.1, -5.1 ppm. FTIP (film): 3545, 2956, 2927, 2856, 1736, 1613, 1514, 1462, 1381, 1303, 1249, 1096, 1038, 944, 835, 777 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{46}\text{H}_{74}\text{O}_{10}\text{NaSi}_2^+$ 865.4713; found 865.4723. **18 (C-Michael product)**: $[\alpha]_D^{22} = -10.0$ (c 0.95, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.14 (1H, d, $J = 8.4$ Hz), 6.82 (1H, d, $J = 9.0$ Hz), 4.18 (1H, d, $J = 12.0$ Hz), 4.18-4.15 (1H, m), 4.12 (1H, d, $J = 12.0$ Hz), 4.06 (1H, dd, $J = 4.8, 1.8$ Hz), 3.95-3.92 (1H, m), 3.83-3.78 (1H, m), 3.77-3.71 (1H, m), 3.34 (3H, s), 3.06 (1H, dd, $J = 3.0, 2.4$ Hz), 2.91 (1H, dd, $J = 4.8, 2.4$ Hz), 2.82 (1H, dd, $J = 3.0, 3.0$ Hz), 2.21 (1H, dd, $J = 12.6, 2.4$ Hz), 2.17 (1H, dd, $J = 19.2, 8.4$ Hz), 2.02 (1H, dd, $J = 8.1, 3.3$ Hz), 1.90-1.83 (3H, m), 1.80-1.74 (2H, m), 1.70 (1H, ddd, $J = 14.4, 9.6, 4.8$ Hz), 1.56-1.52 (1H, m), 1.35-1.30 (2H, m), 1.03 (9H, s), 0.99 (9H, s), 0.81 (3H, d, $J = 6.6$ Hz), 0.78 (1H, d, $J = 6.0$ Hz), 0.16 (6H, s), 0.11 (3H, s), 0.10 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 169.1, 159.8, 131.0, 129.3, 129.3, 114.1, 114.1, 103.9, 81.1, 80.5, 79.7, 72.9, 71.8, 71.1, 70.8, 67.7, 54.8, 40.7, 38.2, 35.1, 33.0, 30.6, 29.5, 29.2, 27.4, 26.2 ($\times 6$), 16.8, 14.4, -4.2, -4.5, -5.1, -5.2 ppm. FTIP (film): 3507, 2954, 2929, 2879, 2856, 1732, 1612, 1513, 1462, 1382, 1363, 1210, 1158, 1068, 1004, 944, 834, 777 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{46}\text{H}_{74}\text{O}_{10}\text{NaSi}_2^+$ 865.4713; found 865.4721.

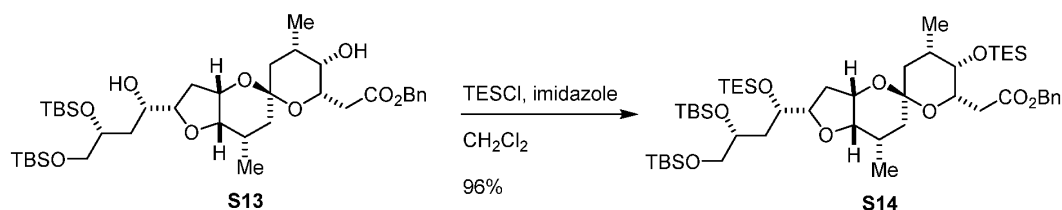


[00810] To a solution of **15** (60 mg, 0.071 mmol, 1 eq.), thiourea catalyst **M** (12.7 mg, 0.036 mmol, 0.5 eq.), BnOAc (12 μL , 0.0710 mmol, 1 eq.), and LiCl (30.2 mg, 0.711 mmol, 10 eq.) in MeCN (1.4 mL, 0.05 M) was added DBU (0.21 mL, 1.42 mmol, 20 eq.). After being

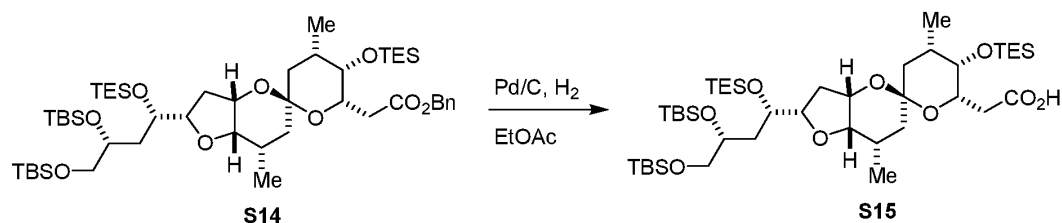
stirred for 2 hr at room temperature, the reaction mixture was loaded to a SiO₂ pad and washed by 33% EtOAc in Hexanes. The mixture was concentrated and dissolved in 1.4 mL of dichloromethane before the addition of BnOAc (12 μ L, 0.0710 mmol, 1 eq.), LiCl (30.2 mg, 0.711 mmol, 10 eq.) and DBU (0.21 mL, 1.42 mmol, 20 eq.). After being stirred for 24 hr at room temperature, the reaction mixture was loaded to a SiO₂ pad and washed by EtOAc. The mixture was concentrated and purified over SiO₂ (25% EtOAc in Hexanes) to afford **16** (55.8 mg, 0.066 mmol, 93%, *dr* > 25:1).



[00811] To a solution of **16** (650 mg, 0.77 mmol) in dichloromethane (15 mL) and phosphate buffer (pH=7, 2.5 mL) was added DDQ (437 mg, 1.93 mmol). After being stirred for 40 min, the reaction was quenched with 10% aqueous Na₂S₂O₃ and saturated aqueous NaHCO₃. The mixture was extracted twice with DCM, dried over Na₂SO₄, filtered, and concentrated. The residue was purified by flash column chromatography on neutral silica gel (33% EtOAc in Hexanes) to afford **S13** (518 mg, 0.72 mmol, 93%) as a colorless oil. **S13**: [α]_D²² = -31.2 (c 4.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.23 (2H, d, *J* = 7.1 Hz), 7.11 (2H, dd, *J* = 7.5, 7.1 Hz), 7.06 (1H, dd, *J* = 7.5, 7.5 Hz), 5.09 (1H, d, *J* = 12.0 Hz), 5.00 (1H, d, *J* = 12.0 Hz), 4.22-4.18 (2H, m), 4.02 (1H, ddd, *J* = 6.6, 3.6, 1.2 Hz), 3.95-3.74 (4H, m), 3.26 (1H, d, *J* = 2.4 Hz), 3.22 (1H, dd, *J* = 2.4, 2.4 Hz), 2.87 (1H, d, *J* = 8.4 Hz), 2.78 (1H, dd, *J* = 15.9, 10.9 Hz), 2.21 (1H, dd, *J* = 14.6, 3.0 Hz), 2.11-2.06 (1H, m), 2.00-1.92 (3H, m), 1.85-1.78 (1H, m), 1.49 (1H, t, *J* = 12.9 Hz), 1.30 (1H, dd, *J* = 14.4, 4.8 Hz), 1.17 (1H, dd, *J* = 14.4, 4.8 Hz), 1.15 (1H, d, *J* = 4.0 Hz), 1.11 (1H, d, *J* = 12.6 Hz), 1.00 (3H, d, *J* = 6.6 Hz), 0.99 (9H, s), 0.96 (9H, s), 0.75 (3H, d, *J* = 6.6 Hz), 0.15 (3H, s), 0.12 (3H, s), 0.08 (3H, s), 0.07 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 171.4, 128.8, 128.7, 128.4, 97.2, 80.5, 79.6, 72.1, 71.9, 70.7, 70.1, 69.9, 67.8, 66.2, 38.9, 37.3, 37.0, 37.0, 35.6, 30.0, 26.3 ($\times$ 6), 18.6, 18.4, 18.2, 17.5, -4.1, -4.5, -5.1, -5.1 ppm. FTIP (film): 3560, 2955, 2928, 2856, 1737, 1471, 1376, 1361, 1252, 1099, 1016, 835, 777 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₃₈H₆₆O₉NaSi₂⁺ 745.4138; found 745.4143.

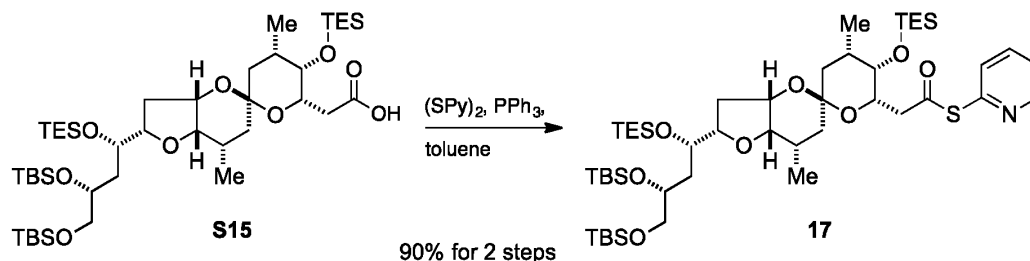


[00812] To a solution of **S13** (495 mg, 0.685 mmol) in dichloromethane (7 mL) was added imidazole (233 mg, 3.43 mmol) and TESCl (0.35 mL, 2.06 mmol). After being for 2 hr at room temperature, the reaction was quenched with MeOH. The mixture was concentrated and purified by flash column chromatography on neutral silica gel (10% EtOAc in Hexanes) to give **S14** (632 mg, 0.66 mmol, 96%) as a colorless oil. **S14**: $[\alpha]_D^{22} = -44.3$ (c 1.0, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.23 (2H, d, $J = 7.1$ Hz), 7.11 (2H, dd, $J = 7.5, 7.1$ Hz), 7.06 (1H, dd, $J = 7.5, 7.5$ Hz), 5.09 (1H, d, $J = 12.0$ Hz), 5.00 (1H, d, $J = 12.0$ Hz), 4.23 (1H, dd, $J = 5.7, 2.1$ Hz), 4.20-4.16 (1H, m), 4.11 (1H, dd, $J = 9.6, 2.4$ Hz), 4.08 (1H, ddd, $J = 9.6, 7.2, 3.0$ Hz), 3.83 (1H, ddd, $J = 8.4, 5.4, 5.4$ Hz), 3.76 (1H, dd, $J = 10.2, 3.0$ Hz), 3.70 (1H, dd, $J = 10.2, 5.4$ Hz), 3.21 (1H, s), 3.19 (1H, d, $J = 3.0$ Hz), 2.84 (1H, dd, $J = 15.6, 10.2$ Hz), 2.25 (1H, dd, $J = 15.0, 3.6$ Hz), 2.22-2.16 (1H, m), 2.15-2.11 (1H, m), 1.96 (1H, ddd, $J = 13.8, 8.4, 3.0$ Hz), 1.88 (1H, ddd, $J = 13.6, 9.6, 6.0$ Hz), 1.83 (1H, dd, $J = 13.8, 6.0$ Hz), 1.73 (1H, ddd, $J = 13.8, 9.0, 4.2$ Hz), 1.59 (1H, t, $J = 13.2$ Hz), 1.52 (1H, t, $J = 13.2$ Hz), 1.40 (1H, t, $J = 4.2$ Hz), 1.37 (1H, t, $J = 4.2$ Hz), 1.06 (9H, t, $J = 7.8$ Hz), 1.04 (9H, s), 1.00 (3H, d, $J = 6.6$ Hz), 0.98 (9H, s), 0.92 (1H, t, $J = 8.4$ Hz), 0.87 (3H, d, $J = 7.2$ Hz), 0.74 (6H, t, $J = 8.4$ Hz), 0.53 (6H, t, $J = 8.8$ Hz), 0.22 (3H, s), 0.22 (3H, s), 0.09 (6H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 171.4, 136.7, 128.9, 128.7, 128.4, 97.0, 81.4, 80.3, 72.2, 72.1, 71.7, 71.5, 70.3, 68.3, 66.3, 38.3, 38.2, 37.6, 37.6, 35.4, 30.7, 26.3 ($\times 6$), 18.6, 18.4, 18.2, 18.2, 7.4, 7.3, 5.8, 5.7, -4.1, -4.2, -5.1, -5.2 ppm. FTIP (film): 2954, 2928, 2877, 2857, 1740, 1471, 1388, 1361, 1211, 1160, 1099, 1081, 1039, 944, 834 cm⁻¹. HRMS (ESI) m/z : [M+Na]⁺ calcd for C₅₀H₉₆O₈NaSi₄⁺ 959.6074; found 959.6070.

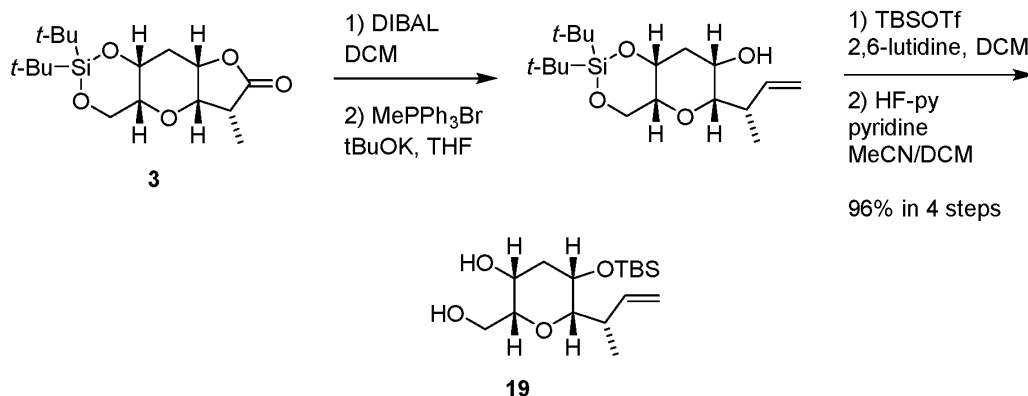


[00813] To a solution of **S14** (330 mg, 0.347 mmol) in EtOAc (3.5 mL) was added Pd/C (33 mg, 10 wt%). The reaction flask was filled with H₂ with lab balloon and stirred for 45 min at

room temperature. The mixture was passed through a pad of SiO₂ with EtOAc, and concentrated. Without further purification the crude acid **S15** was used in the next reaction.



[00814] To a solution of crude acid **S15** (estimated as 0.347 mmol) in toluene (3.5 mL) was added PPh₃ (118 mg, 0.45 mmol, 1.3 eq.) and (PyS)₂ (107 mg, 0.49 mmol, 1.4 eq.).² After being stirred for 3 hr at room temperature, the reaction mixture was concentrated. Flash column chromatography of the residue (neutral SiO₂, hexanes/EtOAc = 1/0, 10/1, 5/1) to give **17** (297 mg, 0.312 mmol, 90% for 2 steps) as a colorless oil. **17**: $[\alpha]_D^{22} = -75.7$ (c 0.4, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 8.32 (1H, dd, $J = 4.2, 1.8$ Hz), 7.54 (1H, d, $J = 7.8$ Hz), 6.96 (1H, ddd, $J = 7.8, 7.8, 1.4$ Hz), 6.47 (1H, dd, $J = 7.8, 4.2$ Hz), 4.34 (1H, dd, $J = 6.0, 1.8$ Hz), 4.24-4.21 (1H, m), 4.17 (1H, dd, $J = 10.2, 2.4$ Hz), 4.12 (1H, dd, $J = 9.0, 6.6, 3.0$ Hz), 3.85 (1H, dt, $J = 9.0, 6.0$ Hz), 3.80 (1H, dd, $J = 10.8, 3.0$ Hz), 3.74 (1H, dd, $J = 10.8, 4.0$ Hz), 3.36 (1H, dd, $J = 2.4, 2.4$ Hz), 3.23 (1H, dd, $J = 14.7, 9.6$ Hz), 3.15 (1H, s), 2.59-2.55 (1H, m), 2.53 (1H, dd, $J = 14.4, 3.0$ Hz), 2.16-2.11 (1H, m), 2.00 (1H, ddd, $J = 13.8, 9.0, 3.0$ Hz), 1.94 (1H, ddd, $J = 13.8, 9.0, 5.4$ Hz), 1.88 (1H, dd, $J = 13.8, 6.0$ Hz), 1.77 (1H, ddd, $J = 13.2, 8.4, 4.2$ Hz), 1.65 (1H, dd, $J = 13.2, 12.6$ Hz), 1.60 (1H, dd, $J = 13.2, 12.6$ Hz), 1.52 (1H, dd, $J = 13.2, 4.8$ Hz), 1.44 (1H, dd, $J = 12.9, 3.4$ Hz), 1.10 (9H, t, $J = 8.4$ Hz), 1.09 (9H, s), 1.07 (3H, d, $J = 7.2$ Hz), 1.03 (9H, s), 0.99 (9H, t, $J = 7.8$ Hz), 0.91 (3H, d, $J = 6.6$ Hz), 0.77 (6H, q, $J = 7.8$ Hz), 0.57 (6H, q, $J = 7.8$ Hz), 0.27 (6H, s), 0.137 (3H, s), 0.136 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 194.7, 152.6, 150.6, 136.5, 129.8, 123.2, 97.2, 81.4, 80.4, 72.3, 72.0, 71.9, 71.5, 70.5, 68.3, 47.6, 38.2, 37.7, 37.6, 35.3, 30.7, 26.3($\times 3$), 26.3($\times 3$), 18.6, 18.5, 18.5, 18.3, 7.4($\times 6$), 5.8($\times 2$), 5.7($\times 6$), -4.0, -4.2, -5.1, -5.2 ppm. FTIR (film): 2956, 2926, 2877, 1716, 1573, 1471, 1251, 837 775, 775, 728 cm⁻¹. HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for C₄₈H₉₂NO₈SSi₄, 954.5615; found, 954.5612.

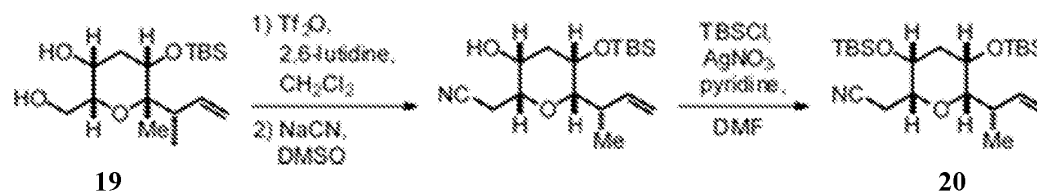
Homohalichondrin Left Halves

[00815] To a stirred solution of lactone **3** (5.01 g, 14.6 mmol, 1 eq.) in CH₂Cl₂ (70 mL) was added DIBAL solution (19.0 mL of 1 M in Hexanes, 19.0 mmol, 1.3 eq.) at -78 °C over 15 min. After being stirred for 15 min at the same temperature, MeOH (2.0 mL), sat. Rochelle's salt aq. (70 mL), and EtOAc (70 mL) were added sequentially. The resulting solution was stirred for 2 h at room temperature to give a clear biphasic solution. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure to give a crude lactol as a colorless solid, which was used in the next reaction without further purification.

[00816] To a stirred suspension of MePPh₃Br (21.0 g, 58.8 mmol, 4 eq.) in THF (50 mL) was added tBuOK (4.9 g, 43.7 mmol, 3 eq.) at 0 °C. After being stirred for 1 h at room temperature, the yellow solution was re-cooled to 0 °C. To this ylide solution was added a solution of the crude lactol (calculated as 14.6 mmol, 1 eq.) in THF (25 mL). After being stirred for 20 min at room temperature, the reaction was quenched with sat. NH₄Cl aq. (50 mL) and H₂O (20 mL). After adding Et₂O (100 mL), the organic layer was separated. The aqueous layer was extracted with Et₂O. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was dissolved in minimum amount of CH₂Cl₂ (ca. 5 mL) and passed through a pad of silica gel (20% EtOAc in Hexanes). The filtrate was concentrated under reduced pressure to give a crude alcohol as a colorless solid, which was used in the next reaction without further purification.

[00817] To a stirred solution of the crude alcohol (calculated as 14.6 mmol, 1 eq.) and 2,6-lutidine (3.4 mL, 29.2 mmol, 2 eq.) in CH₂Cl₂ (70 mL) was added TBSOTf (4.4 mL, 19.2 mmol 1.3 eq.) at 0 °C. After being stirred for 30 min at 0 °C, the ice bath was removed and the reaction mixture was stirred for additional 30 min at room temperature. The reaction was quenched with brine and diluted with Et₂O (200 mL). The organic layer was separated and

[00818] A buffered HF·Py solution was prepared by adding HF·Py (3.0 mL of ~70% HF in pyridine, ca. 114 mmol, ca. 8 eq.) to a mixture of pyridine (15 mL) and MeCN (20 mL) at 0 °C. To a stirred solution of the crude TBS ether (calculated as 14.6 mmol, 1 eq.) in MeCN (50 mL) and CH₂Cl₂ (30 mL) was added the buffered HF·Py solution at -10 °C over 15 min. After being stirred for 30 min at the same temperature, the reaction was warmed up to room temperature. After being stirred for 1 h at the same temperature, the mixture was cooled to 0 °C and quenched with sat. NaHCO₃ aq. carefully. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel (25% then 50% EtOAc in Hexanes) to give diol **19** (4.43 g, 14.0 mmol, 96% for 4 steps) as a colorless amorphous solid. **19**: [α]²⁰_D -36.7 (*c* 1.06, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 6.01 (1H, ddd, *J* = 16.9, 10.5, 6.2 Hz), 5.10 (1H, d, *J* = 16.9 Hz), 5.05 (1H, d, *J* = 10.5 Hz), 4.04 (1H, ddd, *J* = 11.1, 7.8, 3.0 Hz), 3.77 (1H, ddd, *J* = 11.1, 9.2, 4.4 Hz), 3.57 (1H, brs), 3.51-3.46 (2H, m), 3.18 (1H, dd, *J* = 7.5, 4.5 Hz), 2.68-2.66 (2H, m), 2.01 (1H, dd, *J* = 8.4, 3.0 Hz), 1.95 (1H, ddd, *J* = 14.6, 2.7, 2.7 Hz), 1.12 (1H, ddd, *J* = 14.6, 2.7, 2.7 Hz), 0.90 (9H, s), 0.82 (3H, d, *J* = 5.4 Hz), 0.07 (3H, s), -0.06 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 141.9, 114.0, 84.2, 81.7, 66.9, 65.7, 63.4, 38.1, 37.1, 25.9, 18.2, 15.7, -3.8, -5.1 ppm. FTIR (film): 3527, 3259, 2960, 2929, 2858, 1256, 1090, 1056, 879, 776 cm⁻¹. HRMS (ESI) *m/z*: [M+H]⁺ calcd for C₁₆H₃₃O₄Si, 317.2143; found, 317.2145.



[00819] To a stirred solution of diol **19** (7.30 g, 23.1 mmol, 1 eq.) and 2,6-lutidine (10.8 mL, 92.7 mmol, 4 eq.) in CH₂Cl₂ (150 mL) was added Tf₂O (4.7 mL, 27.9 mmol, 1.2 eq.) at -78 °C. After being stirred for 10 min at the same temperature, the reaction was quenched with MeOH (1.0 mL) and brine (100 mL). After adding Et₂O (500 mL), the organic layer was separated from the aqueous layer and washed with 1N HCl and brine sequentially. The

organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure to give a crude triflate as wine red oil, which was dissolved in DMSO (150 mL) immediately without further purification.

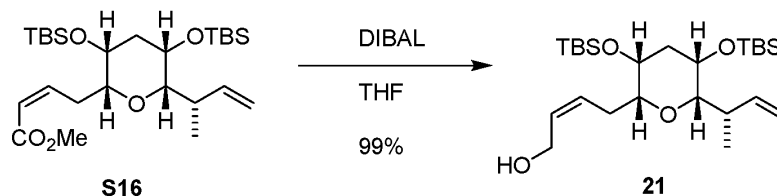
[00820] To the DMSO solution of crude triflate (calculated as 23.1 mmol, 1 eq.) was added NaCN (11.3 g, 230 mmol, 10 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was filtered through a paper and the filter cake was washed with EtOAc thoroughly. The filtrate was washed with H₂O and brine. The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was passed through a pad of silica gel (20% EtOAc in Hexanes) to give a crude nitrile as a pink oil, which was used in the next step without further purification.

[00821] To a stirred solution of the crude nitrile (calculated as 23.1 mmol, 1 eq.) and pyridine (15.0 mL, 186 mmol, 8 eq.) in DMF (80 mL) were added TBSCl (10.4 g, 69.0 mmol, 3 eq.) and AgNO₃ (11.8 g, 69.5 mmol, 3 eq.) at 0 °C. After being stirred for 30 min at 0 °C, the cooling bath was removed and the reaction mixture was stirred for 18 h at room temperature. The reaction mixture was diluted with Et₂O (100 mL) and filtered through a bed of Celite. The filter cake was washed thoroughly with Et₂O. The filtrate was washed with H₂O and brine. The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel (0% then 10% EtOAc in Hexanes) to give bis-TBS **20** (8.87 g, 20.2 mmol, 87% for 3 steps) as a colorless solid. **20**: $[\alpha]_D^{20}$ -6.7 (*c* 1.07, CHCl₃). MP: 65-67 °C (recrystallized from Et₂O). ¹H NMR (600 MHz, C₆D₆) δ: 6.31 (1H, ddd, *J* = 17.4, 10.8, 6.6 Hz), 5.19 (1H, d, *J* = 10.8 Hz), 5.15 (1H, d, *J* = 17.4 Hz), 3.52 (1H, brs), 3.23-3.22 (1H, m), 3.11 (1H, ddd, *J* = 8.0, 5.9, 2.0 Hz), 2.82-2.77 (1H, m), 2.66 (1H, d, *J* = 9.6 Hz), 2.40 (1H, dd, *J* = 16.8, 7.8 Hz), 2.07 (1H, dd, *J* = 16.8, 5.4 Hz), 1.80 (1H, ddd, *J* = 15.4, 2.7, 2.7 Hz), 1.32 (1H, ddd, *J* = 15.4, 4.7, 4.7 Hz), 0.98 (9H, s), 0.93 (3H, d, *J* = 6.6 Hz), 0.89 (9H, s), 0.10 (3H, s), 0.01 (3H, s), -0.01 (3H, s), -0.07 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ: 141.7, 117.9, 113.9, 85.2, 77.1, 65.0, 64.4, 38.3, 37.2, 26.4, 20.6, 18.4, 18.3, 15.9, -2.5, -3.5, -5.00, -5.03 ppm. FTIR (film): 2955, 2930, 2886, 2857, 1473, 1388, 1254, 1139, 1099, 1022, 835, cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₃H₄₅NO₃Si₂Na, 462.2830; found, 462.2831.

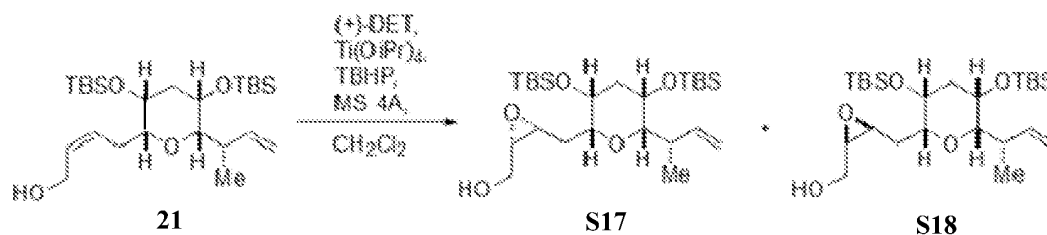


[00823] To a stirred solution of methyl bis(2,2,2-trifluoroethyl) phosphonoacetate (6.3 mL, 29.8 mmol, 1.5 eq.) and 18-crown-6 ether (42.0 g, 159 mmol, 8 eq.) in THF (400 mL) was added KHMDS solution (60 mL of 0.5 M in toluene, 30 mmol, 1.5 eq.) at -78 °C. After being stirred for 30 min at -78 °C, the resulting mixture was added to a solution of the crude aldehyde (calculated as 19.9 mmol, 1 eq.) in THF (100 mL) and stirred for 30 min at the same temperature. The reaction was quenched with sat. NH₄Cl aq. and diluted with Hexanes (400 mL). The organic layer was separated and the aqueous layer was extracted with Et₂O/Hexanes (1:1). The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel (0% then 2% EtOAc in Hexanes) to give α,β-unsaturated ester **S16** (8.37 g, 16.8 mmol, 84% for 2 steps) as a colorless oil. Only Z-isomer was obtained exclusively. **S16**: [α]_D²⁰ +42.0 (*c* 1.03, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ: 6.67-6.63 (1H, m), 6.16 (1H, ddd, *J* = 17.6, 10.2, 6.6 Hz), 5.91 (1H, dd, *J* = 11.1, 1.5 Hz), 5.11 (1H, dd, *J* = 17.6, 1.5 Hz), 5.09 (1H, dd, *J* = 10.2, 1.5 Hz), 3.60 (1H, d, 1.8 Hz), 3.454-3.447 (1H, m), 3.34 (3H, s), 3.34-3.30 (1H, m), 3.13 (1H, d, 10.8 Hz), 3.03-2.97 (1H, m), 2.89-2.83 (1H, m), 2.68 (1H, d, 9.0 Hz), 1.92 (1H, d, 14.6 Hz), 1.45 (1H, ddd, 14.6, 4.5, 4.5 Hz), 1.03 (9H, s), 1.01 (9H, s), 0.93 (3H, d, *J* = 7.2 Hz), 0.19 (3H, s), 0.16 (3H, s), 0.12 (3H, s), 0.04 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ: 166.6, 149.3, 142.6, 119.9, 113.4, 85.3, 81.0, 66.7, 64.7, 50.5, 39.0, 37.6, 32.5, 26.7, 26.4, 18.6, 18.5, 15.9, -2.3, -3.6, -4.6, -5.0 ppm.

FTIR (film): 2951, 2929, 2857, 1723, 1644, 1253, 1091, 951, 836, 771 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{26}\text{H}_{50}\text{O}_5\text{Si}_2\text{Na}$, 521.3089; found, 521.3087.

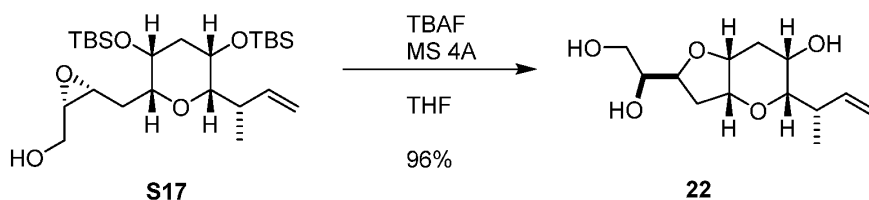


[00824] To a stirred solution of ester **S16** (8.30 g, 16.6 mmol, 1 eq.) in THF (200 mL) was added DIBAL solution (66 mL of 1 M in hexanes, 66.0 mmol, 4 eq.) at -78 °C. After being stirred for 10 min at the same temperature, the reaction mixture was warmed to 0 °C and stirred for additional 30 min. The reaction was quenched with acetone (5.0 mL) and sat Rochelle's salt aq. (200 mL). The resulting mixture was stirred for 2 h at room temperature to give a clear biphasic solution. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel (11% EtOAc in Hexanes) to give allyl alcohol **21** (7.72 g, 16.4 mmol, 99%) as a colorless solid. **21**: [α]_D²⁰ -16.6 (*c* 1.00, CHCl₃). MP: 60-62 °C (recrystallized from Et₂O). ¹H NMR (600 MHz, C₆D₆) δ : 6.24 (1H, ddd, *J* = 17.4, 10.5, 6.9 Hz), 5.83-5.81 (1H, m), 5.77-5.74 (1H, m), 5.18-5.14 (2H, m), 4.23-4.20 (1H, m), 4.14-4.10 (1H, m), 3.62 (1H, d, *J* = 1.2 Hz), 3.349-3.345 (1H, m), 3.05 (1H, d, *J* = 9.6 Hz), 2.89-2.80 (2H, m), 2.72 (1H, d, *J* = 9.0 Hz), 1.96 (1H, dd, *J* = 14.4, 6.6 Hz), 1.91 (1H, ddd, *J* = 14.6, 2.4, 2.4 Hz), 1.78-1.77 (1H, m), 1.49 (1H, ddd, *J* = 14.6, 4.7, 4.7 Hz), 1.01 (18H, s), 0.95 (3H, d, *J* = 7.2 Hz), 0.15 (3H, s), 0.08 (3H, s), 0.05 (3H, s), 0.004 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 142.4, 131.4, 129.2, 113.6, 85.1, 80.6, 66.7, 65.0, 58.8, 39.1, 37.5, 30.5, 26.5, 26.4, 18.53, 18.50, 16.0, -2.3, -3.5, -4.7, -5.0 ppm. FTIR (film): 2952, 2929, 2855, 1463, 1252, 1138, 1003, 833, 768 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₅H₅₀O₄Si₂Na, 493.3140; found, 493.3142.



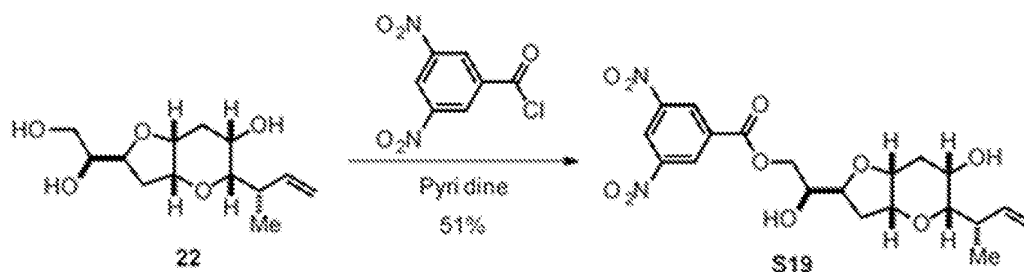
[00825] To a stirred suspension of molecular sieve 4A (1.5 g, activated powder) in CH_2Cl_2 (50 mL) were added Ti(OiPr)_4 (0.72 mL, 2.43 mmol, 15 mol%) and a solution of (+)-DET (0.55 mL, 3.21 mmol, 20 mol%) in CH_2Cl_2 (5.0 mL) at -20°C . After being stirred for 15 min at the same temperature, to the reaction mixture was added TBHP solution (4.5 mL of ~ 5.5 M in decane over molecular sieve 4A, ca. 24.8 mmol, ca. 1.5 eq.) and stirred for 30 min. In a separate flask, a solution of allyl alcohol **21** (7.65 g, 16.2 mmol, 1 eq.) in CH_2Cl_2 (80 mL) was cooled to -78°C . To this cooled allyl alcohol solution was added the above catalyst solution via cannula and rinsed with CH_2Cl_2 (30 mL). After being stirred for 10 min at -78°C , the reaction mixture was warmed to -10°C and stirred for 15 h. The reaction flask was removed from cooling bath, and Et_2O (200 mL) was added to the cold reaction mixture with stirring, followed by addition of sat. Na_2SO_4 aq. (2.5 mL). After being stirred for 2 h at room temperature, the mixture was filtered through a pad of Celite, and the filter cake was washed with EtOAc thoroughly. After removal of solvent under reduced pressure, the obtained crude material was purified by flash column chromatography on silica gel (0%, 5%, 10%, 11%, then 20% EtOAc in Hexanes) to give epoxy alcohol **S17** (6.81 g, 14.0 mmol, 86%) as a colorless solid and its undesired diastereomer **S18** (862 mg, 1.77 mmol, 11%) as a colorless solid. **S17**: $[\alpha]_D^{20} -26.0$ (c 1.08, CHCl_3). MP: $78-79^\circ\text{C}$ (recrystallized from Hexanes/ EtOAc). ^1H NMR (600 MHz, C_6D_6) δ : 6.09-6.03 (1H, m), 5.21 (1H, d, $J = 18.0$ Hz), 5.11 (1H, d, $J = 10.8$ Hz), 3.92-3.88 (1H, m), 3.65 (1H, ddd, $J = 11.4, 7.8, 3.0$ Hz), 3.58 (1H, s), 3.239-3.235 (1H, m), 3.17-3.12 (2H, m), 3.08 (1H, $J = 4.1, 4.1, 4.1$ Hz), 2.78-2.71 (2H, m), 2.64 (1H, d, $J = 9.6$ Hz), 2.13 (1H, ddd, $J = 14.1, 11.4, 8.4$ Hz), 1.84 (1H, ddd, $J = 15.0, 2.4, 2.4$ Hz), 1.57-1.53 (1H, m), 1.41 (1H, ddd, $J = 15.0, 4.2, 4.2$ Hz), 1.01 (9H, s), 0.98 (9H, s), 0.90 (3H, d, $J = 6.6$ Hz), 0.13 (3H, s), 0.03 (3H, s), 0.02 (3H, s), -0.06 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 142.1, 114.1, 85.35, 85.33, 78.8, 66.2, 64.5, 60.9, 56.1, 54.7, 38.8, 37.6, 30.3, 26.5, 26.3, 18.4, 16.0, $-2.3, -3.7, -4.8, -5.1$ ppm. FTIR (film): 2952, 2929, 2856, 1252, 1011, 833, 768 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{NH}_4]^+$ calcd for $\text{C}_{25}\text{H}_{54}\text{NO}_5\text{Si}_2$, 504.3535; found, 504.3527. **Diastereomer S18**: $[\alpha]_D^{20} +11.9$ (c 1.00, CHCl_3). MP: $89-91^\circ\text{C}$ (recrystallized from Et_2O). ^1H NMR (600 MHz, C_6D_6) δ : 6.19 (1H, ddd, $J = 17.4, 10.7, 6.9$ Hz), 5.17 (1H, ddd, $J = 17.4,$

1.8, 1.8 Hz), 5.12 (1H, d, $J = 10.7$ Hz), 3.70-3.68 (1H, m), 3.65 (1H, ddd, $J = 11.4, 6.0, 6.0$ Hz), 3.59 (1H, ddd, $J = 11.4, 5.4, 5.4$ Hz), 3.49 (1H, dd, $J = 8.4, 4.8$ Hz), 3.44 (1H, ddd, $J = 7.8, 4.2, 4.2$ Hz), 3.41 (1H, ddd, $J = 8.4, 3.6, 3.6$ Hz), 3.02 (1H, dd, $J = 10.8, 5.4$ Hz), 2.89-2.83 (1H, m), 2.76 (1H, dd, $J = 10.2, 2.4$ Hz), 2.44 (1H, ddd, $J = 13.8, 8.4, 5.1$ Hz), 1.95 (1H, ddd, $J = 14.4, 4.8, 3.6$ Hz), 1.65 (1H, ddd, $J = 14.4, 4.8, 4.8$ Hz), 1.57 (1H, ddd, $J = 14.4, 7.8, 3.6$ Hz), 1.59-1.50 (1H, m), 1.04 (9H, s), 0.98-0.97 (12H, m), 0.16 (3H, s), 0.07 (3H, s), 0.06 (3H, s), 0.00 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 142.3, 113.7, 84.4, 79.2, 66.5, 65.3, 60.7, 56.0, 54.5, 39.1, 37.4, 30.9, 26.4, 26.3, 18.5, 18.4, 15.9, -2.5, -3.8, -4.8, -5.0 ppm. FTIR (film): 3448, 2953, 2929, 2857, 1463, 1253, 1128, 1088, 1005, 902, 833, 770, 735 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{25}\text{H}_{50}\text{O}_5\text{Si}_2\text{Na}$, 509.3089; found, 509.3088.

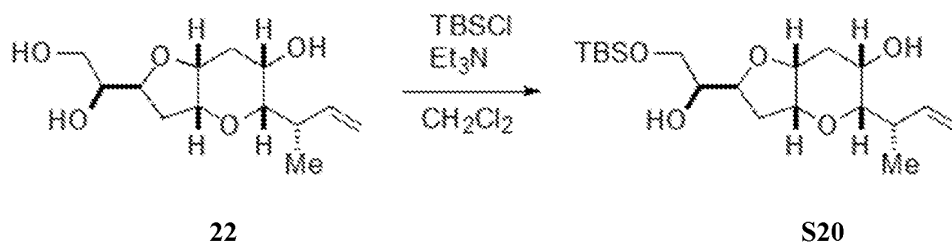


[00826] To a stirred solution of epoxy alcohol **S17** (6.73 g, 13.8 mmol, 1 eq.) in THF (20.0 mL) were added molecular sieves 4A (4.0 g, activated pelet) and TBAF solution (80.0 mL of 1 M in THF, 80.0 mmol, 5.8 eq.) at room temperature. After being stirred for 12 h at the same temperature, CaCO_3 (20.0 g), DOWEX 50WX8-400 (60.0 g), MeOH (10 mL), and THF (100 mL) were added sequentially. After being stirred for 1.5 h at room temperature, the mixture was filtered through a pad of Celite, and the filtrand was washed with EtOAc thoroughly. After removal of solvent under reduced pressure, the obtained crude material was purified by flash column chromatography on silica gel (CH_2Cl_2 then EtOAc) to give triol **22** (3.73 g, 13.3 mmol, 96%) as a colorless amorphous solid. **22**: $[\alpha]_D^{20}$ -64.8 (c 1.00, CHCl_3). ^1H NMR (600 MHz, CD_3OD) δ : 5.96 (1H, ddd, $J = 17.5, 10.5, 7.1$ Hz), 5.03 (1H, ddd, 17.5, 1.6, 1.6 Hz), 4.97 (1H, ddd, $J = 10.5, 1.6, 1.5$ Hz), 4.35 (1H, ddd, 9.5, 6.9, 4.1 Hz), 4.05 (1H, d, $J = 3.0$ Hz), 3.93 (1H, s), 3.73 (1H, dd, $J = 3.0, 3.0$ Hz), 3.61 (1H, dd, $J = 11.1, 5.7$ Hz), 3.57 (1H, dd, 11.1, 5.7 Hz), 3.46 (1H, dd, $J = 6.0, 6.0, 4.2$ Hz), 3.01 (1H, d, $J = 9.0$ Hz), 2.50-2.44 (1H, m), 2.31 (1H, ddd, $J = 15.5, 2.4, 2.4$ Hz), 2.09 (1H, ddd, 13.2, 9.2, 3.8 Hz), 2.04 (1H, dd, $J = 13.2, 6.6$ Hz), 1.89 (1H, ddd, $J = 15.5, 3.8, 3.8$ Hz), 0.99 (3H, d, $J = 7.2$ Hz) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 143.5, 113.8, 82.8, 79.3, 79.0, 77.4, 74.9, 64.8, 64.6, 39.4, 36.6, 33.1, 15.6 ppm. FTIR (film): 3387, 2935, 2886, 1638, 1415, 1093, 1032, 838 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{13}\text{H}_{22}\text{O}_5\text{Na}$, 281.1359; found, 281.1359. The stereochemistry of triol

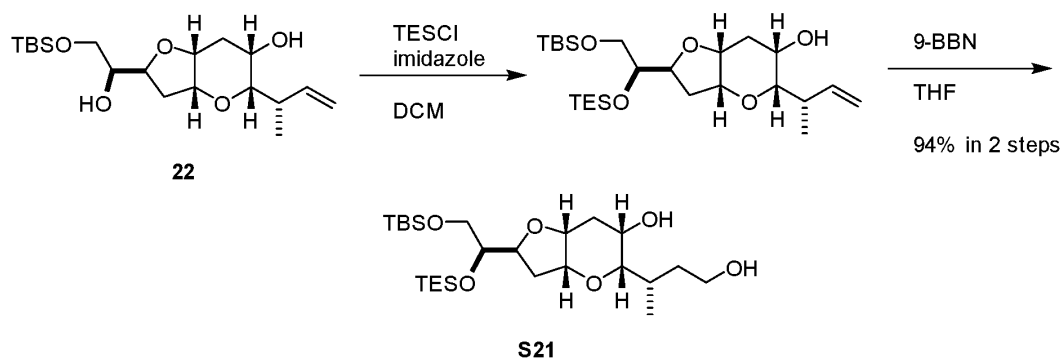
22 was confirmed by X-ray crystallographic analysis on its 3,5-dinitrobenzoate derivative **S19**.



[00827] To a stirred solution of triol **22** (16.4 mg, 0.0635 mmol) in pyridine (0.50 mL) was added 3,5-dinitrobenzoyl chloride (20.0 mg, 0.0867 mmol, 1.4 eq.) at room temperature. After being stirred for 5 h at the same temperature, the resultant reaction mixture was diluted with EtOAc and was washed with 1N HCl, sat. NaHCO₃ aq., and brine sequentially. The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by PTLC (EtOAc) to give dinitrobenzoate **S19** (14.7 mg, 0.0325 mmol, 51%) as a colorless solid. Recrystallization from Hexanes/EtOAc gave a single crystal suitable for X-ray diffraction studies. **S19**: $[\alpha]_D^{20}$ -51.3 (*c* 0.735, CHCl₃). MP: 127-129 °C (recrystallized from Hexanes/EtOAc). ¹H NMR (600 MHz, C₆D₆) δ : 8.70 (2H, d, *J* = 2.1 Hz), 8.48 (1H, t, *J* = 2.1 Hz), 6.13 (1H, ddd, *J* = 16.5, 10.3, 6.9 Hz), 5.18 (1H, d, *J* = 16.5 Hz), 5.14 (1H, d, *J* = 10.3 Hz), 4.28 (1H, dd, *J* = 11.4, 7.2 Hz), 4.00-3.95 (2H, m), 3.55 (1H, ddd, *J* = 10.2, 3.0, 3.0 Hz), 3.48 (1H, d, *J* = 3.0 Hz), 3.38 (1H, s), 3.29-3.26 (1H, m), 2.97 (1H, d, *J* = 10.8 Hz), 2.86-2.82 (1H, m), 2.61 (1H, d, *J* = 9.0 Hz), 2.22 (1H, d, *J* = 15.3 Hz), 2.01-1.96 (1H, m), 1.86 (1H, dd, *J* = 12.8, 6.9 Hz), 1.68 (1H, ddd, *J* = 12.8, 9.6, 3.6 Hz), 1.20 (1H, ddd, *J* = 15.3, 3.3, 3.3 Hz), 1.03 (3H, d, *J* = 7.2 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 162.7, 148.3, 142.5, 133.0, 128.8, 122.1, 113.8, 82.1, 78.0, 77.6, 77.2, 71.4, 68.4, 63.5, 38.7, 36.1, 32.3, 15.3 ppm. FTIR (film): 3489, 3423, 3100, 2919, 1733, 1545, 1345, 1282, 1170, 1095, 921, 721 cm⁻¹. HRMS (ESI) *m/z*: [M+NH₄]⁺ calcd for C₂₀H₂₈N₃O₁₀, 470.1769; found, 470.1779.



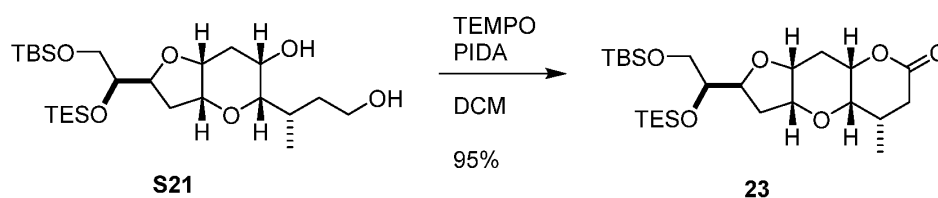
[00828] To a stirred solution of triol **22** (480 mg, 1.71 mmol, 1 eq.) in CH_2Cl_2 (9.0 mL) were added Et_3N (0.95 mL, 6.82 mmol, 4 eq.) and TBSCl (385 mg, 2.55 mmol, 1.5 eq.) at room temperature. After being stirred for 5 h, to the reaction mixture was added TBSCl (200 mg, 1.33 mmol, 0.8 eq.) and stirred for an additional 15 h. The reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on silica gel (17% then 25% EtOAc in Hexanes) to give TBS ether **S20** (633 mg, 1.70 mmol, 99%) as a colorless solid. **S20**: $[\alpha]_D^{20}$ -41.6 (c 1.00, CHCl_3). MP: 55-58 °C (recrystallized from Et_2O). ^1H NMR (600 MHz, CD_3OD) δ : 5.97 (1H, ddd, $J = 17.4, 10.5, 7.5$ Hz), 5.04 (1H, d, $J = 17.4$ Hz), 4.98 (1H, d, $J = 10.5$ Hz), 4.41-4.38 (1H, m), 4.05 (1H, s), 3.93 (1H, s), 3.73 (1H, s), 3.72-3.66 (2H, m), 3.45 (1H, dd, 9.3, 5.7 Hz), 3.02 (1H, d, $J = 9.0$ Hz), 2.50-2.44 (1H, m), 2.32 (1H, d, $J = 15.2$ Hz), 2.13 (1H, ddd, $J = 13.1, 9.6, 3.4$ Hz), 2.04 (1H, dd, $J = 13.1, 7.2$ Hz), 1.89 (1H, ddd, $J = 15.2, 3.3, 3.3$ Hz), 0.99 (3H, d, $J = 7.2$ Hz), 0.92 (9H, s), 0.09 (6H, s) ppm. ^{13}C NMR (125 MHz, CD_3OD) δ : 143.4, 113.9, 82.8, 79.3, 78.6, 77.6, 74.8, 65.7, 65.0, 39.5, 36.5, 33.0, 26.4, 19.2, 15.6, -5.2, -5.3 ppm. FTIR (film): 3504, 2953, 2930, 2886, 2857, 1255, 1095, 837, 755 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{19}\text{H}_{36}\text{O}_5\text{SiNa}$, 395.2224; found, 395.2225.



[00829] To a stirred solution of TBS ether **S20** (540 mg, 1.45 mmol, 1 eq.) in CH_2Cl_2 (20 mL) were added imidazole (300 mg, 4.41 mmol, 3 eq.) and TESCl (0.29 mL, 1.73 mmol, 1.2

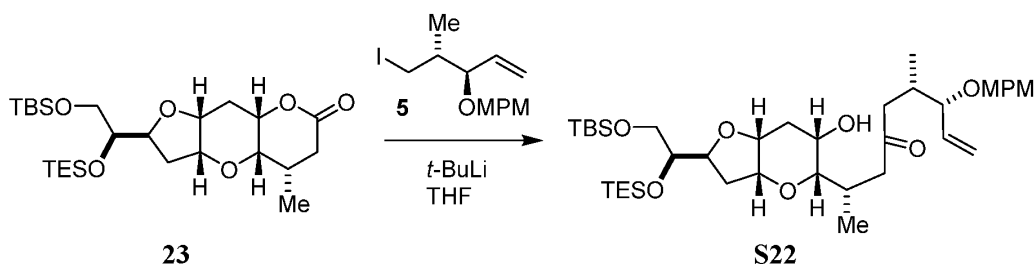
eq.) at 0 °C. After being stirred for 15 min at 0 °C, the reaction was quenched with brine. After separation of the organic layer, the aqueous layer was extracted with Et₂O. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was dissolved in hexanes and filtered through a plug of silica gel (20% EtOAc in Hexanes) to give a crude TES ether as pale yellow oil, which was used in the next step without further purification.

[00830] The crude TES ether (calculated as 1.45 mmol, 1 eq.) was cooled to 0 °C and dissolved in pre-cooled 9-BBN solution (9.0 mL of 0.5 M in THF, 4.50 mmol, 3 eq., ca. 5 °C). After being stirred for 5 min at 0 °C, the ice bath was removed and the reaction mixture was stirred for 1 h at room temperature. The reaction mixture was cooled to 0 °C and quenched with H₂O (10 mL). After adding NaBO₃•H₂O (2.60 g, 26.0 mmol, 18 eq.) at the same temperature, the resulting mixture was stirred vigorously for 1 h at room temperature. The mixture was diluted with brine (10 mL) and EtOAc (20 mL). The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (CH₂Cl₂ then 0%, 9%, 17%, 20%, 25%, then 33% EtOAc in Hexanes) to give diol **S21** (689 mg, 1.36 mmol, 94% for 2 steps) as a colorless oil. **S21**: [α]_D²⁰ +3.27 (*c* 1.04, CHCl₃). ¹H NMR (600 MHz, CD₃OD) δ : 4.36 (1H, ddd, *J* = 8.1, 8.1, 3.6 Hz), 4.07 (1H, d, *J* = 3.0 Hz), 3.90 (1H, s), 3.73 (1H, s), 3.70 (1H, dd, *J* = 9.9, 6.3 Hz), 3.68-3.59 (4H, m), 2.96 (1H, d, *J* = 9.6 Hz), 2.29 (1H, d, *J* = 15.0 Hz), 2.09 (1H, ddd, *J* = 13.2, 9.0, 4.2 Hz), 2.04 (1H, dd, *J* = 13.2, 6.6 Hz), 2.03-1.98 (1H, m), 1.91-1.87 (2H, m), 1.40-1.34 (1H, m), 0.98 (9H, t, *J* = 7.8 Hz), 0.92-0.90 (12H, m), 0.65 (6H, q, *J* = 7.8 Hz), 0.084 (3H, s), 0.079 (3H, s) ppm. ¹³C NMR (125 MHz, CD₃OD) δ : 83.3, 78.4, 79.2, 77.5, 76.8, 66.2, 65.1, 61.5, 37.4, 36.5, 33.0, 32.3, 26.44, 26.38, 19.2, 15.7, 7.3, 6.0, -5.2 ppm. FTIR (film): 3518, 2952, 2931, 2876, 1252, 1092, 835, 777, 741 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₂₅H₅₂O₆Si₂Na, 527.3195; found, 527.3194.



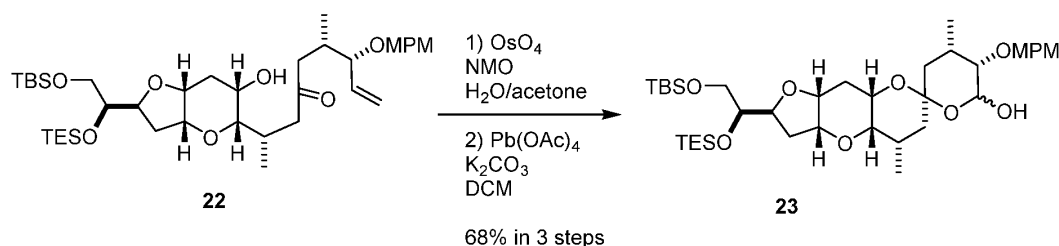
[00831] To a stirred solution of diol **S21** (202 mg, 0.400 mmol) in CH₂Cl₂ (5.0 mL) were added TEMPO (12.5 mg, 20 mol%) and PIDA (390 mg, 3 eq.) at room temperature. After

being stirred for 36 h, the reaction was quenched with sat. NaHCO_3 aq. and sat. $\text{Na}_2\text{S}_2\text{O}_3$ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 9%, 20%, 25%, then 33% EtOAc in Hexanes) to give lactone **23** (191 mg, 0.381 mmol, 95%) as a colorless oil. **23**: $[\alpha]_D^{20}$ -30.0 (*c* 1.02, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 4.44-4.41 (1H, m), 3.95-3.92 (1H, m), 3.85-3.82 (1H, m), 3.70-3.66 (2H, m), 3.54 (1H, d, J = 1.8 Hz), 3.46 (1H, brs), 2.52 (1H, s), 2.36-2.31 (2H, m), 2.16 (1H, dd, J = 18.0, 6.0 Hz), 2.05-2.00 (1H, m), 1.90 (1H, dd, J = 12.9, 6.6 Hz), 1.34-1.30 (1H, m), 1.17 (1H, ddd, J = 15.8, 3.9, 3.9 Hz), 1.04 (9H, t, J = 7.8 Hz), 1.00 (9H, s), 0.76 (3H, d, J = 6.6 Hz), 0.68 (6H, q, J = 7.8 Hz), 0.18 (3H, s), 0.15 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 168.4, 78.2, 77.9, 75.8, 73.8, 73.0, 71.4, 65.8, 35.6, 33.0, 31.3, 30.9, 26.2, 18.6, 16.7, 7.2, 5.6, -5.2 -5.3 ppm. FTIR (film): 2953, 2929, 2877, 1730, 1461, 1246, 1117, 1085, 1002, 856, 776, 741, 668 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{25}\text{H}_{49}\text{O}_6\text{Si}_2$, 501.3062; found, 501.3063.



[00832] To a stirred solution of lactone **23** (467 mg, 0.932 mmol) and iodide (420 mg, 1.3 eq.) in THF (6.0 mL) was added *t*BuLi solution (1.3 mL of 1.7 M in pentane, 2.21 mmol, 2.5 eq.) over 20 min at -78 °C. After being stirred for 30 min, the reaction was quenched with sat. NH_4Cl aq. and stirred for 10 min at room temperature. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 5%, 7%, then 17% EtOAc in Hexanes) to give ketone, which contained a small amount of impurity. The pure compound, which exists as an equilibrium mixture of ketone and hemiacetal (ca. 10:1 ratio in C_6D_6) was obtained for analytical purpose by PTLC (Hexanes/EtOAc = 2:1). Spectral data only for major ketone form of **S22** are shown here. **S22**: $[\alpha]_D^{20}$ -1.4 (*c* 1.13, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.28 (2H, d, J = 8.1 Hz), 6.85 (2H, d, J = 8.1 Hz), 5.64 (1H, ddd, J =

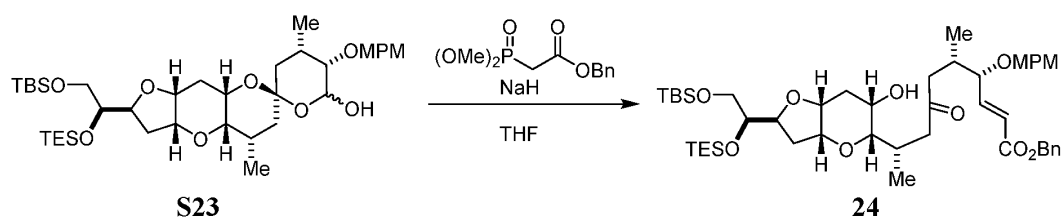
16.8, 9.0, 7.5 Hz), 5.13 (1H, d, $J = 9.0$ Hz), 5.11 (1H, d, $J = 16.8$ Hz), 4.57 (1H, d, $J = 11.4$ Hz), 4.49 (1H, ddd, $J = 6.3, 6.3, 3.3$ Hz), 4.23 (1H, d, $J = 10.8$ Hz), 3.83 (1H, dd, $J = 9.9, 6.3$ Hz), 3.75 (1H, dd, $J = 10.8, 6.0$ Hz), 3.74 (1H, s), 3.63 (1H, d, $J = 3.0$ Hz), 3.60-3.55 (2H, m), 3.47 (1H, dd, $J = 7.5, 7.5$ Hz), 3.37 (1H, d, $J = 10.8$ Hz), 3.33 (3H, s), 2.79 (1H, dd, $J = 16.8, 4.2$ Hz), 2.77-2.69 (3H, m), 2.52-2.50 (1H, m), 2.30 (1H, d, $J = 15.6$ Hz), 2.24 (1H, dd, $J = 16.8, 7.5$ Hz), 2.20 (1H, dd, $J = 16.8, 8.7$ Hz), 2.04 (1H, dd, $J = 12.9, 6.9$ Hz), 1.96 (1H, ddd, $J = 12.6, 9.0, 4.2$ Hz), 1.24 (1H, ddd, $J = 15.3, 3.0, 3.0$ Hz), 1.12 (1H, dd, $J = 8.4, 8.4$ Hz), 1.01 (9H, t, $J = 7.8$ Hz), 10.1-0.95 (6H, m), 0.95 (9H, s), 0.63 (6H, q, $J = 7.8$ Hz), 0.09 (3H, s), 0.08 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 208.7, 159.6, 137.9, 131.3, 129.5, 119.0, 114.0, 84.1, 82.0, 78.2, 78.1, 76.7, 75.8, 70.2, 65.5, 63.7, 54.8, 47.8, 46.0, 35.6, 34.0, 32.4, 31.4, 26.1, 18.5, 16.6, 16.3, 7.2, 5.5, -5.3 ppm. FTIR (film): 3518, 2953, 2932, 2876, 1709, 1613, 1514, 1463, 1413, 1248, 1092, 1037, 835, 778, 741 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{39}\text{H}_{68}\text{O}_8\text{Si}_2\text{Na}$, 743.4345; found, 743.4414.



[00833] To a stirred solution of the crude ketone (calculated as 0.932 mmol, 1 eq.) in acetone (5.0 mL) were added OsO_4 solution (5.0 mL of 0.02 M in H_2O , 0.100 mmol, 10 mol%) and NMO (220 mg, 1.89 mmol, 2 eq.) at room temperature. After being stirred for 4 h at the same temperature, the reaction was quenched with Na_2SO_3 (1.5 g) and diluted with brine. After being stirred for 30 min at room temperature, the organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure to give a crude diol as a brown oil, which was used in the next step without further purification.

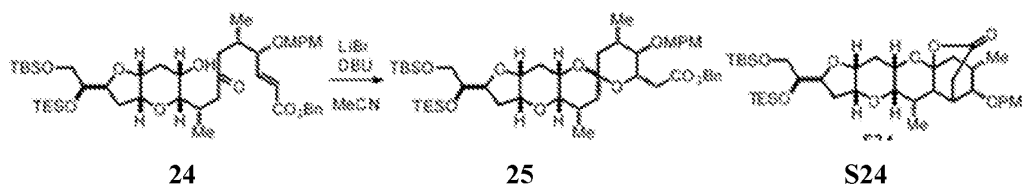
[00834] To a solution of the crude diol (calculated as 0.932 mmol, 1 eq.) in CH_2Cl_2 (19 mL) were added K_2CO_3 (1.3 g, 9.41 mmol, 10 eq.) and $\text{Pb}(\text{OAc})_4$ (620 mg, 1.40 mmol, 1.5 eq.) at room temperature. After being stirred for 15 min at the same temperature, the reaction was diluted with Hexanes/EtOAc (2:1) and filtered through a pad of SiO_2 (33% EtOAc in Hexanes). After evaporation of organic solvent, the obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 5%, 9%, 13%, then 17% EtOAc in

Hexanes) to give hemiacetal **S23** (461 mg, 0.638 mmol, 68% for 3 steps) as pale brown oil. The product was obtained as an equilibrium mixture of hemiacetal (ca. 3:7 ratio in C₆D₆). **S23**: [α]_D²⁰ -35.8 (*c* 1.04, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.17-7.15 (2H, m), 6.80-6.74 (2H, m), 5.44 (0.3H, dd, *J* = 11.7, 1.5 Hz), 4.92 (0.7H, dd, *J* = 12.6, 1.5 Hz), 4.52 (0.7H, d, *J* = 11.1 Hz), 4.45-4.39 (1.3H, m), 4.32 (0.7H, d, *J* = 11.1 Hz), 4.23 (0.3H, d, *J* = 11.4 Hz), 4.09 (0.3H, d, *J* = 3.0 Hz), 3.95 (0.3H, d, *J* = 10.8 Hz), 3.83-3.80 (1H, m), 3.77-3.74 (1H, m), 3.69-3.60 (3H, m), 3.56 (0.7H, d, *J* = 4.2 Hz), 3.33 (0.3H, s), 3.29-3.27 (4H, m), 3.16 (0.7H, s), 2.74-2.72 (1H, m), 2.62-2.56 (0.3H, m), 2.35-2.25 (1.7H, m), 2.18 (0.7H, d, *J* = 15.0 Hz), 2.10-2.00 (1.3H, m), 1.92-1.80 (1.3H, m), 1.72-1.59 (1.7H, m), 1.54 (0.7H, dd, *J* = 12.6, 4.2 Hz), 1.51-1.48 (1H, m), 1.45-1.38 (1H, m), 1.11-1.07 (9.9H, m), 1.04-0.98 (14.1H, m), 0.77-0.72 (6H, m), 0.121-0.09 (6H, m) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 159.9, 159.7, 131.2, 129.7, 129.5, 114.2, 114.0, 98.9, 98.2, 93.8, 91.3, 80.1, 79.1, 79.0, 78.5, 77.8, 77.7, 76.8, 76.0, 74.1, 73.9, 73.4, 73.1, 72.2, 66.1, 64.8, 64.0, 54.7, 38.4, 38.2, 37.7, 37.3, 36.4, 31.5, 31.3, 29.7, 29.2, 26.2, 26.1, 23.2, 18.6, 17.8, 17.5, 17.44, 17.39, 7.3, 7.2, 5.7, 5.5, -5.1, -5.3 ppm. FTIR (film): 3507, 2953, 2928, 2874, 1514, 1462, 1248, 1090, 1035, 1013, 835, 776, 742 cm⁻¹. HRMS (ESI) *m/z*: [M+NH₄]⁺ calcd for C₃₈H₇₀NO₉Si₂, 740.4584; found, 740.4608.



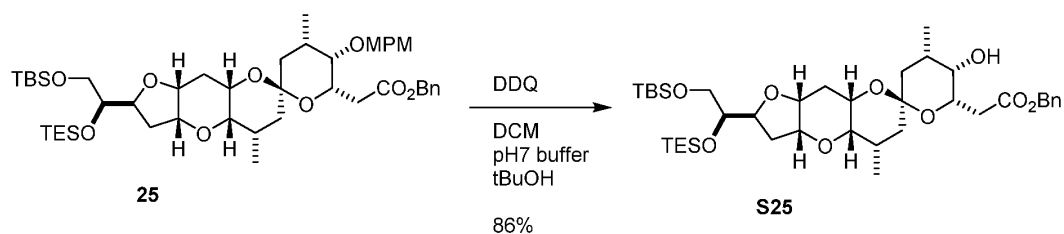
[00835] To a stirred solution of hemiacetal **S23** (461 mg, 0.638 mmol, 1 eq.) and benzyl dimethylphosphonoacetate (0.67 mL, 3.19 mmol, 5 eq.) in THF (26.0 mL) was added NaH (100 mg of 60% in mineral oil, 2.50 mmol, 4 eq.) at 0 °C. After being stirred for 3 h at the same temperature, the reaction was quenched with sat. NH₄Cl aq. The organic layer was separated and the aqueous layer was extracted with Hexanes/EtOAc (1:1). The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 5%, 9%, 13%, then 17% EtOAc in Hexanes) to give unsaturated ester **24** (479 mg, 0.560 mmol, 88%) as colorless oil. **24**: [α]_D²⁰ -21.7 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.24 (2H, d, *J* = 7.2 Hz), 7.15 (2H, d, *J* = 8.4 Hz), 7.10 (2H, dd, *J* = 7.2, 7.2 Hz), 7.05 (1H, t, *J* = 7.2 Hz), 7.04 (1H, dd, *J* = 15.5, 6.3 Hz), 6.79 (2H, d, *J* = 8.4 Hz), 6.20 (1H, d, *J* = 15.5 Hz), 5.14 (1H, d, *J* = 12.0 Hz), 5.10 (1H, d, *J* = 12.0 Hz), 4.48 (1H, ddd, *J* = 8.1, 8.1,

3.4 Hz), 4.38 (1H, d, $J = 12.0$ Hz), 4.10 (1H, d, $J = 12.0$ Hz), 3.83 (1H, dd, $J = 9.9, 6.3$ Hz), 3.77-3.74 (2H, m), 3.63-3.59 (3H, m), 3.55 (1H, ddd, $J = 10.8, 3.0, 3.0$ Hz), 3.35-3.32 (4H, m), 2.70-2.64 (3H, m), 2.54 (1H, dd, $J = 16.8, 4.8$ Hz), 2.47-2.43 (1H, m), 2.29 (1H, d, $J = 15.6$ Hz), 2.15-2.08 (2H, m), 2.04-1.95 (2H, m), 1.25 (1H, ddd, $J = 15.0, 3.6, 3.6$ Hz), 1.10 (1H, dd, $J = 7.8, 7.8$ Hz), 1.01 (9H, t, $J = 8.1$ Hz), 0.95 (9H, s), 0.92 (3H, d, $J = 6.6$ Hz), 0.86 (3H, d, $J = 6.6$ Hz), 0.64 (6H, q, $J = 8.1$ Hz), 0.08 (3H, s), 0.07 (3H, s) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 208.2, 165.6, 159.8, 147.8, 136.6, 130.6, 129.7, 128.70, 128.66, 128.3, 123.3, 114.1, 81.9, 81.5, 78.2, 78.1, 76.7, 75.9, 71.1, 66.4, 65.5, 63.7, 54.7, 47.6, 45.5, 35.6, 33.6, 32.4, 31.4, 26.1, 18.5, 16.4, 16.3, 7.2, 5.5, -5.3 ppm. FTIR (film): 3526, 2953, 2933, 2876, 1719, 1612, 1513, 1462, 1249, 1160, 1093, 1036, 836, 778, 741, 697 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{47}\text{H}_{74}\text{O}_{10}\text{Si}_2\text{Na}$, 877.4713; found, 877.4713.



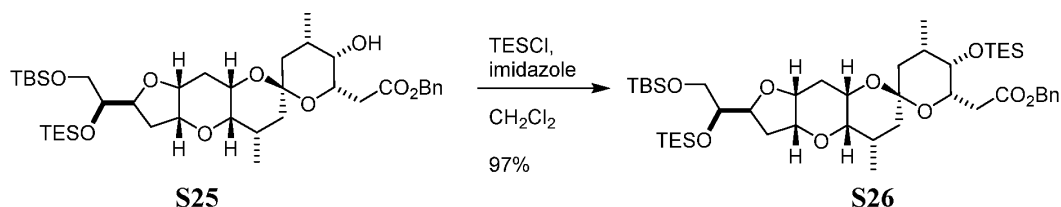
[00836] To a stirred solution of unsaturated ester **24** (132 mg, 0.154 mmol, 1 eq.) in MeCN (3.0 mL) were added LiBr (134 mg, 1.54 mmol, 10 eq.) and DBU (0.46 mL, 3.08 mmol, 20 eq.) at room temperature. After being stirred for 11 h at the same temperature, hexanes (3.0 mL) and H_2O (3.0 mL) were added to the reaction mixture. The organic layer was separated and the aqueous layer was extracted with hexanes. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 5%, 6%, then 9% EtOAc in Hexanes) to give spiro ketal **25** (92.1 mg, 0.108 mmol, 70%) as colorless oil and C-Michael addition product **S24** (6.7 mg, 0.00897 mmol, 6%) as colorless oil. **25**: $[\alpha]_{\text{D}}^{20} -32.5$ (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.23 (2H, d, $J = 7.1$ Hz), 7.19 (2H, d, $J = 8.4$ Hz), 7.11 (2H, dd, $J = 7.5, 7.1$ Hz), 7.06 (1H, dd, $J = 7.5, 7.5$ Hz), 6.76 (2H, d, $J = 8.4$ Hz), 5.09 (1H, d, $J = 12.0$ Hz), 5.00 (1H, d, $J = 12.0$ Hz), 4.44 (1H, ddd, $J = 10.2, 4.8, 4.8$ Hz), 4.31 (1H, d, $J = 10.2$ Hz), 4.23-4.21 (2H, m), 3.83-3.79 (2H, m), 3.75 (dd, $J = 10.2, 6.0$ Hz), 3.72-3.68 (3H, m), 3.30 (3H, s), 2.96 (1H, dd, $J = 15.6, 9.6$ Hz), 2.85 (1H, s), 2.80 (1H, d, $J = 2.4$ Hz), 2.38-2.35 (2H, m), 2.28 (1H, dd, $J = 15.6, 3.6$ Hz), 2.17-2.13 (1H, m), 2.05 (1H, dd, $J = 12.3, 6.3$ Hz), 1.85 (1H, ddd, $J = 13.2, 9.6, 4.2$ Hz), 1.70-1.62 (2H, m), 1.59-1.55 (2H, m), 1.43 (1H, dd, $J = 13.2, 4.2$ Hz), 1.11 (9H, t, $J = 8.1$ Hz), 1.01 (3H, d, $J = 4.2$ Hz), 1.00-0.99 (12H, m), 0.76 (6H, q, $J = 8.1$ Hz), 0.120 (3H, s), 0.116 (3H, s) ppm. ^{13}C NMR (125 MHz,

C₆D₆) δ : 171.6, 159.7, 136.8, 131.3, 129.7, 128.72, 128.66, 114.0, 97.0, 79.1, 78.5, 77.9, 76.8, 75.3, 74.2, 73.5, 69.6, 66.15, 66.06, 63.7, 54.7, 38.2, 37.7, 37.3, 36.5, 31.5, 30.6, 29.3, 26.2, 18.6, 18.2, 17.6, 7.4, 5.7, -5.1, -5.3 ppm. FTIR (film): 2954, 2928, 2874, 1737, 1514, 1462, 1249, 1088, 1018, 836, 742 cm⁻¹. HRMS (ESI) m/z : [M+Na]⁺ calcd for C₄₇H₇₄O₁₀Si₂Na, 877.4713; found, 877.4712. **C-Michael product S25**: [α]_D²⁰ -79.5 (*c* 0.855, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.10 (2H, d, *J* = 8.7 Hz), 6.79 (2H, d, *J* = 8.7 Hz), 4.42 (1H, ddd, *J* = 9.6, 6.2, 3.5 Hz), 4.20 (1H, d, *J* = 12.0 Hz), 4.11 (1H, d, *J* = 12.0 Hz), 3.87 (1H, dd, *J* = 10.2, 6.0 Hz), 3.78 (1H, dd, *J* = 10.2, 5.4 Hz), 3.69-3.66 (3H, m), 3.61 (1H, dd, *J* = 2.7, 2.7 Hz), 3.32 (3H, s), 2.86 (1H, dd, *J* = 3.3, 3.3 Hz), 2.62 (1H, d, *J* = 2.4 Hz), 2.38 (1H, dd, *J* = 12.6, 3.0 Hz), 2.30 (1H, d, *J* = 15.6 Hz), 2.23 (1H, dd, *J* = 19.2, 7.8 Hz), 2.05 (1H, ddd, *J* = 13.6, 3.6, 3.6 Hz), 1.99-1.89 (4H, m), 1.81 (1H, d, *J* = 19.2 Hz), 1.63-1.58 (1H, m), 1.30-1.25 (2H, m), 1.06 (9H, t, *J* = 7.8 Hz), 0.98 (9H, s), 0.84 (3H, d, *J* = 6.6 Hz), 0.82 (3H, d, *J* = 6.0 Hz), 0.70 (6H, q, *J* = 7.8 Hz), 0.114 (3H, s), 0.112 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 169.7, 159.7, 131.1, 129.4, 114.0, 104.9, 80.6, 78.7, 78.2, 76.1, 74.0, 73.7, 71.1, 65.9, 65.2, 54.7, 40.8, 36.0, 33.1, 30.9, 30.7, 30.5, 29.5, 29.4, 26.2, 18.6, 16.9, 13.9, 7.3, 5.6, -5.1, -5.2 ppm. FTIR (film): 2954, 2931, 2876, 1729, 1612, 1514, 1462, 1302, 1247, 1205, 1157, 1133, 1090, 1038, 992, 937, 835, 777, 742 cm⁻¹. HRMS (ESI) m/z : [M+Na]⁺ calcd for C₄₀H₆₆O₉Si₂Na, 769.4138; found, 769.4146.



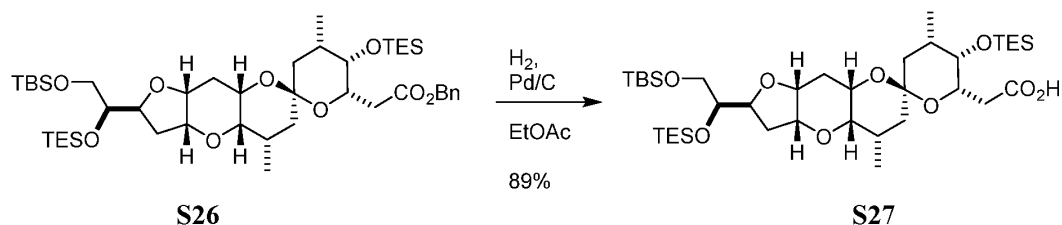
[00837] To a stirred solution of MPM ether **25** (248 mg, 0.290 mmol, 1 eq.) in CH₂Cl₂ (6.0 mL), phosphate buffer (0.60 mL, pH7), and tBuOH (0.60 mL) was added DDQ (200 mg, 0.881 mmol, 3 eq.) at room temperature. After being stirred for 15 min at the same temperature, the reaction was quenched with sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with CH₂Cl₂. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 7%, then 9% EtOAc in Hexanes) to give alcohol **S25** (183 mg, 0.249 mmol, 86%) as a colorless oil. **S25**: [α]_D²⁰ -45.6 (*c* 1.01, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.21 (2H, d, *J* = 7.5 Hz),

7.11 (2H, dd, $J = 7.5, 7.5$ Hz), 7.06 (1H, t, $J = 7.5$ Hz), 5.07 (1H, d, $J = 12.6$ Hz), 4.97 (1H, d, $J = 12.6$ Hz), 4.43 (1H, ddd, $J = 10.2, 5.3, 5.3$ Hz), 4.14 (1H, dd, $J = 9.6, 3.0$ Hz), 3.82 (1H, dd, $J = 9.6, 5.4$ Hz), 3.77-3.74 (2H, m), 3.70-3.67 (3H, m), 2.96 (1H, d, $J = 8.4$ Hz), 2.86 (1H, dd, $J = 15.6, 10.2$ Hz), 2.77 (1H, d, $J = 1.8$ Hz), 2.33-2.28 (2H, m), 2.22-2.17 (1H, m), 2.12-2.04 (2H, m), 1.86 (1H, ddd, $J = 12.8, 10.2, 4.2$ Hz), 1.61 (1H, dd, $J = 12.9, 12.9$ Hz), 1.09 (9H, t, $J = 7.9$ Hz), 1.05 (3H, d, $J = 6.6$ Hz), 0.99 (9H, s), 0.93 (3H, d, $J = 7.2$ Hz), 0.75 (6H, q, $J = 7.9$ Hz), 0.12 (3H, s), 0.12 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 171.5, 136.7, 128.64, 128.57, 97.0, 79.0, 77.9, 76.8, 74.1, 73.3, 70.5, 69.3, 66.09, 66.06, 63.7, 37.4, 37.31, 37.27, 36.4, 30.2, 29.2, 26.2, 18.6, 17.65, 17.62, 7.3, 5.7, -5.1, -5.3 ppm. FTIR (film): 3469, 2953, 2928, 2874, 1736, 1498, 1251, 1128, 1017, 835, 776, 740 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{39}\text{H}_{66}\text{O}_9\text{Si}_2\text{Na}$, 757.4138; found, 757.4135.

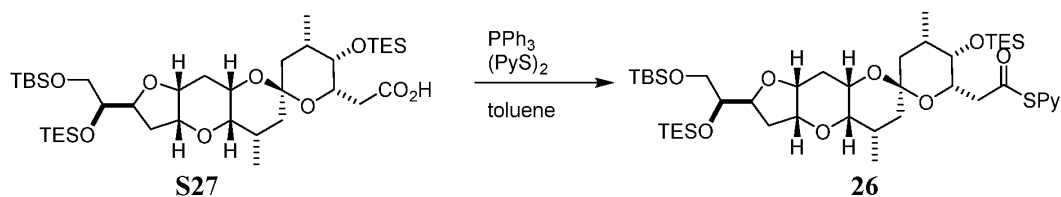


[00838] To a stirred solution of alcohol **S25** (183 mg, 0.249 mmol, 1 eq.) in CH_2Cl_2 (3.0 mL) were added imidazole (50.0 mg, 0.744 mmol, 3 eq.) and TESCl (60 μL , 0.357 mmol, 1.5 eq.) at room temperature. After being stirred for 4 h at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 3%, then 5% EtOAc in Hexanes) to give TES ether **S26** (205 mg, 0.241 mmol, 97%) as a colorless oil. **S26**: $[\alpha]_D^{20}$ -49.1 (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.26 (2H, d, $J = 7.2$ Hz), 7.10 (2H, dd, $J = 7.2, 7.2$ Hz), 7.05 (1H, t, $J = 7.2$ Hz), 5.14 (1H, d, $J = 12.3$ Hz), 5.02 (1H, d, $J = 12.3$ Hz), 4.46 (1H, ddd, $J = 9.8, 5.1, 5.1$ Hz), 4.17 (1H, dd, $J = 10.2, 3.0$ Hz), 3.84-3.80 (2H, m), 3.77 (1H, dd, $J = 10.2, 5.4$ Hz), 3.73 (1H, d, $J = 1.8$ Hz), 3.71-3.69 (2H, m), 3.22 (1H, s), 2.92 (1H, dd, $J = 15.3, 9.1$ Hz), 2.80 (1H, s), 2.38 (1H, d, $J = 15.6$ Hz), 2.30-2.27 (2H, m), 2.16-2.11 (1H, m), 2.06 (1H, dd, $J = 12.9, 6.3$ Hz), 1.87 (1H, ddd, $J = 13.2, 9.6, 4.2$ Hz), 1.69-1.61 (2H, m), 1.59 (1H, ddd, $J = 15.2, 4.7, 4.7$ Hz), 1.53 (1H, dd, $J = 12.6, 3.6$ Hz), 1.46 (1H, dd, $J = 12.3, 3.9$ Hz), 1.11 (9H, t, $J = 7.9$ Hz), 1.00 (9H, s), 1.00-0.95 (12H, m), 0.93 (3H, d, $J = 7.2$ Hz), 0.76 (6H, q, $J = 7.9$ Hz), 0.56 (6H, q, $J = 7.9$ Hz), 0.131 (3H, s), 0.127 (3H, s) ppm. ^{13}C NMR

(125 MHz, C₆D₆) δ : 171.6, 136.8, 128.75, 128.67, 128.3, 96.9, 79.0, 77.9, 76.8, 74.2, 73.6, 72.4, 69.9, 66.2, 66.1, 63.6, 38.3, 37.5, 37.4, 36.4, 31.5, 30.7, 29.3, 26.2, 18.6, 17.5, 7.4, 5.8, 5.7, -5.1, -5.3 ppm. FTIR (film): 2953, 2928, 2875, 1738, 1498, 1240, 1033, 1001, 974, 834, 737, 677 cm⁻¹. HRMS (ESI) m/z : [M+H]⁺ calcd for C₄₅H₈₁O₉Si₃, 849.5183; found, 849.5184.

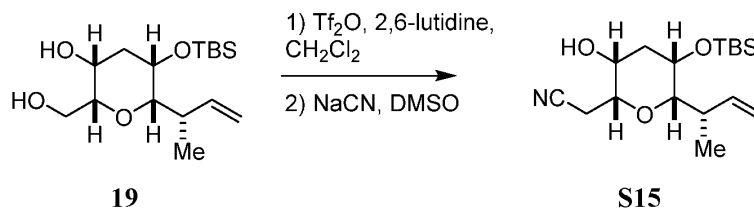


[00839] To a stirred solution of Benzyl ester **S26** (230 mg, 0.271 mmol, 1 eq.) in EtOAc (10.0 mL) was added wet 10% Pd/C (23 mg, 10%w/w) at room temperature. The reaction was stirred under 1 atmosphere of hydrogen for 2 h. The resulting mixture was filtered through a pad of Celite (EtOAc). The organic solvent was removed under reduced pressure to give a crude mixture, which was purified by flash column chromatography on neutral silica gel (0%, 5%, then 33% EtOAc in Hexanes) to give carboxylic acid **S27** (183 mg, 0.241 mmol, 89%) as a colorless oil. **S27**: [α]_D²⁰ -54.8 (*c* 1.06, CHCl₃). ¹H NMR (500 MHz, C₆D₆) δ : 4.45 (1H, ddd, *J* = 10.0, 5.0, 5.0 Hz), 4.12 (1H, dd, *J* = 9.8, 2.8 Hz), 3.89 (1H, d, *J* = 3.5 Hz), 3.81 (1H, dd, *J* = 11.0, 5.0 Hz), 3.76-3.73 (2H, m), 3.70-3.67 (2H, m), 3.24 (1H, s), 2.89 (1H, d, *J* = 1.5 Hz), 2.87 (1H, dd, *J* = 16.0, 3.0 Hz), 2.40 (1H, d, *J* = 15.5 Hz), 2.34 (1H, dd, *J* = 16.0, 3.0 Hz), 2.32-2.21 (2H, m), 2.04 (1H, dd, *J* = 12.8, 5.8 Hz), 1.87 (1H, ddd, *J* = 13.5, 9.5, 4.0 Hz), 1.70-1.62 (3H, m), 1.53 (1H, dd, *J* = 12.5, 4.0 Hz), 1.47 (1H, dd, *J* = 13.0, 4.0 Hz), 1.09 (9H, t, *J* = 8.0 Hz), 1.00 (3H, d, *J* = 7.0 Hz), 0.99 (9H, s), 0.97 (9H, t, *J* = 8.0 Hz), 0.92 (3H, d, *J* = 7.0 Hz), 0.75 (6H, q, *J* = 8.0 Hz), 0.57 (6H, q, *J* = 8.0 Hz), 0.120 (3H, s), 0.115 (3H, s) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 178.1, 97.1, 79.0, 77.9, 76.7, 74.3, 73.6, 72.3, 69.6, 66.1, 63.8, 38.0, 37.5, 37.4, 36.4, 31.5, 30.7, 29.4, 26.2, 18.6, 17.5, 7.3, 5.8, 5.7, -5.1, -5.3 ppm. FTIR (film): 3120, 2953, 2928, 2875, 1731, 1713, 1461, 1415, 1309, 1078, 1033, 1004, 833, 776, 724 cm⁻¹. HRMS (ESI) m/z : [M+H]⁺ calcd for C₃₈H₇₅O₉Si₃, 759.4713; found, 759.4724.



[00840] To a stirred solution of carboxylic acid **S27** (183 mg, 0.241 mmol, 1 eq.) in toluene (1.2 mL) were added PPh_3 (190 mg, 0.724 mmol, 3 eq.) and $(\text{PyS})_2$ (64.0 mg, 0.290 mmol, 1.2 eq.) at room temperature. After being stirred for 12 h at the same temperature, the resulting reaction mixture was directly subjected to column chromatography on neutral silica gel (0%, 3%, 5%, then 6% EtOAc in Hexanes) to give pyridinethiol ester **26** (199 mg, 0.233 mmol, 97%) as pale yellow oil. **26**: $[\alpha]_D^{20}$ -65.7 (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 8.33 (1H, dd, $J = 5.0, 1.4$ Hz), 7.54 (1H, d, $J = 8.1$ Hz), 6.96 (1H, ddd, $J = 8.1, 7.6, 1.4$ Hz), 6.47 (1H, dd, $J = 7.6, 5.0$ Hz), 4.45 (1H, ddd, $J = 9.8, 5.1, 5.1$ Hz), 4.17 (1H, dd, $J = 10.2, 2.4$ Hz), 3.88 (1H, d, $J = 3.6$ Hz), 3.81 (1H, dd, $J = 10.5, 5.1$ Hz), 3.75 (1H, dd, $J = 10.5, 5.7$ Hz), 3.71-3.65 (3H, m), 3.25 (1H, dd, $J = 15.0, 10.2$ Hz), 3.10 (1H, s), 2.82 (1H, d, $J = 1.8$ Hz), 2.53 (1H, dd, $J = 15.0, 2.4$ Hz), 2.39-2.34 (2H, m), 2.27-2.23 (1H, m), 2.04 (1H, dd, $J = 12.9, 5.7$ Hz), 1.85 (1H, ddd, $J = 13.4, 9.8, 3.8$ Hz), 1.70-1.60 (3H, m), 1.54-1.50 (2H, m), 1.10 (9H, t, $J = 7.8$ Hz), 0.99-0.97 (21H, m), 0.91 (3H, d, $J = 6.0$ Hz), 0.75 (6H, q, $J = 8.0$ Hz), 0.56 (6H, q, $J = 7.8$ Hz), 0.121 (3H, s), 0.119 (3H, s) ppm. ^{13}C NMR (125 MHz, C_6D_6) δ : 194.6, 152.8, 150.6, 136.5, 129.8, 123.1, 97.1, 79.0, 77.9, 76.8, 74.2, 73.6, 72.5, 69.9, 66.1, 63.8, 47.8, 37.5, 37.4, 36.4, 31.4, 30.7, 29.3, 26.2, 18.6, 18.5, 17.5, 7.4, 7.3, 5.9, 5.7, -5.1, -5.3 ppm. FTIR (film): 2953, 2928, 2875, 1708, 1572, 1420, 1250, 1033, 1001, 834, 775, 736, 723 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{43}\text{H}_{77}\text{NO}_8\text{SSi}_3\text{Na}$, 874.4570; found, 874.4573.

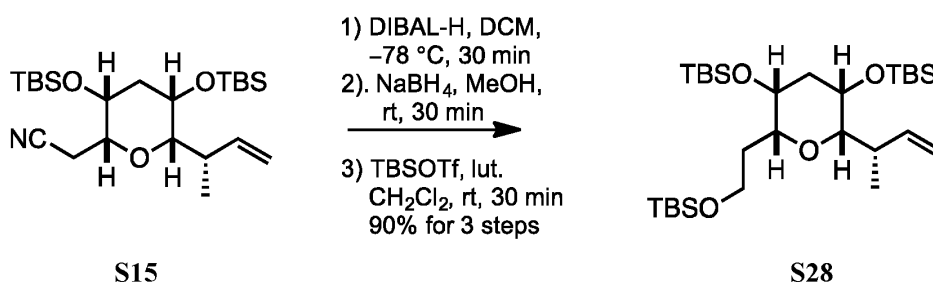
Norhalichondrin Left Halves



[00841] To a stirred solution of diol **19** (3.65 g, 11.6 mmol, 1 eq.) and 2,6-lutidine (5.4 mL, 46.4 mmol, 4 eq.) in CH_2Cl_2 (75 mL) was added Tf_2O (2.4 mL, 27.9 mmol, 1.2 eq.) at -78°C . After being stirred for 10 min at the same temperature, the reaction was quenched with

MeOH (0.5 mL) and brine (50 mL). After being added Et₂O (250 mL), the organic layer was separated from aqueous layer and washed with 1N HCl and brine sequentially. The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure to give a crude triflate as wine red oil, which was dissolved in DMSO (75 mL) immediately without further purification.

[00842] To the DMSO solution of crude triflate (estimated as 11.6 mmol, 1 eq.) was added NaCN (5.7 g, 115 mmol, 10 eq.) at room temperature. After being stirred for 1 hr at the same temperature, the reaction mixture was filtered through a paper and the filter cake was washed with EtOAc thoroughly. The filtrate was washed with H₂O and brine. The organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 5%, 10%, then 25% EtOAc in Hexanes) to give **S15** (3.26 g, 10.1 mmol, 87% for 2 steps) as a colorless oil. **S15**: [α]_D²⁰ -24.0 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 6.18 (1H, m), 5.15 (1H, d, *J* = 9.6 Hz), 5.11 (1H, d, *J* = 15.4 Hz), 3.48 (1H, s), 3.39 (1H, d, *J* = 11.4 Hz), 3.21 (1H, d, *J* = 11.4 Hz), 2.97 (1H, dd, *J* = 6.6, 6.4 Hz), 2.61-2.55 (2H, m), 2.33 (1H, dd, *J* = 16.2, 8.4 Hz), 2.09 (1H, dd, *J* = 15.0, 7.8 Hz), 1.88 (1H, d, *J* = 15.2 Hz), 0.99 (1H, d, *J* = 12.6 Hz), 0.88 (9H, s), 0.80 (3H, d, *J* = 5.8 Hz), 0.03 (3H, s), -0.09 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 141.2, 117.6, 114.1, 84.7, 76.9, 66.4, 65.8, 37.5, 36.3, 25.8, 20.4, 18.1, 15.3, -4.0, -5.1 ppm. FTIR (film): 3515, 2956, 2930, 2897, 2858, 1473, 1433, 1365, 1255, 1168, 1122, 1082, 1046, 995, 978, 940, 834, 775, 736, 688, 474 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₁₇H₃₁NO₃SiNa, 348.1965; found, 348.1969.

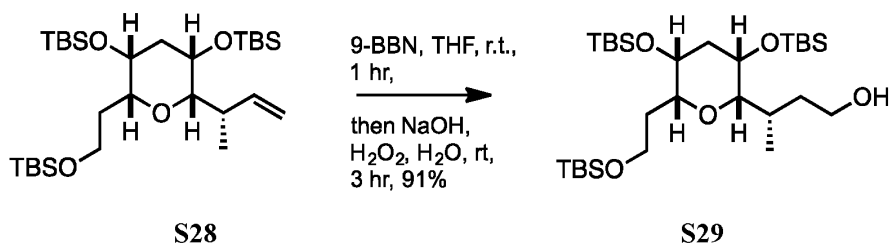


[00843] To a stirred solution of **S15** (3.25 g, 10.0 mmol, 1 eq.) in CH₂Cl₂ (150 mL) were added DIBAL solution (45 ml of 1 M in hexanes, 45.0 mmol, 4.5 eq.) at -78 °C under Ar atmosphere. After being stirred for 30 min at the same temperature, 150 mL of sat. Rochelle's salt aq. was added and the temperature was allowed to increase to room temperature. After being stirred for 1 h at the same temperature, the mixture was diluted with EtOAc. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined

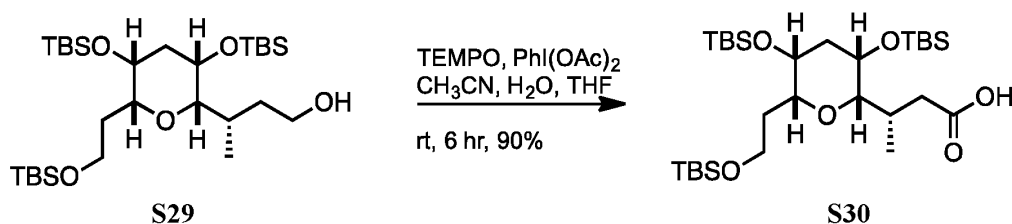
organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure to give a crude aldehyde, which was used in the next reaction without further purification.

[00844] To a stirred solution of the crude aldehyde (calculated as 10.0 mmol, 1 eq.) in CH₂Cl₂ (150 mL) were added NaBH₄ (1.89 g, 50.0 mmol, 5 eq.) at 0 °C. After being stirred for 30 min at room temperature, the reaction was quenched with AcOH (2.8 mL) and the resultant mixture was concentrated under reduced pressure. The residue was passed through a pad of silica gel (5% MeOH in CH₂Cl₂) to give a crude alcohol, which was used in the next reaction after concentration without further purification.

[00845] To a stirred solution of the crude alcohol (calculated as 10.0 mmol, 1 eq.) in CH₂Cl₂ (100 mL) were added 2,6-lutidine (3.75 g, 35.0 mmol, 3.5 eq.) and TESOTf (7.93 g, 30.0 mmol, 3 eq.) at 0 °C. After being stirred for 30 min at room temperature, MeOH (1 mL) was added and the resultant mixture was diluted with 200 mL of CH₂Cl₂. The resultant mixture was washed with 1N HCl and sat. NaHCO₃ aq. The organic layer was dried over MgSO₄, filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 5%, 10%, then 15% EtOAc in Hexanes) to give **S28** (5.03 g, 9.00 mmol, 90% for 3 steps) as a colorless oil. **S28**: $[\alpha]_D^{20} +3.4$ (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 6.27 (1H, m), 5.17 (1H, ddd, *J* = 17.4, 1.8, 1.6 Hz), 5.16 (1H, dd, *J* = 10.8, 1.8 Hz), 4.00 (1H, m), 3.82 (1H, m), 3.67 (1H, quin, *J* = 2.0 Hz), 3.49 (1H, ddd, *J* = 10.8, 1.8, 1.6 Hz), 2.89 (1H, m), 2.84 (1H, dd, *J* = 9.0, 1.8 Hz), 2.18 (1H, m), 1.94 (1H, ddd, *J* = 15.8, 2.3, 2.0 Hz), 1.65 (1H, m), 1.54 (1H, dt, *J* = 15.6, 4.4 Hz), 1.03 (9H, s), 1.022 (9H, s), 1.020 (9H, s), 0.98 (3H, d, *J* = 6.6 Hz), 0.17 (3H, s), 0.15 (3H, s), 0.12 (3H, s), 0.08 (3H, s), 0.06 (3H, s), 0.02 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 143.0, 113.4, 85.1, 77.4, 66.3, 65.0, 60.3, 39.3, 37.8, 36.1, 18.6, 18.5, 16.1, -2.2, -3.4, -4.7, -5.00, -5.02, -5.10 ppm. FTIR (film): 2953, 2928, 2844, 2856, 1472, 1462, 1387, 1251, 1020, 958, 938, 886, 808, 768, 705, 674, 662 cm⁻¹. HRMS (ESI) *m/z*: [M+H]⁺ calcd for C₂₉H₆₃O₄Si₃, 559.4034; found, 559.4063.

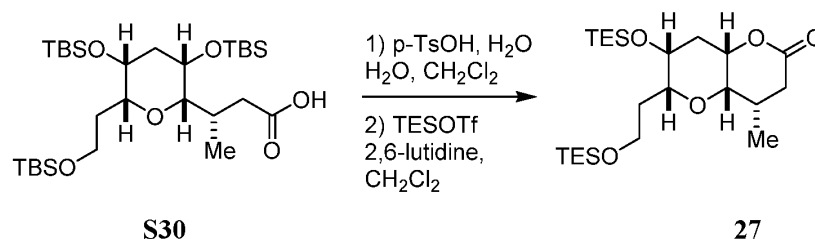


[00846] To a stirred solution of the **S28** (4.9 g, 8.78 mmol, 1 eq.) in THF (16 mL) was added 9-BBN solution (17.6 mL of 0.5 M in THF, 8.8 mmol, 2 eq.) at 0 °C. After being stirred for 1 h at room temperature, 2 N NaOH aq. (21 mL) was added, followed by 9 M H₂O₂ aq. very slowly at 0 °C. After being stirred for 30 min at room temperature, the resultant mixture was quenched with sat. Na₂S₂O₃ aq. The organic layer was separated and the aqueous layer was extracted with Et₂O. The combined organic layer was dried over MgSO₄, filtered, and concentrated under reduced pressure. The crude product was purified by flash column chromatography on silica gel (0%, 5%, 10%, then 20% EtOAc in Hexanes) to give **S29** (4.61 g, 7.99 mmol, 91%) as a white solid.



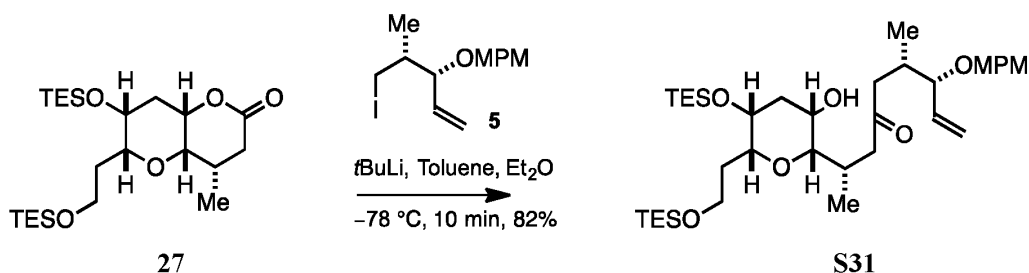
[00847] To a stirred solution of alcohol **S29** (1.6 g, 2.81 mmol, 1 eq.) in THF/CH₃CN/H₂O (3.5 mL/13 mL/13 mL) were added TEMPO (220 mg, 1.41 mmol, 0.5 eq.) and PhI(OAc)₂ (4.53 g, 14.1 mmol, 5 eq.) at room temperature. After being stirred for 5 h at the same temperature, the resultant mixture was quenched with sat. Na₂S₂O₃ aq. The organic layer was separated and the aqueous layer was extracted with Et₂O. The combined organic layer was dried over MgSO₄, filtered, and concentrated under reduced pressure. The crude product was purified by flash column chromatography on silica gel (0%, 5%, 10%, then 25% EtOAc in Hexanes) to give acid **S30** (1.49 g, 2.53 mmol, 90%) as an amorphous solid. **S30**: [α]_D²⁰ +34.0 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ: 3.90 (1H, td, *J* = 9.6, 4.2 Hz), 3.79 (1H, m), 3.61 (1H, s), 3.46 (1H, d, *J* = 10.8 Hz), 3.40 (1H, s), 3.01 (1H, dd, *J* = 16.2, 4.2 Hz), 2.82 (1H, d, *J* = 4.2 Hz), 2.66 (1H, m), 2.35 (1H, dd, *J* = 16.2, 7.2 Hz), 2.15 (1H, m), 1.89 (1H, d, *J* = 14.4 Hz), 1.65 (1H, m), 1.49 (1H, dt, *J* = 14.4, 4.9 Hz), 1.04 (9H, s), 1.03 (9H, s), 1.02 (9H, s), 0.96 (3H, d, *J* = 7.2 Hz), 0.15 (3H, s), 0.14 (3H, s), 0.13 (3H, s), 0.08 (3H, s), 0.06 (3H, s), 0.01 (3H, s) ppm. ¹³C NMR (150 MHz, C₆D₆) δ: 180.2, 84.1, 77.6, 67.3, 64.9, 60.6,

39.4, 37.8, 36.0, 30.8, 26.7, 26.5, 26.3, 18.6, 18.5, 18.4, 16.8, -2.6, -3.4, -4.8, -5.0, -5.1, -5.2 ppm. FTIR (film): 2953, 2929, 2885, 2857, 1708, 1472, 1463, 1386, 1253, 1098, 1020, 938, 886, 772 cm^{-1} . HRMS (ESI) m/z : $[M+H]^+$ calcd for $\text{C}_{29}\text{H}_{63}\text{O}_6\text{Si}_3$, 591.3927; found, 591.3921.

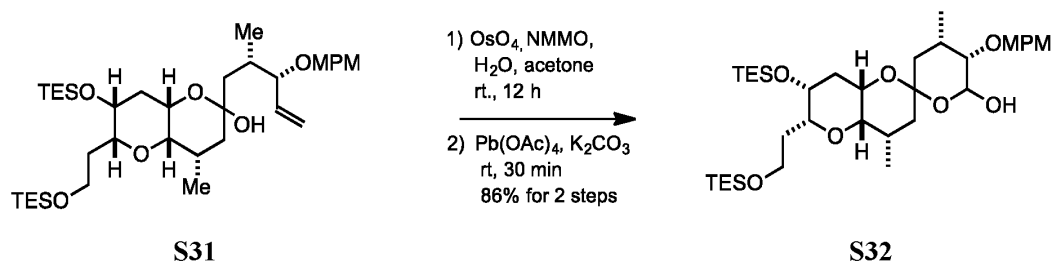


[00848] To a stirred solution of acid **S30** (2.50 g, 4.23 mmol, 1 eq.) in CH_2Cl_2 (40 mL) were added $p\text{-TsOH}\cdot\text{H}_2\text{O}$ (805 mg, 4.23 mmol, 1 eq.) and H_2O (0.76 mL, 42.3 mmol, 10 eq.) at room temperature. After being stirred for 24 h at the same temperature, the resultant mixture was directly concentrated under reduced pressure to give a crude lactone. The crude product was used in the next step without any purification.

[00849] To a stirred solution of the crude lactone (calculated as 4.23 mmol, 1 eq.) in CH_2Cl_2 (100 mL) were added 2,6-lutidine (5.43 g, 50.8 mmol, 12 eq.) and TESOTf (11.2 g, 42.3 mmol, 10 eq.) at 0 °C. After being stirred for 1 h at room temperature, MeOH (3 mL) was added and the resultant mixture was diluted with 200 mL of CH_2Cl_2 . The resultant mixture was washed with sat. NH_4Cl aq. and sat. NaHCO_3 aq. The organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 5%, 10%, then 15% EtOAc in Hexanes) to give bis-TES **27** (1.46 g, 3.20 mmol, 76% for 2 steps) as a colorless oil. **27**: $[\alpha]_D^{20} +25.5$ (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 3.80 (1H, td, $J = 9.6, 4.4$ Hz), 3.70 (1H, m), 3.61 (1H, s), 3.37 (1H, dd, $J = 10.2, 3.0$ Hz), 3.33 (1H, s), 2.82 (1H, s), 2.46 (1H, dd, $J = 17.2, 12.0$ Hz), 2.13 (1H, dd, $J = 17.2, 4.6$ Hz), 2.09-2.02 (2H, m), 1.67 (1H, m), 1.32 (1H, m), 1.20 (1H, dt, $J = 14.4, 4.5$ Hz), 1.09-1.01 (18H, m), 0.79 (3H, d, $J = 6.6$ Hz), 0.75-0.61 (12H, m) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 168.5, 75.6, 74.3, 73.4, 66.2, 59.2, 36.3, 36.0, 33.0, 31.4, 26.2, 16.6, 7.2, 5.3, 4.8 ppm. FTIR (film): 2954, 2934, 2911, 2876, 1734, 1459, 1414, 1376, 1239, 1197, 1097, 1032, 938, 780, 726 cm^{-1} . HRMS (ESI) m/z : $[M+H]^+$ calcd for $\text{C}_{23}\text{H}_{47}\text{O}_5\text{Si}_2$, 459.2957; found, 459.2946.

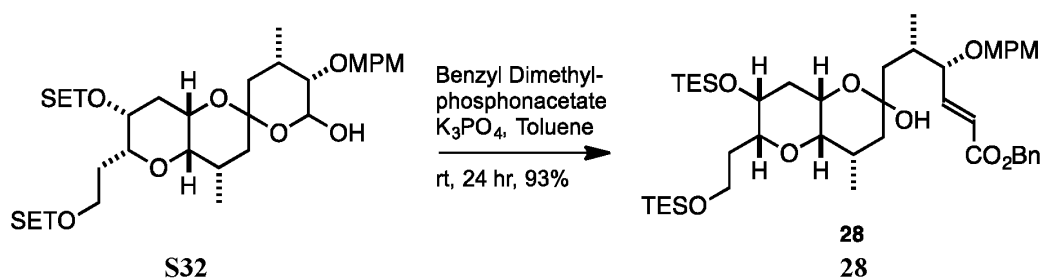


[00850] To a stirred solution of the iodide **5** (1.21 g, 3.50 mmol, 1.3 eq.) in Et₂O (30 mL, bubbled with Ar gas) were added *t*-BuLi solution (3.5 mL of 1.7 M in pentane, 5.92 mmol, 2.2 eq.) at -78 °C under Ar atmosphere. After being stirred for 5 min at the same temperature, the resultant mixture was transferred via cannula to a stirred solution of the lactone **27** (1.23 g, 2.69 mmol, 1 eq.) in toluene (30 mL, bubbled with Ar gas) at -78 °C under Ar atmosphere. After being stirred for 10 min at the same temperature, the resultant mixture was quenched with sat. NaHCO₃ aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The crude product was purified by flash column chromatography on neutral silica gel (0%, 5%, 10%, then 15% EtOAc in Hexanes) to give ketone/hemiketal mixture **S31** (1.48g, 2.19 mmol, 82%) as a colorless oil. **S31**: $[\alpha]^{20}_{\text{D}} +2.8$ (*c* 1.00, CHCl₃). ¹H NMR (500 MHz, C₆D₆) δ : 7.27 (2H, d, *J* = 9.6 Hz), 6.85 (2H, d, *J* = 9.6 Hz), 5.64 (1H, m), 5.16-5.05 (2H, m), 4.55 (1H, d, *J* = 13.8 Hz), 4.23 (1H, d, *J* = 13.8 Hz), 3.94-3.87 (1H, m), 3.84-3.78 (1H, m), 3.70-3.61 (2H, m), 3.49 (1H, t, *J* = 7.8 Hz), 3.44 (1H, d, *J* = 11.4 Hz), 3.40 (1H, s), 3.34 (3H, s), 2.98 (1H, dd, *J* = 18.0, 3.6 Hz), 2.91-2.85 (1H, m), 2.81 (1H, d, *J* = 10.8 Hz), 2.74 (1H, dd, *J* = 18.0, 3.6 Hz), 2.55-2.48 (1H, m), 2.30-2.18 (2H, m), 2.13-2.05 (1H, m), 2.02 (1H, dt, *J* = 17.4, 2.0 Hz), 1.68-1.60 (1H, m), 1.28 (1H, dt, *J* = 17.4, 2.0 Hz), 1.08 (9H, t, *J* = 8.4 Hz), 0.99 (3H, d, *J* = 7.8 Hz), 0.97 (3H, d, *J* = 7.8 Hz), 0.93 (9H, t, *J* = 8.4 Hz), 0.70 (6H, q, *J* = 8.4 Hz), 0.51 (6H, q, *J* = 8.4 Hz) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 208.9, 159.7, 137.9, 131.4, 129.5, 117.9, 114.0, 84.6, 84.0, 77.0, 70.2, 69.7, 65.0, 59.7, 54.8, 47.9, 46.0, 37.7, 36.5, 34.0, 31.3, 16.5, 16.2, 7.2, 7.0, 5.0, 4.9 ppm. FTIR (film): 3523, 2955, 2935, 2912, 2876, 1711, 1613, 1514, 1459, 1413, 1376, 1301, 1247, 1085, 1003, 819, 743 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₃₇H₆₆O₇Si₂Na, 701.4245; found, 701.4268.

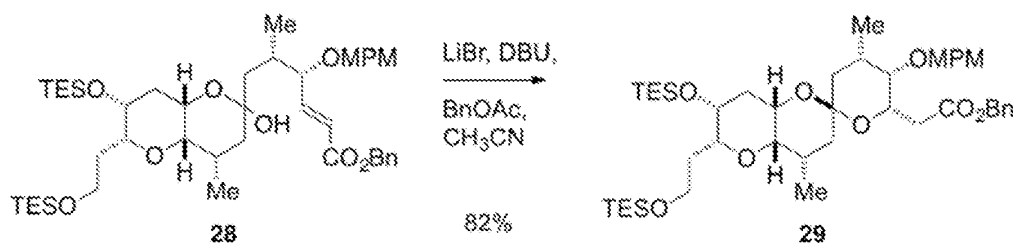


[00851] To a stirred solution of **S31** (1.17 g, 1.73 mmol, 1 eq.) in acetone (17.3 mL) were added NMMO (405 mg, 3.46 mmol, 2 eq.) and OsO₄ solution (4.4 mL of 0.02 M in H₂O, 0.088 mmol, 5 mol%) at room temperature. After being stirred for 12 h at the same temperature, the mixture was diluted with water. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The residue was passed through a pad of silica gel (EtOAc) and concentrated under reduced pressure to give a crude diol, which was used in the next reaction without further purification.

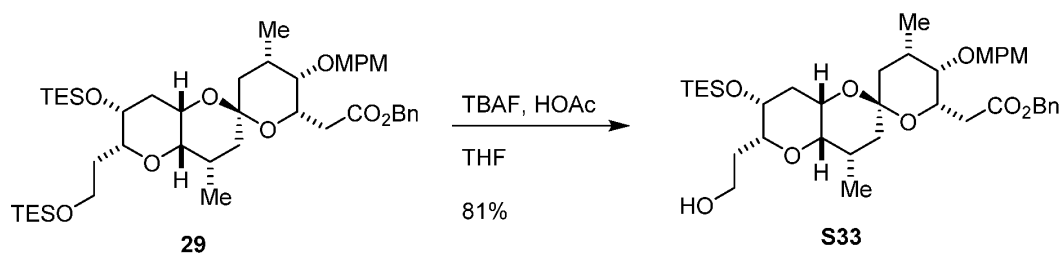
[00852] To a stirred solution of diol (calculated as 1.73 mmol, 1 eq.) in CH₂Cl₂ (17.3 mL) was added K₂CO₃ (2.39 g, 17.3 mmol, 10 eq.) and Pb(OAc)₄ (1.53 g, 3.46 mmol, 2 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction mixture was passed through a pad of silica gel (EtOAc). The filtrate was concentrated under reduced pressure. The residue was purified by flash column chromatography on silica gel (0%, 9%, 17%, then 25% EtOAc in Hexanes) to give **S32** (1.0 g, 1.48 mmol, 86% for 2 steps) as colorless oil. **S32** was obtained as an equilibrium mixture of hemiacetal. **S32**: [α]_D²⁰ -20.0 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.16 (2H, d, *J* = 8.4 Hz), 6.76 (2H, d, *J* = 8.4 Hz), 5.44 (0.28H, d, *J* = 9.6 Hz), 4.91 (0.72H, d, *J* = 9.6 Hz), 4.54 (0.72H, d, *J* = 9.6 Hz), 4.46 (0.28H, d, *J* = 9.6 Hz), 4.36 (0.72H, d, *J* = 9.6 Hz), 4.26 (0.28H, d, *J* = 9.6 Hz), 4.11 (0.28H, s), 3.98-3.86 (1.28H, m), 3.85-3.76 (1H, m), 3.60 (0.78H, s), 3.46-3.36 (2H, m), 3.30 (0.84H, s), 3.29 (2.16H, s), 3.22 (0.28H, s), 3.20 (0.72H, s), 2.99 (0.28H, s), 2.96 (0.72H, s), 2.64 (0.28H, m), 2.34 (1.72H, m), 2.23-2.11 (1H, m), 1.92-1.76 (1.28H, m), 1.76-1.60 (3.27H, m), 1.54 (1H, m), 1.50-1.40 (2H, m), 1.15-0.98 (24H, m), 0.74-0.54 (12H, m) ppm. ¹³C NMR (125 MHz, C₆D₆) δ : 159.9, 131.2, 129.7, 129.6, 114.3, 114.0, 99.0, 98.2, 93.9, 91.3, 80.1, 78.5, 76.4, 76.3, 76.0, 75.9, 75.8, 72.1, 67.0, 66.9, 66.0, 65.2, 59.6, 59.5, 54.7, 38.5, 38.4, 37.8, 37.7, 36.6, 36.4, 30.1, 29.0, 23.5, 17.8, 17.5, 17.3, 7.3, 7.2, 5.5, 5.4, 4.8 ppm. FTIR (film): 2953, 2931, 2910, 2874, 1514, 1459, 1247, 1208, 1192, 1166, 1134, 1095, 1029, 1007, 967, 819, 742, 726 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₃₆H₆₄O₈Si₂Na, 703.4037; found, 703.4047.



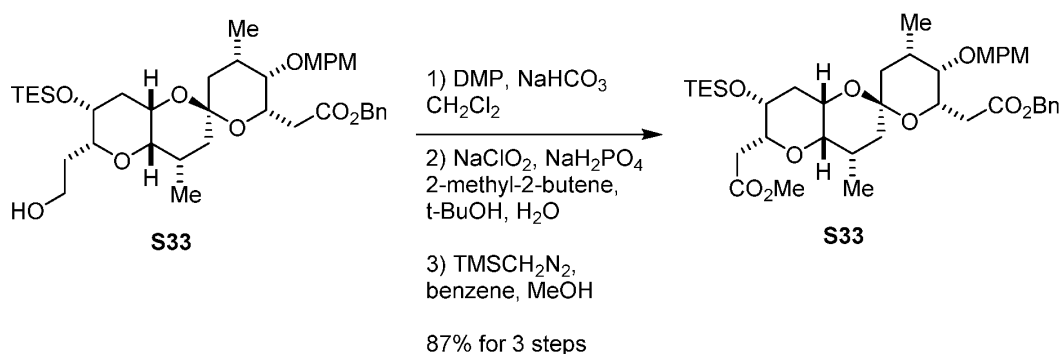
[00853] To a stirred solution of **S32** (700 mg, 1.029 mmol, 1 eq.) in toluene (10 mL) were added benzyl dimethylphosphonoacetate (0.86 mL, 4.116 mmol, 4 eq.) and K_3PO_4 (660 mg, 3.087 mmol, 3 eq.) at room temperature. After being stirred for 36 h at the same temperature, the reaction mixture was passed through a pad of silica gel (50% EtOAc in Hexanes) to give crude **28**, which was further purified by flash column chromatography on neutral silica gel (0%, 5%, 6%, then 9% EtOAc in Hexanes) to give a ~8:1 mixture of *E/Z* isomers **28** (776 mg, 0.956 mmol, 93%) as a colorless oil. Some of the pure *E* isomer was isolated for characterization purpose by further chromatography. The Mixture of *E/Z* isomer was directly used in the next step. **28**: $[\alpha]_D^{20}$ -2.2 (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.24 (2H, d, *J* = 7.8 Hz), 7.15 (2H, d, *J* = 7.8 Hz), 7.11 (2H, t, *J* = 7.8 Hz), 7.05 (1H, m), 6.79 (2H, d, *J* = 7.8 Hz), 6.19 (1H, d, *J* = 15.6 Hz), 5.12 (2H, q, *J* = 12.0 Hz), 4.37 (1H, d, *J* = 12.0 Hz), 4.11 (1H, d, *J* = 12.0 Hz), 3.92-3.87 (1H, m), 3.83-3.78 (1H, m), 3.68-3.57 (3H, m), 3.46-3.41 (2H, m), 3.33 (3H, s), 2.91 (1H, dd, *J* = 15.0, 2.0 Hz), 2.85-2.78 (2H, m), 2.52 (1H, dd, *J* = 17.4, 4.5 Hz), 2.49-2.44 (1H, m), 2.21-2.10 (2H, m), 2.08 (1H, td, *J* = 10.2, 6.0 Hz), 2.03 (1H, d, *J* = 14.4 Hz), 1.68-1.61 (1H, m), 1.29 (1H, d, *J* = 14.4 Hz), 1.07 (9H, t, *J* = 8.4 Hz), 0.97 (3H, d, *J* = 7.8 Hz), 0.93 (9H, t, *J* = 8.4 Hz), 0.86 (3H, d, *J* = 7.8 Hz), 0.69 (6H, q, *J* = 8.4 Hz), 0.52 (6H, q, *J* = 8.4 Hz) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 208.3, 165.6, 159.8, 147.8, 136.7, 130.7, 129.7, 128.7, 128.6, 128.3, 123.3, 114.0, 84.5, 81.6, 76.9, 71.2, 69.6, 66.4, 65.0, 59.6, 54.8, 47.8, 45.6, 37.8, 36.5, 33.7, 31.2, 16.3, 16.2, 7.2, 7.0, 5.1, 4.9 ppm. FTIR (film): 3511, 2954, 2912, 2876, 1720, 1514, 1458, 1413, 1376, 1248, 1165, 1085, 1004, 820, 743, 729, 698 cm^{-1} . HRMS (ESI) *m/z*: $[\text{M}+\text{Na}]^+$ calcd for $\text{C}_{45}\text{H}_{72}\text{O}_9\text{Si}_2\text{Na}$, 835.4613; found, 835.4622.



[00854] To a stirred solution of unsaturated ester **28** (800 mg, 0.985 mmol, 1 eq.) in MeCN (3.0 mL) were added BnOAc (296 mg, 1.97 mmol, 2 eq.), LiBr (857 mg, 9.85 mmol, 10 eq.), and DBU (0.74 mL, 4.93 mmol, 5 eq.) at room temperature. After being stirred for 12 h at the same temperature, hexanes (3.0 mL) and H₂O (3.0 mL) were added to the reaction mixture. The organic layer was separated and the aqueous layer was extracted with hexanes. The combined organic layer was dried over Na₂SO₄, filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 5%, 6%, then 9% EtOAc in Hexanes) to give spiro ketal **29** (658 mg, 0.810 mmol, 82%) as colorless oil. **29**: $[\alpha]_D^{20}$ -28.1 (*c* 1.00, CHCl₃). ¹H NMR (600 MHz, C₆D₆) δ : 7.24 (2H, d, *J* = 7.8 Hz), 7.20 (2H, d, *J* = 7.8 Hz), 7.13 (2H, t, *J* = 7.8 Hz), 7.09 (1H, q, *J* = 7.8 Hz), 6.77 (2H, d, *J* = 7.8 Hz), 5.09 (1H, d, *J* = 14.4 Hz), 5.00 (1H, d, *J* = 14.4 Hz), 4.33 (1H, d, *J* = 11.0 Hz), 4.28-4.21 (2H, m), 3.94 (1H, m), 3.88-3.78 (2H, m), 3.48 (1H, d, *J* = 10.8 Hz), 3.42 (1H, m), 3.30 (3H, s), 3.04 (1H, s), 2.97 (1H, dd, *J* = 14.4, 10.8 Hz), 2.90 (1H, s), 2.38 (1H, m), 2.31 (1H, dd, *J* = 17.0, 2.5 Hz), 2.26-2.16 (2H, m), 2.04 (1H, d, *J* = 15.0 Hz), 1.77-1.64 (3H, m), 1.61-1.55 (2H, m), 1.42 (1H, dd, *J* = 12.6, 3.1 Hz), 1.12-1.04 (21H, m), 1.02 (3H, d, *J* = 6.6 Hz), 0.67 (12H, q, *J* = 8.4 Hz) ppm. ¹³C NMR (150 MHz, C₆D₆) δ : 171.7, 159.7, 136.8, 131.3, 129.7, 128.8, 128.6, 128.3, 114.0, 97.0, 78.6, 76.4, 76.2, 75.3, 69.8, 67.2, 66.1, 64.9, 59.6, 54.7, 38.2, 37.7, 37.4, 36.7, 36.4, 30.9, 29.2, 18.2, 17.4, 7.3, 7.2, 5.5, 4.9 ppm. FTIR (film): 2953, 2978, 2911, 2874, 1736, 1613, 1514, 1458, 1414, 1396, 1371, 1303, 1246, 11246, 1207, 1190, 1165, 1128, 1095, 1032, 1010, 974, 822, 741, 727, 698, 671 cm⁻¹. HRMS (ESI) *m/z*: [M+Na]⁺ calcd for C₄₅H₇₂O₉Si₂Na, 835.4613; found, 835.4598.

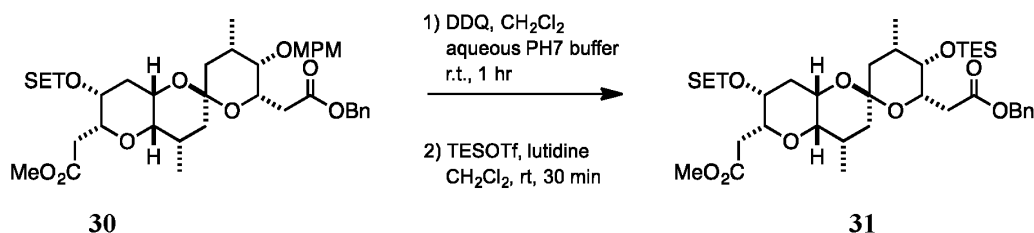


[00855] To a stirred solution of TES ether **29** (600 mg 0.738 mmol, 1 eq.) in CH_2Cl_2 (5 mL) was added TBAF (1.107 mL, 1 M solution in THF, 1.107 mmol, 1.5 eq.) at 0 °C. After being stirred for 5 hr at the same temperature, without concentration, the reaction mixture was directly subjected to the flash column chromatography on neutral silica gel (0%, 5%, then 10% EtOAc in Hexanes) to give ester **S33** (418 mg, 0.598 mmol, 81%) as a colorless oil. **S33**: $[\alpha]_D^{20} -46.0$ (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.24 (2H, d, $J = 7.8$ Hz), 7.20 (2H, d, $J = 7.8$ Hz), 7.13 (2H, t, $J = 7.8$ Hz), 7.09 (1H, q, $J = 7.8$ Hz), 6.77 (2H, d, $J = 7.8$ Hz), 5.09 (1H, d, $J = 14.4$ Hz), 5.00 (1H, d, $J = 14.4$ Hz), 4.33 (1H, d, $J = 11.0$ Hz), 4.28-4.21 (2H, m), 3.85-3.79 (2H, m), 3.77-3.72 (1H, m), 3.29 (3H, s), 3.21 (1H, d, $J = 8.4$ Hz), 2.99-2.93 (2H, m), 2.88 (1H, s), 2.36 (1H, s), 2.28 (1H, dd, $J = 16.2, 3.0$ Hz), 2.24 (1H, m), 2.16-2.08 (2H, m), 2.01 (1H, d, $J = 13.8$ Hz), 1.69-1.54 (3H, m), 1.50 (1H, dt, $J = 13.8, 2.1$ Hz), 1.36 (2H, dd, $J = 12.6, 3.0$ Hz), 1.08 (9H, t, $J = 8.4$ Hz), 1.03 (3H, d, $J = 7.2$ Hz), 0.97 (3H, d, $J = 7.2$ Hz), 0.65 (6H, q, $J = 8.4$ Hz) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 171.7, 159.8, 136.8, 131.2, 129.7, 128.7, 128.6, 128.3, 114.0, 97.0, 80.1, 78.5, 76.6, 75.3, 69.8, 66.9, 66.1, 64.6, 61.3, 54.8, 38.1, 37.7, 37.3, 36.4, 35.1, 30.9, 28.9, 18.2, 17.6, 7.3, 5.5 ppm. FTIR (film): 2954, 2928, 2911, 2874, 1736, 1612, 1514, 1457, 1420, 1304, 1247, 1208, 1163, 1086, 1064, 1038, 1010, 821, 740, 699 cm^{-1} . HRMS (ESI) m/z : $[\text{M}+\text{H}]^+$ calcd for $\text{C}_{39}\text{H}_{59}\text{O}_9\text{Si}$, 699.3923; found, 699.3937.



[00856] To a stirred solution of alcohol **S33** (400 mg, 0.573 mmol, 1 eq.) in CH_2Cl_2 (10 mL) were added NaHCO_3 (480 mg, 5.73 mmol, 10 eq.) and Dess-Martin periodinane (486 mg, 1.146 mmol, 2 eq.) subsequently at room temperature. After being stirred for 30 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was passed through a SiO_2 to give the crude aldehyde which was used in the next step without further purification. To a stirred solution of the crude aldehyde (estimated as 0.573 mmol, 1 eq.) in a mixed solvent (9 mL of *t*-BuOH, 2 mL of water and 2 mL of 2-methyl-2-butene) were added $\text{NaH}_2\text{PO}_3 \cdot \text{H}_2\text{O}$ (316 mg, 2.292 mmol, 4 eq.) and NaClO_2 (156 mg, 1.719 mmol, 3 eq.) subsequently at room temperature. After being stirred for 30 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude acid was used in the next step without further purification. To a stirred solution of the crude acid (estimated as 0.573 mmol, 1 eq.) in a mixed solvent (15 mL of benzene and 3 mL of MeOH) was added TMSCH_2N_2 solution (0.81 mL of 2 M in Et_2O , 1.72 mmol, 3 eq.) at room temperature. After being stirred for 5 min at the same temperature, the reaction was quenched with HOAc (0.1 mL) at 0 °C. After concentration, the obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 3%, then 5% EtOAc in Hexanes) to give ester **30** (362 mg, 0.499 mmol, 87% for 3 steps) as a colorless oil. **30**: $[\alpha]_D^{20}$ -35.6 (*c* 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.24 (2H, d, $J = 7.8$ Hz), 7.20 (2H, d, $J = 7.8$ Hz), 7.13 (2H, t, $J = 7.8$ Hz), 7.09 (1H, q, $J = 7.8$ Hz), 6.77 (2H, d, $J = 7.8$ Hz), 5.09 (1H, d, $J = 14.4$ Hz), 5.00 (1H, d, $J = 14.4$ Hz), 4.33 (1H, d, $J = 11.0$ Hz), 4.28-4.21 (2H, m), 3.78 (1H, s), 3.74 (1H, d, $J = 8.4$ Hz), 3.59 (1H, s), 3.38 (3H, s), 3.30 (3H, s), 2.99 (1H, s), 2.95 (1H, dd, $J = 12.2, 10.8$ Hz), 2.91 (1H, s), 2.83 (1H, dd, $J = 16.2, 7.2$ Hz), 2.58 (1H, dd, $J = 16.2, 6.0$ Hz), 2.36 (1H, s), 2.31 (1H, dd, $J = 16.2, 3.2$ Hz), 2.13 (1H, s), 2.01 (1H, d, $J = 15.0$ Hz), 1.66 (2H, td, $J = 12.6, 2.0$ Hz), 1.58-1.51 (2H, m), 1.38 (1H, dd, $J = 12.6, 3.2$ Hz), 1.06 (9H, t, $J = 8.4$ Hz), 1.05 (3H, d, $J = 6.0$ Hz), 1.00 (3H, d, $J = 6.0$ Hz), 0.64 (6H, q, $J = 8.4$ Hz) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 171.8, 171.7, 159.7, 136.7, 131.3, 129.7, 128.8, 128.7, 128.3, 114.0, 97.0, 78.5, 77.1, 76.4, 75.3, 69.8, 66.1, 65.7, 64.4, 54.8, 50.9, 38.0, 37.7, 37.3, 37.2, 36.2, 30.9, 28.9, 18.2, 17.6, 7.3, 5.4 ppm. FTIR (film): 2952, 2924, 2874, 2854,

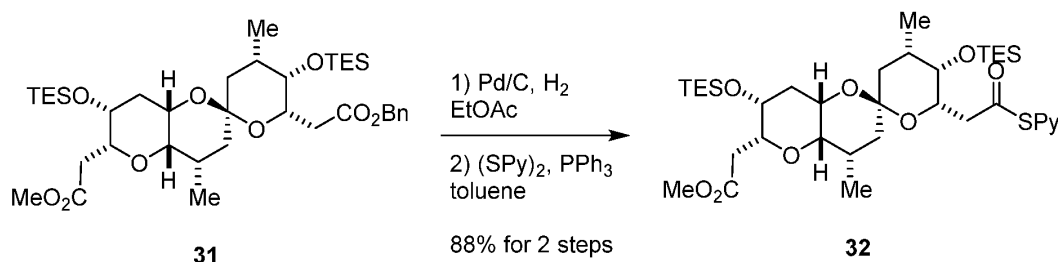
1737, 1613, 1514, 1457, 1436, 1397, 1371, 1302, 1246, 1207, 1163, 1085, 1038, 944, 738, 698 cm^{-1} . HRMS (ESI) m/z : $[M+Na]^+$ calcd for $\text{C}_{40}\text{H}_{58}\text{O}_{10}\text{SiNa}$, 749.3697; found, 749.3720.



[00857] To a stirred solution of MPM ether **30** (350 mg, 0.482 mmol, 1 eq.) in CH_2Cl_2 (10 mL), phosphate buffer (3 mL, pH7) was added DDQ (219 mg, 0.964 mmol, 2 eq.) at room temperature. After being stirred for 1 h at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with CH_2Cl_2 . The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was passed through a SiO_2 to give the crude alcohol, which was used in the next step without further purification.

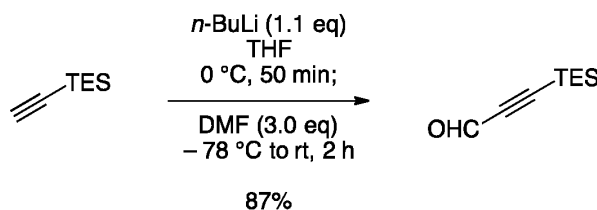
[00858] To a stirred solution of the crude alcohol (calculated as 0.482 mmol, 1 eq.) in CH_2Cl_2 (5.0 mL) were added 2,6-lutidine (129 mg, 1.205 mmol, 2.5 eq.) and TESOTf (255 mg, 0.964 mmol, 2 eq.) at room temperature maintained with a water bath. After being stirred for 30 min at the same temperature, the reaction was quenched with sat. NaHCO_3 aq. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was dried over Na_2SO_4 , filtered, and concentrated under reduced pressure. The obtained crude material was purified by flash column chromatography on neutral silica gel (0%, 3%, then 5% EtOAc in Hexanes) to give TES ether **31** (288 mg, 0.400 mmol, 97%) as a colorless oil. **31**: $[\alpha]_D^{20}$ -52.1 (c 1.00, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 7.26 (2H, d, J = 7.8 Hz), 7.11 (2H, t, J = 7.8 Hz), 7.05 (1H, q, J = 7.8 Hz), 5.09 (1H, d, J = 14.4 Hz), 5.00 (1H, d, J = 14.4 Hz), 4.18 (1H, dd, J = 10.2, 4.0 Hz), 3.80 (1H, s), 3.74 (1H, td, J = 6.6, 1.5 Hz), 3.61 (1H, s), 3.38 (3H, s), 3.29 (1H, s), 2.98 (1H, s), 2.91 (1H, dd, J = 13.8, 12.0 Hz), 2.84 (1H, dd, J = 15.0, 8.4 Hz), 2.60 (1H, dd, J = 15.0, 5.5 Hz), 2.35-2.29 (2H, m), 2.15-2.07 (1H, m), 2.03 (1H, dd, J = 15.0, 3.0 Hz), 1.68 (1H, t, J = 13.8 Hz), 1.64 (1H, t, J = 13.8 Hz), 1.58-1.51 (2H, m), 1.40 (1H, dd, J = 13.2, 4.3 Hz), 1.07 (9H, t, J = 8.4 Hz), 0.99 (3H, d, J = 7.2 Hz), 0.96 (3H, d, J = 7.2 Hz), 0.95 (9H, t, J = 8.4 Hz), 0.65 (6H, q, J = 8.4 Hz), 0.56 (6H, q, J = 8.4 Hz) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 171.8, 171.6, 136.7, 128.8, 128.7, 128.3, 96.9, 77.2, 76.4, 72.5, 70.1, 66.2, 65.6, 64.4, 38.3, 37.5, 37.4, 37.2, 36.3, 31.0, 28.9, 18.5, 17.1, 7.3, 7.2, 5.8, 5.4 ppm. FTIR (film): 2955, 2925, 2876, 1739, 1457, 1372, 1304, 1266, 1240, 1208,

1160, 1130, 1086, 1063, 1036, 947, 856, 740, 697 cm^{-1} . HRMS (ESI) m/z : $[M+Na]^+$ calcd for $\text{C}_{38}\text{H}_{64}\text{O}_9\text{Si}_2\text{Na}$, 743.3987; found, 743.4003.

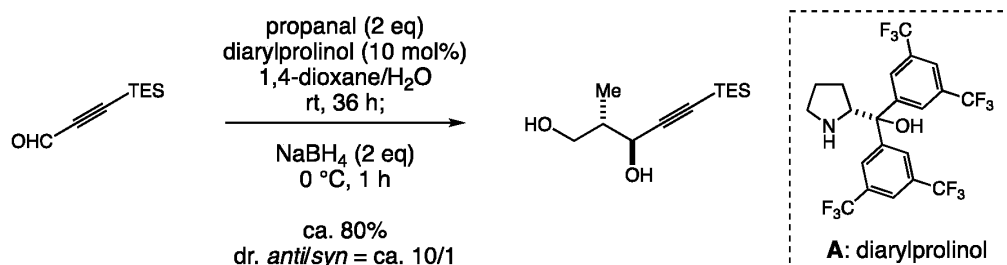


[00859] To a stirred solution of Benzyl ester **31** (250 mg, 0.347 mmol, 1 eq.) in EtOAc (10.0 mL) was added wet 10% Pd/C (25 mg, 10%w/w) at room temperature. The reaction was stirred under 1 atmosphere of hydrogen for 2 h at the same temperature. The resulting mixture was filtered through a pad of Celite (EtOAc). The organic solvent was removed under reduced pressure to give a crude acid, which was used in the next step without further purification.

[00860] To a stirred solution of the crude carboxylic acid (calculated as 0.347 mmol, 1 eq.) in toluene (1.2 mL) were added PPh_3 (110 mg, 0.416 mmol, 1.2 eq.) and $(\text{PyS})_2$ (107 mg, 0.486 mmol, 1.4 eq.) at room temperature. After being stirred for 12 h at the same temperature, the resulting reaction mixture was directly subjected to column chromatography on neutral silica gel (0%, 3%, 5%, then 6% EtOAc in Hexanes) to give pyridinethiol ester **5-32** (221 mg, 0.306 mmol, 88% for 2 steps) as colorless oil. **5-32**: $[\alpha]_D^{20}$ -62.1 (c 1.0, CHCl_3). ^1H NMR (600 MHz, C_6D_6) δ : 8.32 (1H, dd, J = 4.5, 1.3 Hz), 7.52 (1H, d, J = 7.8 Hz), 6.97 (1H, td, J = 7.8, 1.5 Hz), 6.48 (1H, m), 4.19 (1H, dd, J = 9.6, 2.8 Hz), 3.88 (1H, t, J = 1.6 Hz), 3.72 (1H, m), 3.58 (1H, t, J = 1.6 Hz), 3.37 (3H, s), 3.23 (1H, dd, J = 13.8, 10.2 Hz), 3.17 (1H, brs), 3.04 (1H, J = 2.4 Hz, 1H), 2.82 (1H, dd, J = 13.8, 7.2 Hz), 2.59-2.53 (2H, m), 2.41-2.34 (1H, m), 2.27 (1H, m), 2.03 (1H, dt, J = 14.4, 2.4 Hz), 1.70 (1H, t, J = 13.2 Hz), 1.63 (1H, t, J = 13.2 Hz), 1.59 (1H, dt, J = 15.0, 3.5 Hz), 1.53 (1H, dd, J = 13.2, 5.2 Hz), 1.45 (1H, dd, J = 13.2, 5.2 Hz), 1.06 (9H, t, J = 8.4 Hz), 1.00 (3H, d, J = 6.0 Hz), 0.98 (9H, t, J = 8.4 Hz), 0.93 (3H, d, J = 6.6 Hz), 0.64 (6H, q, J = 8.4, 1.1 Hz), 0.56 (6H, q, J = 8.4 Hz) ppm. ^{13}C NMR (150 MHz, C_6D_6) δ : 194.7, 152.7, 150.6, 136.5, 128.5, 123.1, 97.4, 77.7, 77.5, 72.5, 70.0, 68.6, 67.3, 63.9, 47.8, 37.5, 36.7, 36.5, 30.5, 29.1, 27.8, 27.7, 23.4, 21.0, 18.5, 17.3, 7.4, 5.9 ppm. IR (film): 2955, 2931, 2874, 2857, 1708, 1132, 1035, 974 cm^{-1} . HRMS (ESI) m/z : $[M+Na]^+$ calcd for $\text{C}_{36}\text{H}_{61}\text{NO}_8\text{SSi}_2\text{Na}$, 746.3554; found, 746.3578.

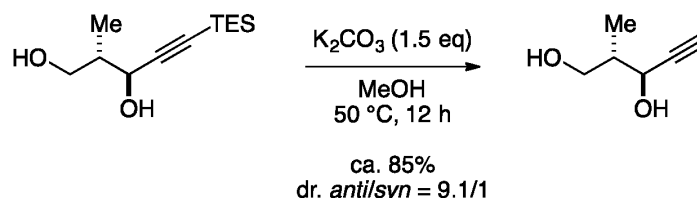
Additional Experimental Procedures for Preparing Compound (1) and Other Compounds

[00861] To a stirred solution of (triethylsilyl)acetylene (25.7 g, 183 mmol) in THF (250 mL) was added *n*-BuLi (2.5 M in hexane, 80.0 mL, 201 mmol) dropwise over 20 min at 0 °C. After stirring for 50 min, DMF (42.0 mL, 549 mmol) was added to a mixture at –78 °C. The reaction mixture was allowed to warm up to room temperature and stirred for 2 h. The reaction was quenched with 3M HCl (185 mL) at 0 °C. The mixture was diluted with EtOAc (260 mL), washed with H₂O (190 mL) four times and brine one time. The organic layer was dried over anhydrous sodium sulfate, filtered, and concentrated under reduced pressure. Finally, crude oil was purified by distillation (~63 °C) to afford desired aldehyde (26.9 g, 160 mmol, 87%) as a colorless oil; ¹H NMR (400 MHz, CDCl₃) δ 9.18 (s, 1H), 1.02 (t, *J* = 7.6 Hz, 9H), 0.69 (q, *J* = 7.6 Hz, 6H); ¹³C NMR (100 MHz, CDCl₃) δ 176.8, 103.5, 101.6, 7.2, 3.7; IR (neat) 2958, 2877, 2151, 1667, 995, 723, 677.

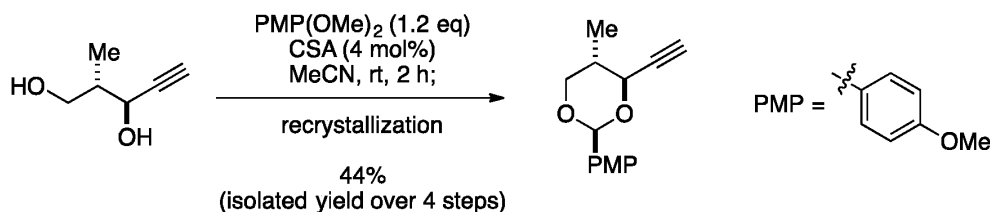


[00862] To a stirred mixture of alkynyl aldehyde (25.4 g, 151 mmol), propanal (21.7 mL, 302 mmol), and water (8.20 mL, 453 mmol) in 1,4-dioxane (150 mL) was added diarylprolinol **A** (7.90 g, 15.1 mmol; Hayashim Y.; Itoh, T.; Aratake, S.; Ishikawa, H. *Angew. Chem. Int. Ed.* **2008**, 47, 2082). After stirring for 36 h at room temperature, NaBH₄ (11.4 g, 302 mmol) was added to a mixture at 0 °C. After stirring for 1 h at 0 °C, the reaction was quenched with H₂O (54 mL), brine (320 mL) and the reaction mixture was extracted with EtOAc five times. The combined organic extracts were dried over anhydrous sodium sulfate, filtered, and concentrated under reduced pressure. The resulting oil was filtered through a pad of silica gel

(450 g) and washed with a mixture of hexanes/EtOAc = 10/1, then 1/1. The filtrate was concentrated again under reduced pressure. The residue (27.7 g, ca. 121 mmol, ca. 80%) was subject to the next reaction without further purification.

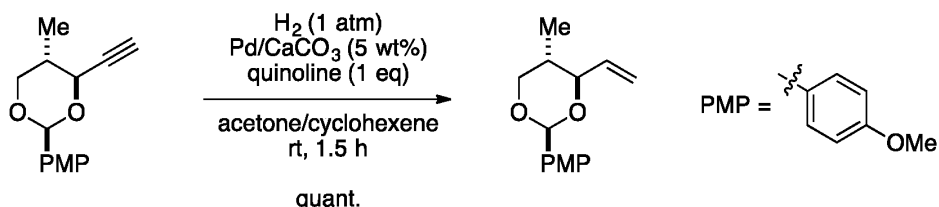


[00863] To a stirred solution of the crude diol (27.7 g, ca. 121 mmol) in MeOH (400 mL) was added K_2CO_3 (25.2 g, 182 mmol) at room temperature. The resulting mixture was heated at 50 °C and stirred for 12 h. After cooling to room temperature, the solvent was removed under reduced pressure. The resulting oil was diluted with CH_2Cl_2 and filtered through a pad of silica gel (130 g) to remove K_2CO_3 . The precipitate was washed with CH_2Cl_2 /MeOH = 10/1 and the filtrate was concentrated under reduced pressure. The resulting oil was filtered through a pad of silica gel (160 g) again and washed with a mixture of hexanes/EtOAc = 5/1, then 1/2. The filtrate was concentrated again under reduced pressure. The residue (11.7 g, ca. 103 mmol, ca. 85%) was subject to the next reaction without further purification.

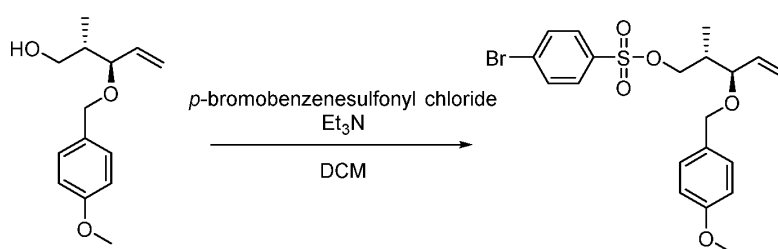


[00864] To a stirred mixture of the crude diol (11.7 g, ca. 103 mmol) and *p*-methoxybenzylidenedimethylacetal (22.0 mL, 124 mmol) in MeCN (180 mL) was added a catalytic amount of CSA (0.96 g, 4.12 mmol) at room temperature. After stirring for 2 h, the reaction mixture was quenched with Et_3N (2.9 mL, 20.6 mmol), filtered through a pad of silica gel (70 g), washed with CH_2Cl_2 and concentrated under reduced pressure. The residue was purified by crystallization from hexanes/ CH_2Cl_2 to afford desired *anti* compound (15.5 g, 66.8 mmol, 44% over 4 steps from alkynyl aldehyde) as a white pure crystal; $[\alpha]_D^{20} +3.00$ (*c* 1.0, MeOH); ^1H NMR (400 MHz, $CDCl_3$) δ 7.42 (d, J = 8.4 Hz, 2H), 6.88 (d, J = 8.4 Hz, 2H), 5.43 (s, 1H), 4.21–4.15 (m, 2H), 3.80 (s, 3H), 3.51 (dd, J = 11.2, 11.2 Hz, 1H), 2.53 (d, J = 2.4 Hz, 1H), 2.19–2.17 (m, 1H), 0.95 (d, J = 7.2 Hz, 3H); ^{13}C NMR (100 MHz, $CDCl_3$)

δ 160.1, 130.2, 127.6, 113.6, 101.5, 80.7, 74.3, 73.7, 72.7, 55.3, 35.1, 12.5; IR (neat) 3263, 2963, 2838, 1614, 1519, 1247, 1024, 997, 834.

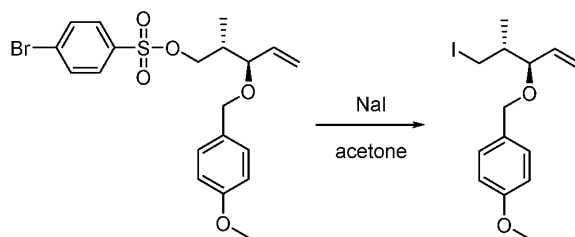


[00865] To a stirred solution of alkyne (15.5 g, 66.8 mmol) in a mixture of acetone (480 mL) and cyclohexene (160 mL) was added quinoline (7.90 mL, 66.8 mmol) and 5wt% Pd/CaCO₃ poisoned with Pb (0.775 g, 5wt%) at room temperature. The flask containing a mixture was charged with hydrogen gas (1 atm) at room temperature, vigorously stirred for 1.5 h and filtered through a pad of Celite. The filter cake was washed with EtOAc and the filtrate was concentrated under reduced pressure. The residue was filtered through a pad of silica gel (230 g) and washed with a mixture of hexanes/EtOAc = 20/1. The filtrate was concentrated again under reduced pressure to afford pure alkene (15.6 g, 66.7 mmol, quant.); a white solid; $[\alpha]_D^{20} +65.4$ (*c* 1.0, CHCl₃); ¹H NMR (400 MHz, CDCl₃) δ 7.44 (d, *J* = 8.8 Hz, 2H), 6.89 (d, *J* = 8.8 Hz, 2H), 5.94–5.85 (m, 1H), 5.51 (s, 1H), 5.36 (d, *J* = 17.2 Hz, 1H), 5.26 (d, *J* = 10.4 Hz, 1H), 4.16 (dd, *J* = 11.6, 4.8 Hz, 1H), 3.89 (dd, *J* = 11.4, 4.8 Hz, 1H), 3.80 (s, 3H), 3.54 (dd, *J* = 11.6, 11.4 Hz, 1H), 1.93–1.87 (m, 1H), 0.79 (d, *J* = 6.8 Hz, 3H); ¹³C NMR (100 MHz, CDCl₃) δ 159.9, 136.0, 131.0, 127.5, 118.0, 113.6, 101.0, 84.7, 73.0, 55.3, 33.9, 12.3; IR (neat) 2965, 1612, 1516, 1248, 1140, 1034, 815.

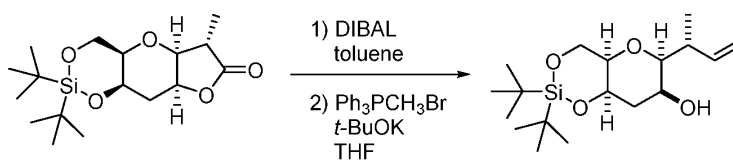


[00866] To a stirred solution of the starting material (445g, 1.885mol) in dichloromethane (750mL) under N₂ atmosphere was added Et₃N (790mL, 5.656mol) at room temperature. To this mixture, a solution of *p*-bromobenzenesulfonyl chloride (730g, 2.828mol) in dichloromethane (600mL) was added dropwise below 41°C. After being stirred for 2hrs at 35–40°C, the mixture was poured into a mixture of *n*-heptane (2L) and water (1.5L) and

stirred for 5min to give a biphasic mixture. The separated aqueous layer was extracted with a mixture of *n*-heptane/EtOAc=2/1(v/v) twice. The combined organic layer was washed with 1M HCl aq. and saturated NaHCO₃ aq. The organic layer was concentrated under reduced pressure to give a crude material as a solid. To this solid was added MTBE (870mL) and *n*-heptane (2.6L) and heated to 49°C to dissolve the solid. The mixture was slowly cooled to room temperature and stirred for 16hrs. After further stirred below 3°C for 1hr, the resulting suspension was filtrated and rinsed with cold *n*-heptane/MTBE=9/1(v/v). The crystals were dried under reduced pressure at 30°C to give the desired compound (740.6g, 1.626mol, 86%). ¹H-NMR (500MHz, CDCl₃) δ ppm 7.75 (2H, d, *J*=8.5Hz), 7.67 (2H, d, *J*=8.5Hz), 7.14 (2H, d, *J*=8.5Hz), 6.86 (2H, d, *J*=8.5Hz), 5.60-5.67 (1H, m), 5.33 (1H, dd, *J*=10.5, 2.0Hz), 5.23 (1H, d, *J*=17.5Hz), 4.46 (1H, d, *J*=11.0Hz), 4.14 (1H, d, *J*=11.0Hz), 4.14 (1H, dd, *J*=9.0, 4.0Hz), 4.10 (1H, dd, *J*=9.0, 4.0Hz), 3.82 (3H, s), 3.57 (1H, dd, *J*=8.0, 8.0Hz), 1.92-2.00 (1H, m), 0.90 (3H, d, *J*=7.0Hz).

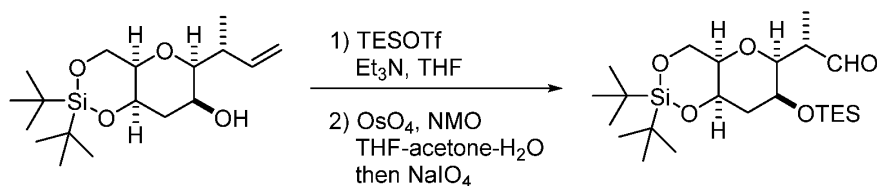


[00867] To a stirred solution of the starting material (730.6g, 1.604mol) in acetone (2.2L) was added NaI (721g, 4.813mol) at room temperature under N₂ atmosphere and heated to 45°C. After being stirred for 2.5hrs, the mixture was cooled below 30°C followed by addition of *n*-heptane (2.2L) and water (2.2L) to give a biphasic mixture. The organic layer was separated and the aqueous layer was extracted with a mixture of *n*-heptane/EtOAc=4/1 (v/v) twice. The combined organic layer was washed with a mixture of aqueous solution containing 6% NaHCO₃ and 3% Na₂S₂O₃ aq. The organic layer was concentrated under reduced pressure and azeotroped with *n*-heptane. The residue was purified by column chromatography on neutral silica gel (10kg, eluent: 0%, 2% then 5% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (551g, 1.591mol, 99%). ¹H-NMR (500MHz, CDCl₃) δ ppm 7.28 (2H, d, *J*=9.0Hz), 6.88 (2H, d, *J*=9.0Hz), 5.69-5.76 (1H, m), 5.35 (1H, dd, *J*=10.0, 1.5Hz), 5.29 (1H, d, *J*=17.5Hz), 4.51 (1H, d, *J*=11.0Hz), 4.28 (1H, d, 11.0Hz), 3.82 (3H, s), 3.53 (1H, dd, *J*=8.0, 8.0Hz), 3.46 (1H, dd, *J*=9.0, 6.0Hz), 3.34 (1H, dd, *J*=9.0, 3.5Hz), 1.51-1.59 (1H, m), 0.94 (3H, d, *J*=7.0Hz).



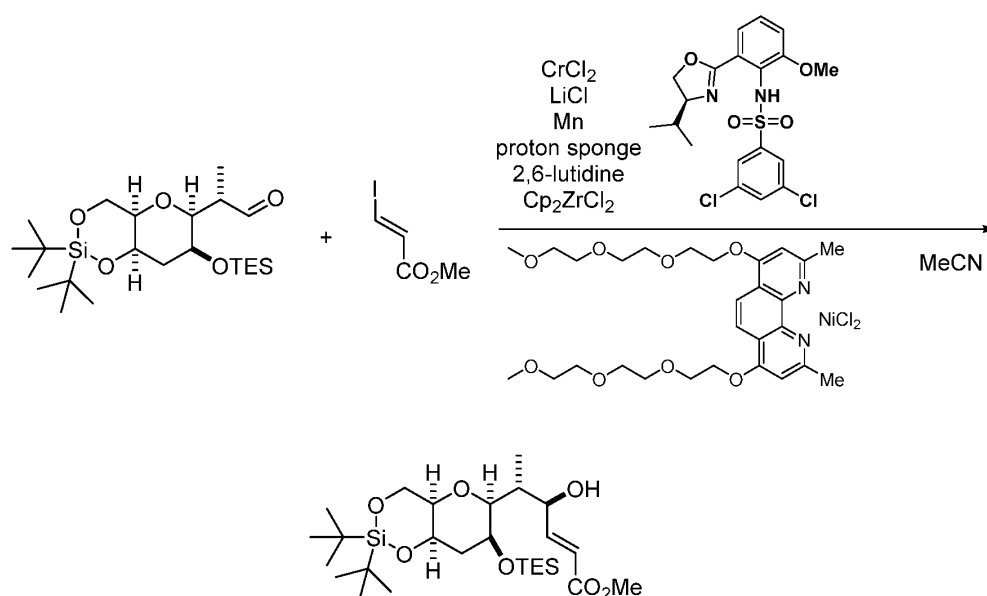
[00868] To a stirred solution of lactone (500g, 1.46mol) in toluene (5L) was added DIBAL solution (1.83L of 1M in toluene, 1.83mol) at $-71 \sim -62^\circ\text{C}$ under N_2 atmosphere. After being stirred for 15min, the reaction was quenched with EtOAc (500mL) at $-69 \sim -65^\circ\text{C}$. Then the mixture was warmed up to -23°C and 30% Rochelle's salt aq. (5L) was added at $-23 \sim 8^\circ\text{C}$. The mixture was stirred for 30min at room temperature to give a biphasic mixture. The organic layer was separated and washed sequentially with water and 5% NaCl aq. The combined aqueous layers were extracted with EtOAc. The combined organic layers were concentrated and azeotroped with toluene twice under reduced pressure to give a crude lactol, which was used in the next reaction without further purification.

[00869] To a suspension of $\text{Ph}_3\text{PCH}_3\text{Br}$ (2086g, 5.84mol) in THF (4L) was added $t\text{-BuOK}$ (491.5g, 4.38mol) at 3°C under N_2 atmosphere. A solution of the crude lactol (calculated as 1.46mol) in THF (1L) was added into the reaction mixture at $1 \sim 9^\circ\text{C}$. After being stirred for 20min at $3 \sim 7^\circ\text{C}$, the reaction was quenched with water (125mL) below 14°C . Then $i\text{PrOAc}$ (5L) and water (2.4L) were added to give a biphasic mixture. The organic layer was separated and washed with 5% NaCl aq. The organic layer was concentrated under reduced pressure. The residue was added toluene (1L) and n -heptane (1L) and stirred for 16hrs at 0°C . The precipitated solid was removed by filtration and filtrated solution was concentrated under reduced pressure. The residue was purified by column chromatography on neutral silica gel (3.0kg, eluent: 0%, 5%, 7%, 9% then 12% EtOAc in n -heptane) to give the desired compound (440g, 1.28mol, 88% in 2steps). $^1\text{H-NMR}$ (500MHz, CDCl_3) δ : 5.70-5.77 (1H, ddd, $J=17, 10.5, 8.0\text{Hz}$), 5.18 (1H, ddd, $J=17, 1.5, 1.5\text{Hz}$), 5.05 (1H, dd, $J=10.5, 1.5\text{Hz}$), 4.42 (1H, dd, $J=3.0, 3.0\text{Hz}$), 4.27 (1H, dd, $J=15, 2.5\text{Hz}$), 4.23 (1H, dd, $J=15, 1.5\text{Hz}$), 3.70-3.74 (1H, m), 3.62 (1H, d, $J=10.5\text{Hz}$), 3.32 (1H, m), 2.97 (1H, dd, $J=9.5, 1.3\text{Hz}$), 2.72-2.80 (1H, m), 2.33 (1H, ddd, $J=15.0, 3.0, 3.0\text{Hz}$), 1.67 (1H, ddd, $J=15.0, 3.0, 3.0\text{Hz}$), 1.12 (3H, d, $J=6.5\text{Hz}$), 1.07 (9H, s), 1.06 (9H, s)



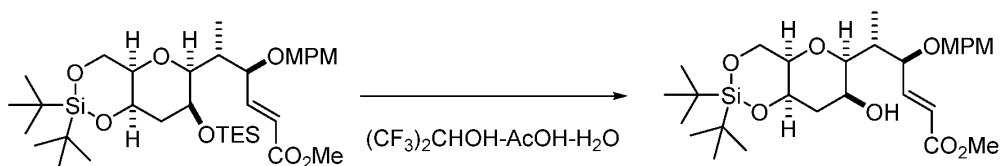
[00870] To a stirred solution of the starting material (440g, 1.28mol) in THF (3.7L) was added triethylamine (367mL, 2.63mol) at 20°C under N₂ atmosphere and cooled to 3°C. To this mixture, TESOTf (354mL, 1.54mol) was added dropwise below 7°C and stirred for 8min at the same temperature. The reaction was quenched with 3.5% NaHCO₃ (4.4L) aq. below 18°C followed by addition of EtOAc (2.2L) and *n*-heptane (2.2L) to give a biphasic mixture. The organic layer was separated and washed with 5% NaCl aq. The organic layer was dried over Na₂SO₄, filtered and concentrated under reduced pressure to give a crude material, which was used in the next reaction without further purification.

[00871] A solution of the crude (calculated as 1.28mol) in THF (2.1L) and acetone (2.1L) was added NMO (464g, 3.96mol) followed by addition of water (0.5L) at room temperature. To this mixture, 4% OsO₄ aq. (194mL, 32mmol) was added and stirred for 19hrs at room temperature. Then water (2.1L) and NaIO₄ (847g, 3.96mol) were added to this mixture and further stirred for 70min at room temperature. To this mixture, *n*-heptane (2.9L) and water (1.2L) were added and stirred for 5min at room temperature to give a biphasic mixture. The organic layer was separated and aqueous layer was extracted with *n*-heptane. The combined organic layers were added to 20% Na₂SO₃ aq. and the resulting biphasic mixture was stirred for 30min at room temperature. The organic layer was separated and washed sequentially with 10% Na₂SO₃ aq., water, 5% NaCl aq. The organic layer was dried over Na₂SO₄, filtered, concentrated under reduced pressure. The residue was purified by column chromatography on neutral silica gel (5.9kg, eluent: 0%, 5%, then 10% EtOAc in *n*-heptane). The collected fractions were concentrated and the resulting aldehyde was solidified by cooling with ice-water bath to give the aldehyde (502.8g, 1.10mol, 86% in 2steps). ¹H-NMR (500MHz, CDCl₃) δ: 9.84 (1H, s), 4.29-4.31 (1H, m), 4.28 (1H, dd, *J*=12.5, 3.0Hz), 4.21 (1H, dd, *J*=12.5, 1.5Hz), 3.86-3.88 (1H, m), 3.57 (1H, dd, *J*=7.5, 2.5Hz), 3.29-3.32 (1H, m), 2.80-2.86 (1H, dq, *J*=7.5Hz, 7.5Hz), 2.21-2.25 (1H, ddd, *J*=15.0, 2.5, 2.5Hz), 1.77-1.81 (1H, ddd, *J*=15.0, 4.0, 4.0Hz), 1.16 (3H, d, *J*=7.5Hz), 1.07 (9H, s), 1.04 (9H, s), 0.96-1.00 (9H, m), 0.58-0.67 (6H, m).

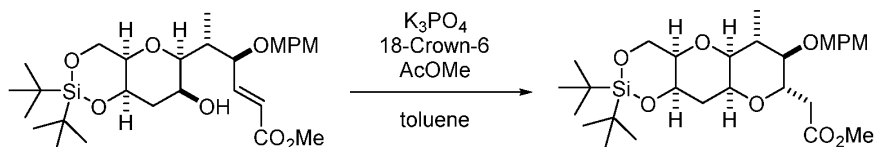


[00872] In a glove box, CrCl_2 (22mg, 0.18mmol), proton sponge (42mg, 0.20mmol), sulfonamide ligand (88mg, 0.20mmol) were added in a well-dried round bottomed flask and added anhydrous MeCN (3.0mL). The mixture was stirred in the glove box at room temperature.

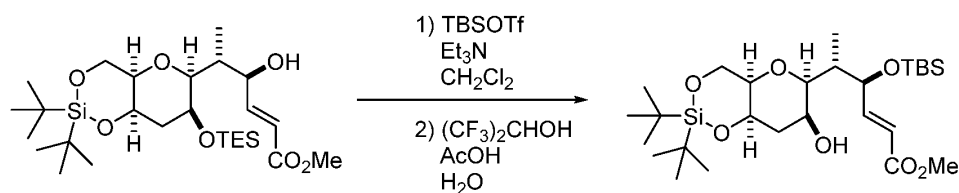
[00873] In another flask, the aldehyde (500mg, 0.90mmol) was added toluene. After azeotropic removal of toluene, the flask was brought in the glove box followed by addition of vinyl iodide (477mg, 2.25mmol), Mn (198mg, 3.60mmol), LiCl (76mg, 1.80mmol), Cp_2ZrCl_2 (289mg, 0.99mmol) and 2,6-lutidine (0.21mL, 1.80mmol). To this flask, the Cr solution prepared above was added followed by addition of Ni catalyst (3.0mg, 4.5 μmol). The mixture was stirred vigorously for 145min at 30°C. The flask was taken out of the glove box and Florisil[®] (1g) was added to this mixture at room temperature. After dilution with *n*-heptane/EtOAc=1/1(v/v) (5mL), the suspension was stirred vigorously for 35min at room temperature and passed through a neutral silica gel (5g) pad. The desired product was eluted with EtOAc and collected solution was concentrated under reduced pressure. The residue was purified by column chromatography on neutral silica gel (10g, eluent; 0%, 10%, 20%, 33% then 50% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (462mg, 0.85mmol, 94%).



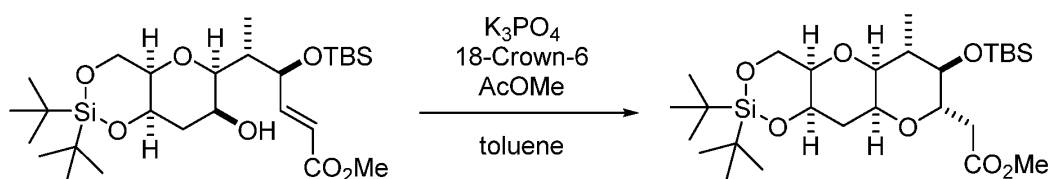
[00874] To a stirred solution of the starting material (636.6g, 0.957mol) in hexafluoroisopropanol (HFIP) (1.6L) and water (160mL) were added AcOH (55mL, 0.957mol) at 5°C under N_2 atmosphere. The mixture was stirred for 30min below 5°C and 3hrs at room temperature. The reaction was quenched with 5% NaHCO_3 aq. (5L) followed by addition of *n*-heptane (5L). After separation of the organic layer, the aqueous and HFIP layer were combined and extracted with *n*-heptane twice. Three organic layers were combined and washed with 5% NaCl aq. The organic layer was dried over Na_2SO_4 , filtered and concentrated. The residue was purified by column chromatography on neutral silica gel (8.5kg, eluent: 0%, 9.1%, 11% then 17% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired alcohol (480g, 0.871mol, 91%).



[00875] The starting material (480.0g, 0.872mol) was azeotroped with toluene (1.4L) twice and added toluene (4.8L). After addition of MeOAc (480mL), the mixture was cooled to 1°C and kept under N_2 atmosphere until used. In another flask was added K_3PO_4 (92.5g, 0.436mol), 18-crown-6 (345.5g, 1.307mol) and MeOH (480mL). This mixture was sonicated for 20min and concentrated under reduced pressure. The resulting mixture was azeotroped with toluene (480mL) three times and toluene (2.4L) was added to the mixture and filtrated. The filtrated solution was added dropwise to the solution of the starting material prepared above below 4°C. After being stirred for 16min below 5°C, the mixture was poured into chilled water (2.4L) followed by addition of *i*PrOAc (2.4L), water (2.4L) and NaCl (100g) to give a biphasic mixture. The separated organic layer was washed with 5% NaCl aq. The organic layer was dried over Na_2SO_4 , filtered and concentrated. The residue was purified by column chromatography on neutral silica gel (8.6kg, eluent: 0%, 9.1%, 17%, 25% then 33% EtOAc in *n*-heptane). The pure fractions were concentrated to give the oxy-Michael product (293g, 0.531mol, 61% in 2steps).



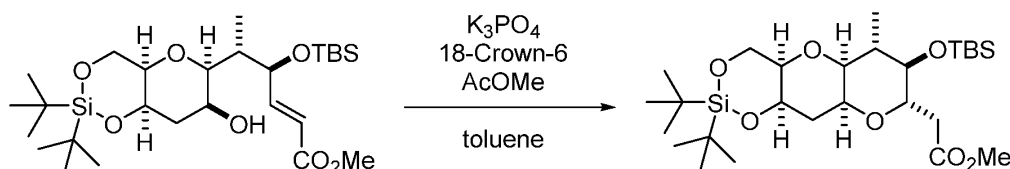
[00876] To a stirred solution of the starting material (41.3g, 75.8mmol) in dichloromethane (206mL) was added Et₃N (25.4mL, 181.9mmol) at 1°C under N₂ atmosphere. To this mixture, TBSOTf (20.9mL, 91.0mmol) was added dropwise below 10°C. After being stirred for 30min at 0°C, the reaction was quenched with 10% NaHCO₃ aq. (206mL) below 17°C. The mixture was extracted with *n*-heptane and the resulting organic phase was washed with water and 5% NaCl aq. The organic layer was dried over Na₂SO₄, filtered and concentrated to give a crude oil, which was used in the next step without further purification. To this crude material (calculated as 75.8mmol) were added HFIP (124mL) and water (12.5mL). After cooling to 4°C, AcOH (4.4mL) was added and the resulting mixture was stirred for 30min at 0°C and another 1.5hrs at room temperature under N₂ atmosphere. The mixture was quenched with 5% NaHCO₃ aq. (400mL) followed by addition of *n*-heptane (400mL) to give a biphasic mixture. The organic phase was separated and washed with 5% NaCl aq. The organic layer was dried over Na₂SO₄, filtered and concentrated. The residue was purified by column chromatography on neutral silica gel (750g, eluent: 0%, 4.8%, 9.1% then 11% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (40.5g, 74.2mmol, 98%). ¹H-NMR (500MHz, CDCl₃) δ: 7.01 (1H, dd, *J*=16.0, 5.0Hz), 5.98 (1H, dd, *J*=16.0, 2.5Hz), 4.43-4.45 (1H, m), 4.40 (1H, dd, *J*=2.5, 2.5Hz), 4.23 (1H, dd, *J*=13.0, 3.0Hz), 4.16 (1H, dd, *J*=13.0, 1.0Hz), 3.75 (3H, s), 3.73-3.76 (1H, m), 3.59 (1H, d, *J*=11.0Hz), 3.28-3.29 (2H, m), 2.33 (1H, ddd, *J*=14.5, 2.5, 2.5Hz), 2.10-2.18 (1H, m), 1.73 (1H, ddd, *J*=15.0, 3.5, 3.5Hz), 1.04-1.06 (21H, m), 0.90 (9H, s), 0.03 (3H, s), 0.02 (3H, s).



[00877] In N₂ atmosphere, K₃PO₄ (0.624g, 2.94mmol), 18-crown-6 (2.33g, 8.82mmol) and MeOH (15mL) were added in a flask and sonicated for 6min. After the mixture was concentrated under reduced pressure, the mixture was azeotroped with toluene (20mL) three

times. The residue was added toluene (15mL) and kept under N₂ atmosphere at room temperature.

[00878] In another flask, the starting material (5.34g, 9.80mmol) was azeotroped with toluene (53mL) twice and the residue was added toluene (53mL) followed by MeOAc (5.3mL). After the mixture was cooled below 3°C, the solution containing K₃PO₄ prepared above was added dropwise to the solution of the starting material below 3°C over 40min. The mixture was poured into a mixture of cold EtOAc/water=1/1 (v/v) (106mL) to give a biphasic mixture. The organic layer was separated and the aqueous layer was extracted with EtOAc. The combined organic layer was washed with water and 5% NaCl aq. The separated organic layer was dried over Na₂SO₄, filtrated and concentrated under reduced pressure. The residue was purified by column chromatography on neutral silica gel (80g, eluent: 0%, 5%, 7%, 10% then 15% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (4.35g, 7.98mmol, 81%). ¹H-NMR (500MHz, CDCl₃) δ: 4.30 (1H, ddd, *J*=8.0, 3.5, 3.5Hz), 4.13 (1H, dd, *J*=12.5, 5.5Hz), 4.09 (1H, dd, *J*=12.5, 3.0Hz), 3.87 (1H, ddd, *J*=9.0, 9.0, 3.0Hz), 3.79 (1H, ddd, *J*=7.5, 4.5, 4.5Hz), 3.70 (3H, s), 3.60 (1H, ddd, *J*=5.5, 3.5, 3.5Hz), 3.34 (1H, dd, *J*=8.0, 5.0Hz), 3.30 (1H, dd, *J*=10.0, 8.5Hz), 2.71 (1H, dd, *J*=14.5, 3.0Hz), 2.41 (1H, dd, *J*=14.5, 10.0Hz), 2.11 (1H, ddd, *J*=13.5, 8.0, 8.0Hz), 1.98 (1H, ddd, *J*=13.5, 4.5, 4.5Hz), 1.77-1.85 (1H, m), 1.10 (3H, d, *J*=6.5Hz), 1.06 (9H, s), 1.02 (9H, s), 0.90 (9H, s), 0.09 (3H, s), 0.06 (3H, s).

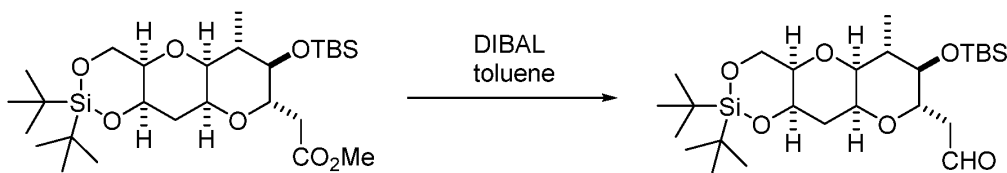


[00879] In N₂ atmosphere, K₃PO₄ (4.73g, 22.27mmol), 18-crown-6 (17.66g, 66.81mmol) and MeOH (120mL) were added in a flask and sonicated for 8min. After the mixture was concentrated under reduced pressure, the mixture was azeotroped with toluene (120mL) three times. The residue was added toluene (150mL) and kept under N₂ atmosphere at room temperature until used.

[00880] In another flask, the starting material (40.45g, 74.24mmol) was azeotroped with toluene (120mL) twice and the residue was added toluene (400mL) followed by MeOAc (40mL). After the mixture was cooled below 3°C, the solution containing K₃PO₄ prepared above was added dropwise to the solution of the starting material below 3°C over 90min. The mixture was poured into a mixture of cold EtOAc/water=1/1 (v/v) (800mL) to give biphasic

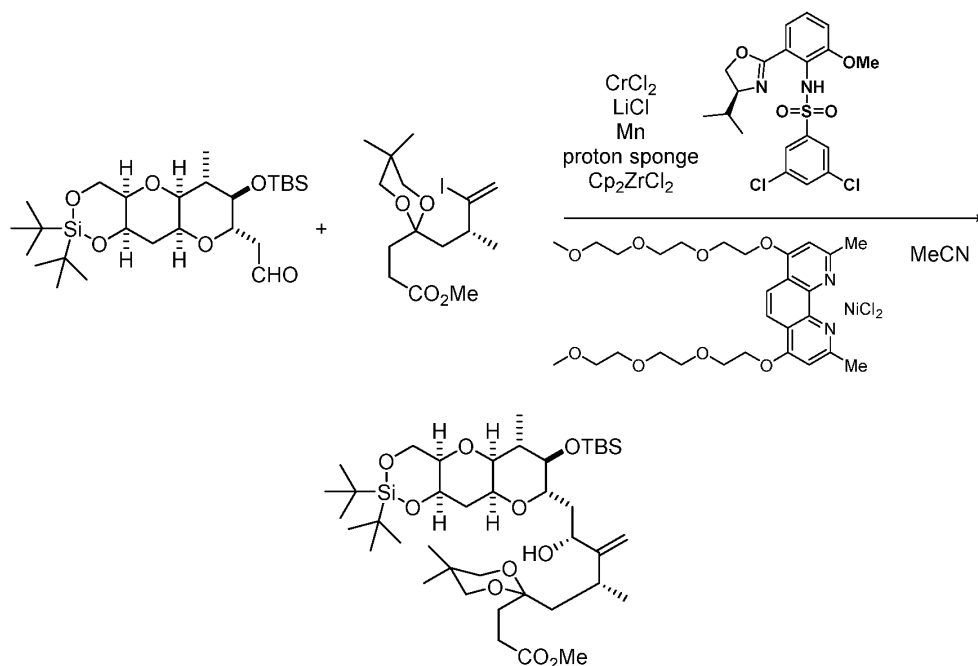
mixture. The organic layer was separated and the aqueous layer was extracted with EtOAc (200mL). The combined organic layer was washed with 5wt% NaCl aq. The separated organic layer was dried over Na₂SO₄, filtrated and concentrated under reduced pressure. The residue was purified by column chromatography on neutral silica gel (600g, eluent: 0%, 9%, 11%, then 14% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (23.36g, 42.9mmol) as a colorless oil. Then, the inseparable fractions were combined and the solvent was removed under reduced pressure. The residue was re-purified by column chromatography on neutral silica gel (300g, eluent: 0%, 9%, 11%, then 14% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (9.90g, 18.17mmol) as a colorless oil. Both desired compound were combined to give the desired product (33.26g, 61.04mmol, 82%).

[00881] The desired product (6.13g, 11.25mmol) was added MeOH (50mL) and the mixture was azeotroped under reduced pressure at 40°C. The residual solid was added MeOH (36.8mL) and the slurry was warmed up to 50°C. After the slurry became a clear solution, the solution was cooled to 35°C and seed was added. After a solution became cloudy, the mixture was stirred at room temperature for 30min and then water (7.4mL) was added to the mixture. After being stirred at room temperature for 3hrs, the generated crystal was collected by suction and washed with cold MeOH/water=5/1 (v/v) (40mL). The crystal was dried at 40°C under reduced pressure for 2hrs to give the desired product (5.48g, 10.06mmol, 89%).
¹H-NMR (500MHz, CDCl₃) δ: 4.30 (1H, ddd, *J*=8.0, 3.5, 3.5Hz), 4.13 (1H, dd, *J*=12.5, 5.5Hz), 4.09 (1H, dd, *J*=12.5, 3.0Hz) 3.87 (1H, ddd, *J*=9.0, 9.0, 3.0Hz), 3.79 (1H, ddd, *J*=7.5, 4.5, 4.5Hz), 3.70 (3H, s), 3.60 (1H, ddd, *J*=5.5, 3.5, 3.5Hz), 3.34 (1H, dd, *J*=8.0, 5.0Hz), 3.30 (1H, dd, *J*=10.0, 8.5Hz), 2.71 (1H, dd, *J*=14.5, 3.0Hz), 2.41 (1H, dd, *J*=14.5, 10.0Hz), 2.11 (1H, ddd, *J*=13.5, 8.0, 8.0Hz), 1.98 (1H, ddd, *J*=13.5, 4.5, 4.5Hz), 1.77-1.85 (1H, m), 1.10 (3H, d, *J*=6.5Hz), 1.06 (9H, s), 1.02 (9H, s), 0.90 (9H, s), 0.09 (3H, s), 0.06 (3H, s).



[00882] The starting material (12g, 22.02mmol) was azeotroped with toluene (36mL) twice and the residue was added toluene (240mL). After the mixture was cooled below -70°C, DIBAL solution (26.4mL of 1M in toluene, 26.43mmol) was added to the mixture at -73 ~ -71°C under N₂ atmosphere. After being stirred for 32min, the reaction was quenched with

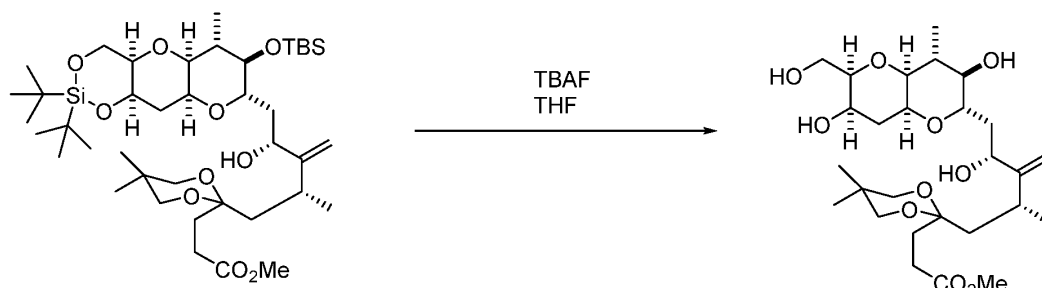
acetone (12mL) at $-75 \sim -74^{\circ}\text{C}$. Then the mixture was warmed up to -30°C and 30wt% Rochelle's salt aq. (180mL) was added at $-29 \sim 4^{\circ}\text{C}$. The mixture was stirred for 3hrs at room temperature to give a biphasic mixture. The organic layer was separated and washed with 5wt% NaCl aq. The separated organic layer was dried over Na_2SO_4 , filtrated and concentrated under reduced pressure to give a crude aldehyde (14.18g), which was used as 100% yield in the next reaction without further purification.



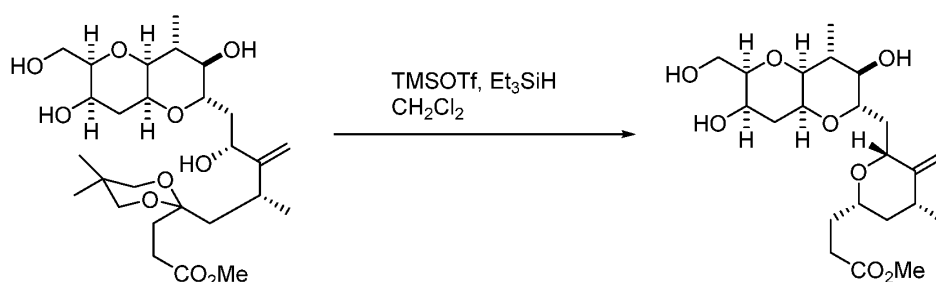
[00883] In a glove bag, CrCl_2 (541mg, 4.41mmol), proton sponge (1.04g, 4.85mmol), sulfonamide ligand (2.15g, 4.85mmol) were added in a well-dried round bottomed flask. After taking flask out from glove bag, anhydrous MeCN (90.7mL) was added under N_2 atmosphere. The mixture was stirred at room temperature until used.

[00884] In another flask, the aldehyde (11.34g, 22.03mmol) and vinyl iodide (10.47g, 26.43mmol) were added toluene. After azeotropic removal of toluene (twice), the residual mixtures were added Mn (4.84g, 88.10mmol), LiCl (1.87g, 44.05mmol) and Cp_2ZrCl_2 (7.08g, 24.23mmol) and then the flask was purged by N_2 gas. To this flask, the Cr solution prepared above was added. After the mixture was warmed up to 40°C , solution of Ni catalyst (730uL (prepared as 100mg/mL MeCN in advance), 0.11 mmol) was added. During stirring vigorously for 19hrs at 40°C , additional Ni catalyst (Y. Kishi et.al, Journal of the American Chemical Society (2015), 137(19), 6219-6225) (365uL, 0.06mmol) was added 6 times to

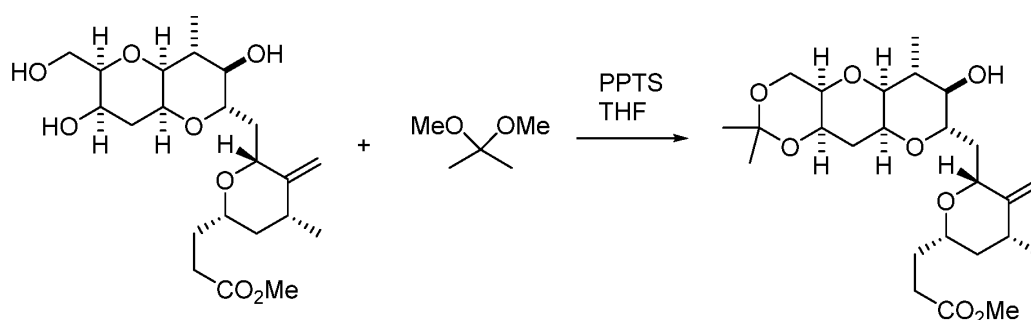
complete the reaction. The mixture was added *i*PrOAc (120mL) followed by a mixture of 5wt% NaHCO₃ and 5wt% D,L-serine aqueous solution (120mL) at 40°C. After stirring for 4hrs at 40°C, the mixture was cooled to room temperature and passed through Celite (25g) with *i*PrOAc (600mL) washing to give biphasic mixture. The organic layer was separated and the organic layer was washed with 5wt% NaHCO₃ aq., water and 5% NaCl aq. The separated organic layer was dried over Na₂SO₄, filtrated and concentrated under reduced pressure to give a crude desired product (25.26g), which was used as 100% yield in the next reaction without further purification.



[00885] To a stirred solution of the starting material (17.3g, 22.03mmol) in THF (52ml) was added TBAF (88mL of 1M in THF, 88.13mmol) at 0°C. After stirring for 15hrs at room temperature, the mixture was added EtOAc (180mL) followed by 10wt% NH₄Cl aq. (180mL) to give biphasic mixture. The organic layer was separated and the aqueous layer was extracted with EtOAc (180mL) twice. The combined organic layer was washed with saturated NaCl aq. The separated organic layer was dried over Na₂SO₄, filtrated and concentrated under reduced pressure. The residue was purified by column chromatography on NH-silica gel (120g, eluent: 0%, 20%, 50%, 100% EtOAc in *n*-heptane and then 5% MeOH in EtOAc). The collected fractions were concentrated to give the desired product (17.46g, including impurities), which was used as 100% yield in the next reaction without further purification.



[00886] The starting material (11.69g, 22.03mmol) was azeotroped with toluene (35mL) twice and the residue was added CH₂Cl₂ (117mL) followed by Et₃SiH (17.6 mL, 110.14mmol) at rt. After the mixture was cooled below -70°C, TMSOTf (16.0mL, 88.11mmol) was added to the mixture at -72 ~ -70°C under N₂ atmosphere. After being stirred for 30min, the reaction mixture was warmed up to 0°C. After being stirred for 27min at 0°C, the reaction mixture was quenched with water (117mL) below 14°C and then the mixture was stirred at room temperature. After being stirred for 25min, saturated NaHCO₃ aq. (58mL) was added and the mixture was stirred for 25min at room temperature to give a biphasic mixture. The organic layer was separated and the aqueous layer was extracted with CH₂Cl₂ (117mL). The combined organic layer was dried over Na₂SO₄, filtrated and concentrated under reduced pressure to give a crude desired product (18.95g), which was used as 100% yield in the next reaction without further purification.



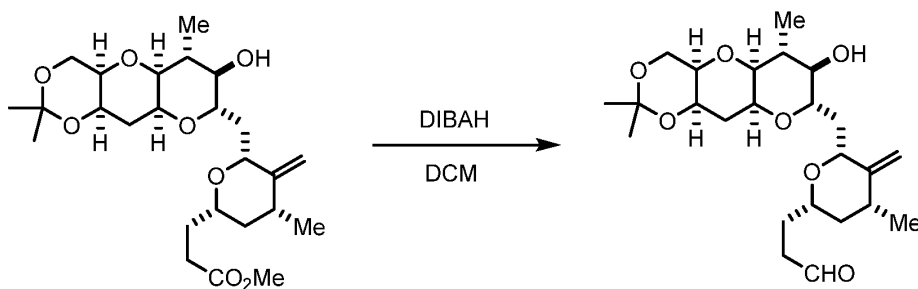
[00887] The starting material (9.44g, 22.03mmol) was azeotroped with toluene (30mL) twice and the residue was added THF (94mL), 2,2-dimethoxypropane (10.84 mL, 88.13 mmol) and PPTS (277mg, 1.10mmol) at room temperature. After being stirred at 40°C under N₂ atmosphere for 4.5hrs, the reaction mixture was cooled to room temperature and then the reaction mixture was diluted with EtOAc (100mL) followed by saturated NaHCO₃ aq. (100mL). After being stirred for 15min, a biphasic mixture was obtained. The organic layer was separated and the aqueous layer was extracted with EtOAc (100mL). The combined

organic layer was washed with saturated NaCl aq. The separated organic layer was dried over Na_2SO_4 , filtrated and concentrated under reduced pressure to give a crude desired product (18.95g)

[00888] A crude desired product (18.95g) was purified by ODS-column chromatography on YMC-GEL(100g, eluent: 3%, 40% MeCN in water). The collected fractions were extracted by CH_2Cl_2 (500mL+250mL) and the combined organic layer was dried over Na_2SO_4 , filtrated and concentrated under reduced pressure to give the desired product (12.00g, including impurities).

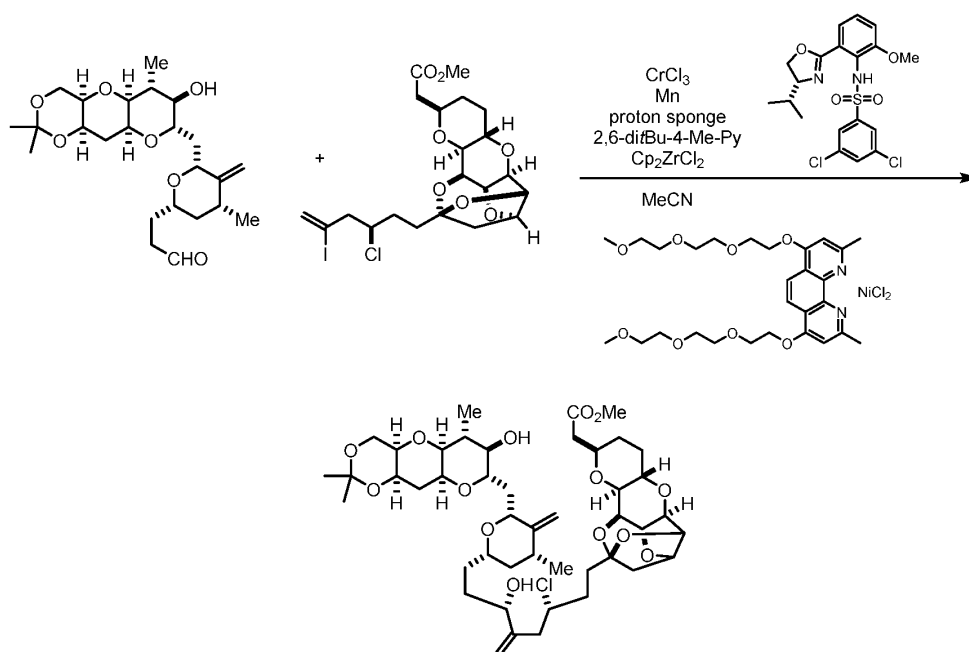
[00889] A crude desired product (12.00g) was purified by column chromatography on neutral silica gel (150g, eluent: 0%, 20%, 33%, 50%, and 66% EtOAc in *n*-heptane). The collected fractions were concentrated to give a crude desired product (4.56g, including impurities).

[00890] A crude desired product (4.56g) was added a mixture of *n*-hexane/EtOAc=9/1 (v/v) (45mL) and the slurry was warmed up to 60°C. After the slurry became a clear solution, the solution was cooled to 45°C and seed was added. After a solution became cloudy, the mixture was stirred at room temperature for 50min and then the mixture was stirred at 0°C for 65min and then stirred at -10°C for 17.5hrs. The generated crystal was collected by suction and washed with cold *n*-hexane (13.5mL). The crystal was dried at 40°C under reduced pressure to give the desired product (3.57g, 7.62mmol, 35% in 5steps). $^1\text{H-NMR}$ (500MHz, CDCl_3) δ : 4.88 (1H, s), 4.82 (1H, d, $J=2.0\text{Hz}$), 4.18 (1H, q, $J=5.5\text{Hz}$), 4.03 (1H, dd, $J=12.5, 3.0\text{Hz}$), 4.00-3.95 (2H, m), 3.88 (1H, dd, $J=6.0, 4.0\text{Hz}$), 3.86 (1H, dd, $J=12.0, 2.5\text{Hz}$), 3.67 (3H, s), 3.64-3.57 (1H, m), 3.55 (1H, d, $J=8.5\text{Hz}$), 3.42 (1H, t, $J=3.0\text{Hz}$), 3.36 (1H, dt, $J=8.0, 5.5\text{Hz}$), 3.26 (1H, q, $J=2.5\text{Hz}$), 2.54-2.40 (2H, m), 2.33-2.23 (1H, m), 2.19-2.11 (2H, m), 2.10-2.00 (2H, m), 1.86-1.68 (4H, m), 1.44 (3H, s), 1.43 (3H, s), 1.16 (3H, d, $J=8.0\text{Hz}$), 1.15-1.09 (1H, m), 1.08 (3H, d, $J=6.0\text{Hz}$).



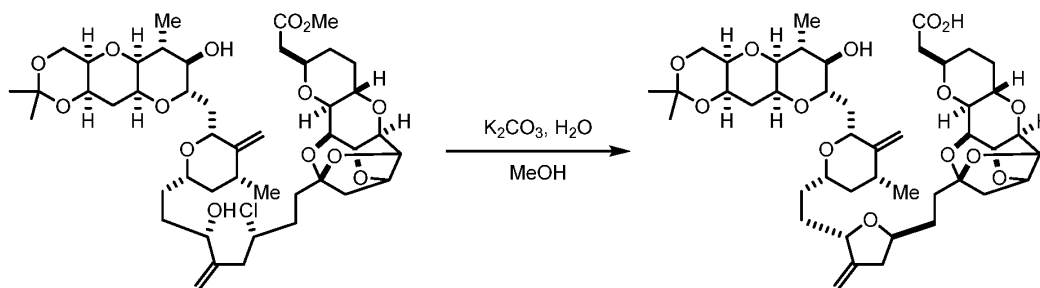
[00891] To a stirred solution of starting material (52.0g, 0.11mol) in dichloromethane (1.04L) was added DIBAL solution (253mL of 1M in toluene, 0.255mol) at -74.3 ~ -71.5°C under N_2 atmosphere. After being stirred for 76min, the reaction was quenched with acetone

(24.5mL) at $-75.2 \sim -73.8^{\circ}\text{C}$. After being stirred for additional 9min, MeOH (27.0mL) was added to the reaction at $-74.2 \sim -73.5^{\circ}\text{C}$, followed by addition of BHT (52.0mg). Then the mixture was warmed up to -20°C and added to 40wt% Rochelle's salt aq. (1.04kg), then EtOAc (936mL) was added to the mixture. The mixture was stirred for 85min at room temperature to give a biphasic mixture. The organic layer was separated and re-extracted with EtOAc (520mL). Combined organic layer was washed with 20wt% NaCl aq. (520g). After addition of BHT (52.0mg), the organic layer was concentrated and azeotroped with toluene (208mL) and MeCN (156mL) under reduced pressure to give a crude aldehyde, which was used in the next reaction without further purification (assumed as 90% yield, 0.1mol).

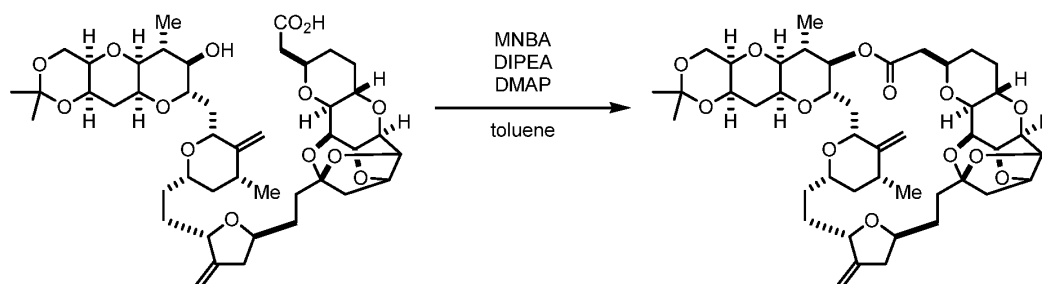


[00892] To a reactor, CrCl_3 (3.16g, 20mmol), proton sponge (4.71g, 22mmol), sulfonamide ligand (9.74g, 22mmol), Mn (11.0g, 200mmol) and Cp_2ZrCl_2 (75.9g, 260mmol) were charged, then reactor was purged with Argon. To the reactor, anhydrous MeCN (263mL) was added and stirred for 70min at 30°C . To a reaction, Ni-complex (662mg, 1.0mmol) in anhydrous MeCN (6.6mL) was added. Then, a mixture of the aldehyde (43.8g, 100mmol), vinyl iodide (63.2g, 114mmol) and 2,6-di-*tert*-butyl-2-methyl pyridine (41.0g, 200mmol) in anhydrous MeCN (131mL), which were azeotroped with MeCN (219mL) in advance, was added. After substrates addition, Ni-complex (661mg, 1.0mmol) in anhydrous MeCN (6.57mL) was added twice to complete the reaction. To the reaction, toluene (876mL) was added, then cooled to 0°C . To the mixture, water (788mL) and 20wt% citric acid aq. (87.5g)

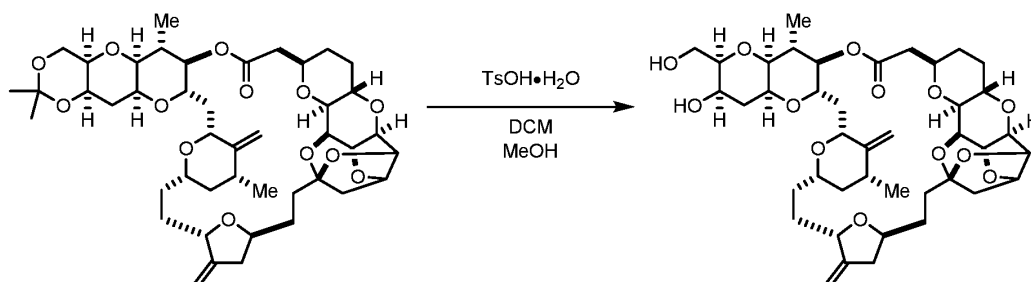
was added at around 14°C for 20min. After removal of the aqueous layer, the organic layer was passed through hyflo super-cel (35.0g) then washed with toluene (876mL). Combined organic layer was washed with 33% MeCN aq. (876mL) followed by a mixture of 5wt% NaHCO₃-10wt% Na₂SO₄ aqueous solution (438g). Organic layer was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (1900g, eluent; 10%, 25%, 50%, 67% then 100% EtOAc in *n*-heptane). The collected fractions were concentrated to give the desired product (79.4g, 92mmol, 82% in two steps). ¹H-NMR (500MHz, CDCl₃) δ: 5.16 (1H, s), 4.97 (1H, s), 4.87 (1H, s), 4.82 (1H, s), 4.70 (1H, t, *J*=4.7Hz), 4.63 (1H, t, *J*=4.5Hz), 4.43 (1H, d, *J*=1.8Hz), 4.18-4.33 (4H, m), 3.93-4.10 (5H, m), 3.78-3.92 (3H, m), 3.68 (3H, s), 3.59 (1H, d, *J*=3.0Hz), 3.19-3.48 (3H, m), 2.85-2.98 (2H, m), 2.56-2.71 (2H, m), 2.48 (1H, dd, 15.1, 6.0Hz), 2.41 (1H, dd, *J*=15.6, 5.8Hz), 1.93-2.33 (11H, m), 1.51-1.92 (10H, m), 1.36-1.50 (8H, m), 1.14 (3H, d, *J*=7.5Hz), 1.08 (3H, d, *J*=6.3Hz).



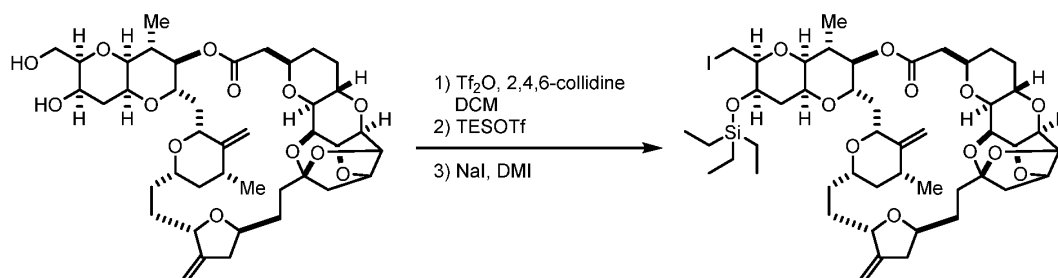
[00893] To a stirred solution of the starting material (79.3g, 91.4mmol) in MeOH (793mL) was added K₂CO₃ (63.2g, 0.457mol) at room temperature under N₂ atmosphere. After heated to 55 °C, the mixture was stirred for 3hr, then water (24.7mL, 1.37mol) was added at same temperature. After being stirred for 20hr, the reaction was cooled to room temperature, then toluene (1.59L) and water (1.59L) was added. Aqueous layer was washed with toluene (1.59L) twice. To the resulted aqueous layer, dichloromethane (1.59L) and 25wt% NH₄Cl aq. (2.38kg) were added. After separation of two layers, the aqueous layer was re-extracted with dichloromethane (2.38L). Combined organic layer was concentrated and azeotroped with toluene (397mL) to give a crude seco acid, which was used in the next reaction without further purification.



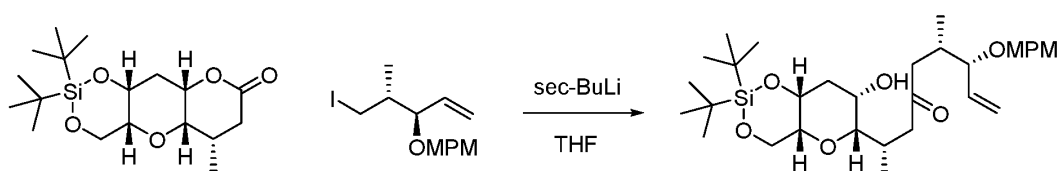
[00894] To a stirred solution of *N,N*-dimethylaminopyridine (DMAP, 67.0g, 549mmol) in toluene (5.60L) was added *N,N*-diisopropylethylamine (DIPEA, 95.6mL, 549mmol) and 2-methyl-6-nitrobenzoic anhydride (MNBA, 94.4g, 274mmol) at 25 °C under N₂ atmosphere, then heated to 80 °C. To the mixture, starting material (74.7g, 91.4mmol) in toluene (1.49L) was added dropwise for 6hr. After cooled to 25 °C, the mixture was washed with 50vol% DMF aq. (1.49L) twice. The second 50% DMF aqueous layer was re-extracted with toluene (500mL). Combined organic layer was washed with 20wt% NH₄Cl aq. (1.49kg) twice, each aqueous layer was re-extracted with EtOAc (750mL). Combined organic layer was washed with 20wt% NaCl aq. (374g) and re-extracted with EtOAc (750mL). Combined organic layer was concentrated and azeotroped with MeCN (200mL). The residue was dissolved in dichloromethane (448mL) and concentrated. The resulted high concentrated dichloromethane solution was added to MeCN (1.05L) at 0 °C under N₂ atmosphere. Resulted precipitation was filtered and washed with MeCN (224mL) followed by drying at 40 °C gave a desired macrolactone (41.5g, 51.9mmol, 56.8% in two steps).



[00895] To a stirred solution of starting material (40.5g, 50.7mmol) in dichloromethane (312mL) and MeOH (579mL) was added *p*-toluenesulfonic acid monohydrate (194mg, 1.01mmol) at 25°C under N₂ atmosphere for 4hr. The reaction was passed through NH-silica gel (40.5g), then rinsed with a mixture of dichloromethane (1.01L) and MeOH (1.01L). Combined filtrate was concentrated and azeotroped with toluene (81mL) twice to give a desired diol, which was used in the next reaction without further purification.

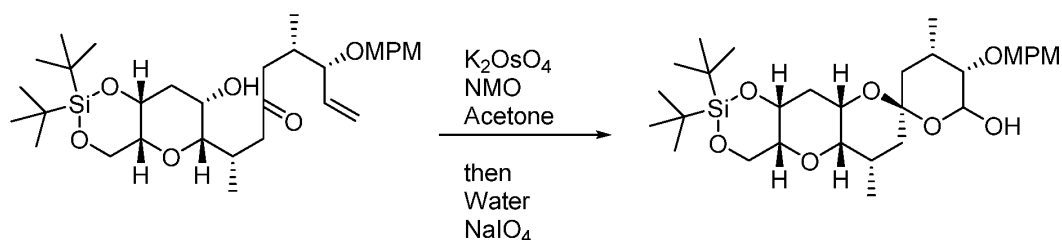


[00896] To a solution of starting material (238mg, 0.313mmol) and 2,4,6-collidine (207 μL , 1.57mmol) in anhydrous dichloromethane (2.38mL) was added trifluoromethanesulfonic anhydride (TiF_2O , 73.7 μL , 0.438mmol) at -78°C under N_2 atmosphere. After stirred for 10min, triethylsilyl trifluoromethanesulfonate (TESOTf, 99 μL , 0.438mmol) was added to the reaction, then warmed to -40°C . After stirred for 70min, NaI (235mg, 1.57mmol) and 1,3-dimethyl-2-imidazolidinone (DMI, 4.75mL) was added, then warmed to room temperature. After stirred for 4hr, toluene (7.13mL) and cold water (4.75mL) was added to the reaction. After separated aqueous layer, organic layer was washed with water (4.75mL), 5wt% citric acid aq. (2.38mL) followed by a mixture of 5wt% NaHCO_3 -10wt% Na_2SO_4 aqueous solution (1.19mL). The organic layer was concentrated under reduced pressure. The residue was purified by column chromatography on silica gel (6g, eluent; 10%, 20%, then 30% EtOAc in *n*-heptane). The collected fractions were concentrated. The residue was crystallized from dichloromethane (0.24mL) and 1-propanol (4.75mL) at 0°C . Resulted precipitation was filtered and washed with 1-propanol to give desired iodide (250mg, 0.254mmol).

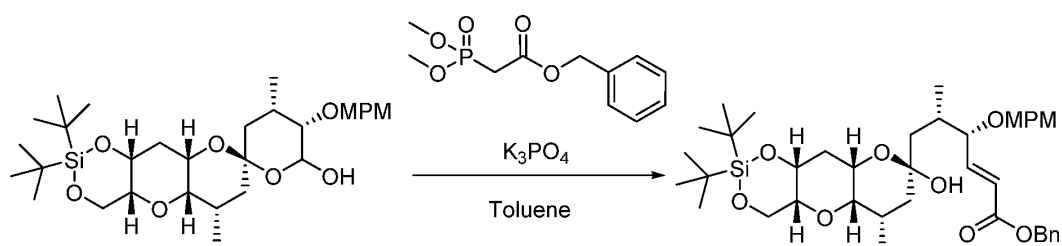


[00897] To a mixture of lactone (55g, 154mmol) and iodide (58.75g, 170mmol) was added THF (550ml) and the reaction mixture was cooled in dry ice-EtOH bath. Sec-BuLi (1.03M solution in cyclohexane, 315mL, 324mmol) was gradually added at $-73.2 \sim -58.5^\circ\text{C}$ during 33min and stirred for 30min. After addition of sat. NH_4Cl (550mL) at $-74.2 \sim -11.2^\circ\text{C}$ during 3min, dry ice bath was removed. The reaction mixture was diluted with AcOEt (550mL) and water (220mL) and stirred in water bath for 40min. The resulting layers were separated and the organic layer was washed with sat. NaCl (165mL) and separated organic layer was concentrated in vacuum. The residue was azeotrope with *n*-heptane (110mL x 3). The

residue was purified by column chromatography on neutral silica gel (1.1kg, eluent: 0%, 5%, 10%, 15% then 25% EtOAc in *n*-heptane) to give desired compound (71.82g, 125mmol, 81%).

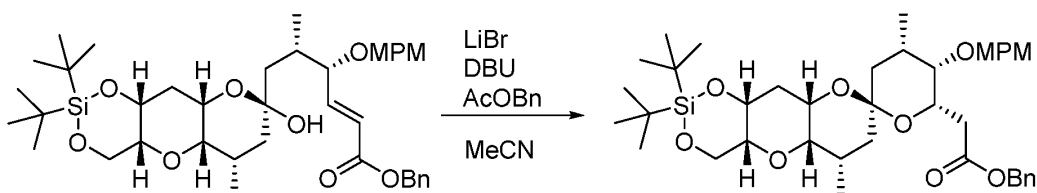


[00898] To a solution of starting material (71.72 g, 124mmol) in acetone (215mL) and water (29mL) was added NMO (29.1g, 249mmol) and $K_2OsO_4 \cdot 2H_2O$ (ca.50%content, 0.182 g, 0.247mmol) at room temperature and stirred for 24.5hrs. The reaction mixture was stirred in water bath and diluted with acetone (143mL) and water (103mL), then $NaIO_4$ (53.2g, 249mmol) and water (40mL for rinsing) was added and stirred at room temperature for 3h. The reaction mixture was diluted with AcOEt (215mL), *n*-heptane (215mL) and water (430mL). The resulting insoluble material was removed by decantation followed by filtration through cotton plug. The filtrate was separated and the aqueous layer was extracted with *n*-heptane/AcOEt = 1/1(v/v) (286mL). The combined organic layer was washed twice with sat. $Na_2S_2O_3$ (215mL) and washed with sat.NaCl (143mL) and concentrated in vacuum. The residue was passed through a pad of neutral silica gel (143g, eluent: 66% EtOAc in *n*-heptane) to give desired compound (71.3g, 123mmol, 99%).



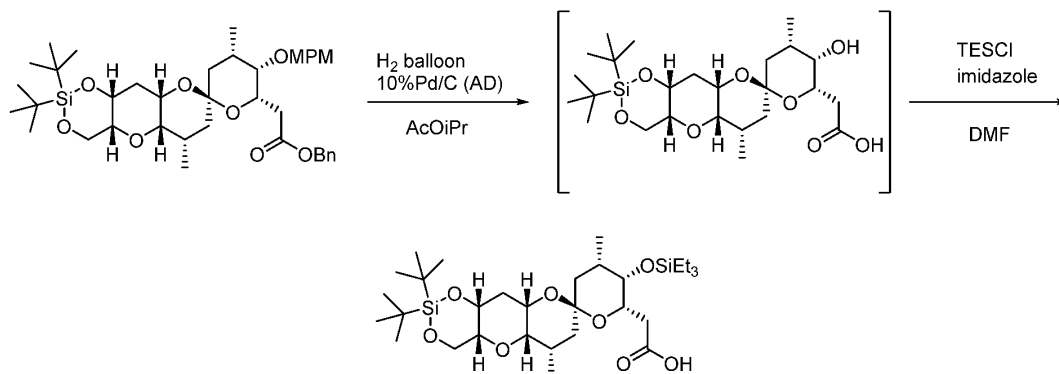
[00899] To a mixture of starting material (71.23 g, 123mmol) and K_3PO_4 (78.4g, 369mmol) and toluene (214mL) was added dimethyl(benzyloxycarbonyl)methyl phosphonate (127.1g, 492mmol). The reaction mixture was purged by N_2 (balloon) and stirred at 30°C for 48hrs. Additional K_3PO_4 (26.13g, 123mmol) and dimethyl(benzyloxycarbonyl)methyl phosphonate (26.1mL, 123mmol) were added and stirred at 30°C for 26hrs. The reaction mixture was

cooled in water bath and diluted with MTBE (712mL) and quenched with sat.NH₄Cl (50mL) at 21~36 °C. The reaction mixture was cooled in ice-water bath and sat.NH₄Cl (662mL) and water (214mL) were successively added. The resulting layers were separated and the aqueous layer was extracted with MTBE (712mL). The combined organic layer was washed with sat.NaCl (356mL) and concentrated in vacuum. The residue was azeotroped with toluene (178mL). The residue was purified by column chromatography on neutral silica gel (1.1kg, eluent: 0%, 8%, 15% then 25% EtOAc in *n*-heptane) to give desired compound (72.63g, 102mmol, 83%, a mixture of *E/Z* isomers). ¹H-NMR (500MHz, CDCl₃) δ: 7.30-7.44 (5H, m), 7.23 (2H, d, *J*=8.6Hz), 6.84-6.92 (3H, m), 6.06 (1H, dd, *J*=15.9, 1.2Hz), 5.18-5.24 (2H, m), 4.50 (1H, d, *J*=11.6Hz), 4.39 (1H, t, *J*=2.8Hz), 4.27 (1H, d, *J*=11Hz), 4.18-4.24 (1H, m), 4.10-4.16 (1H, m), 3.81 (3H, s), 3.72-3.79 (2H, m), 3.60 (1H, d, *J*=10.4Hz), 3.26 (1H, m), 3.01-3.06 (1H, m), 2.73 (1H, dd, *J*= 15.0, 4.6Hz), 2.44-2.59 (2H, m), 2.21-2.39 (4H, m), 1.72 (1H, dt, *J*=14.7, 3.1Hz), 1.05 (18H, s), 0.91 (6H, dd, *J*=14.7, 6.7Hz).



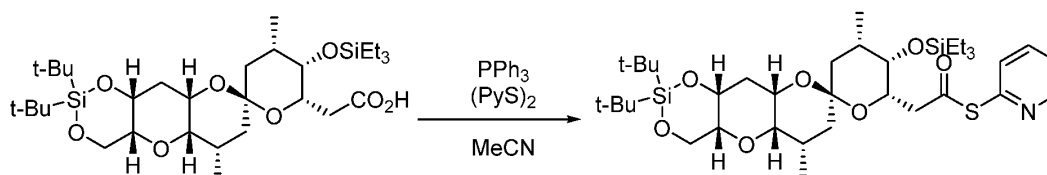
[00900] A solution of the starting material (72.5g, 102mmol) and AcOBn (72.9mL, 510mmol) in MeCN (1087mL) was cooled in ice-water bath. LiBr (88.5g, 1019mmol) was added at 10.1 ~ 16.5°C during 3min and DBU (76.1mL, 510mmol) was added at 11.8 ~ 15.4°C. The reaction mixture was stirred at room temperature for 2hrs. Then, the reaction mixture was stirred at 30°C for 23hrs. The reaction mixture was cooled in ice-water bath and diluted with MTBE (362mL) and quenched with sat.NH₄Cl (362mL) at 7.4 ~ 13.7°C and water (145mL). The resulting layers were separated and the organic layer was washed with sat.NaCl (254mL) and dried over anhydrous Na₂SO₄. After filtration, the filtrate was concentrated in vacuum. The residue was purified by column chromatography on neutral silica gel (1.1kg, eluent: 0%, 3%, 10%, 20% then 35% MTBE in *n*-heptane) to give a desired compound (64.07g, 90mmol, 88%). ¹H-NMR (500MHz, CDCl₃) δ: 7.31-7.40 (5H, m), 7.23-7.27 (2H, m), 6.86 (2H, d, *J*=8.6Hz), 5.01-5.18 (2H, m), 4.41-4.57 (2H, m), 4.24-4.33 (1H, m), 4.16-4.22 (2H, m), 4.08 (1H, ddd, *J*=9.3, 4.7, 1.2Hz), 3.80 (3H, s), 3.51-3.54 (1H, m), 3.25 (1H, brs), 3.15-3.20 (1H, m), 2.97 (1H, d, *J*=2.5Hz), 2.72 (1H, dd, *J*=15.6, 9.5Hz), 2.28-2.42 (2H, m), 2.07 (1H, dt, *J*=15.3, 1.8Hz), 1.87-2.00 (1H, m), 1.69 (1H, dt, *J*=15.3, 4.6Hz),

1.45-1.58 (3H, m), 1.36-1.42 (1H, m), 1.05 (9H, s), 1.03 (9H, s), 0.95 (3H, d, $J=6.7\text{Hz}$), 0.89 (3H, d, $J=6.7\text{Hz}$).

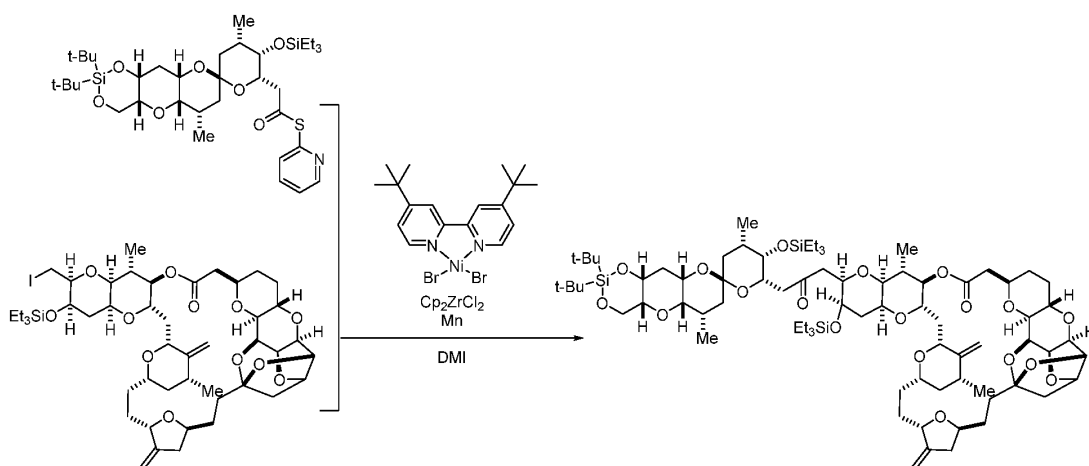


[00901] The starting material (64g, 90mmol) was dissolved in ⁱPrOAc (1280mL) and purged by N₂ (balloon). 10% Pd/C (AD, Kawaken Fine Chemicals, 6.4g) was added and purged by H₂ (balloon) and stirred at room temperature for 24hrs. The catalyst was filtered through Hyflo Supercel (64g) and washed with ⁱPrOAc (1280 mL). The filtrate was concentrated to ca.half volume. ⁱPrOAc (1280 mL) was added and concentrated to ca.half volume. ⁱPrOAc (1280 mL) was added again and concentrated to ca.half volume. DMF (500mL) was added to the residual solution and concentrated in vacuum for 1hr until removal of ⁱPrOAc to give desired seco acid as DMF solution (ca.500mL). This solution of the seco acid (calculated as 90mmol) was diluted with DMF (140mL) and imidazole (36.77g, 540mmol) was added. The reaction mixture was cooled in ice-water bath and TESCl (45.3mL, 270mmol) was slowly added at 2.1 ~ 4.2°C during 13min, then stirred in water bath for 80min. The reaction mixture was cooled in ice-water bath and quenched with sat.NaHCO₃ (384mL) at 4.8 ~ 13.8°C during 24min and stirred in water bath for 1hr. The reaction mixture was diluted with *n*-heptane (640mL) and water (768mL) and the resulting layers were separated. The organic layer was successively washed with sat.NH₄Cl (384mL), water (640mL), 10% NaCl (384mL) and concentrated in vacuum. The residue was purified by column chromatography on neutral silica gel (640g, eluent: 0%, then 67% MTBE in *n*-heptane) to give the desired compound (61.55g). To this compound was added cyclopentyl methyl ether (32mL) and MeCN (94mL) and heated at 50°C to dissolve the solid. To this solution was slowly added MeCN (514mL) during 1hr. The mixture was slowly cooled to room temperature and stirred for 2hrs. After further stirred at 0°C for 1hr followed by -20°C for 14hrs. The resulting suspension was filtrated and rinsed with cold (ca.-20°C) MeCN (92mL). The crystals were dried under

reduced pressure at 40°C to give the desired compound (47.68g, 78mmol, 86%) as a white solid. $^1\text{H-NMR}$ (500MHz, CDCl_3) δ : 4.13-4.31 (3H, m), 3.96-4.06 (1H, m), 3.49-3.65 (2H, m), 3.12-3.24 (2H, m), 2.69 (1H, dd, $J=15.8, 10.0\text{Hz}$), 2.39 (1H, dd, $J=15.8, 3.7\text{Hz}$), 2.20-2.31 (1H, m), 2.08-2.15 (1H, m), 1.98-2.07 (1H, m), 1.81 (1H, dt, $J=15.2, 4.4\text{Hz}$), 1.47-1.57 (3H, m), 1.39-1.45 (1H, m), 1.03 (9H, s), 1.02 (9H, s), 0.94-0.99 (12H, m), 0.86 (3H, d, $J=6.9\text{Hz}$), 0.59-0.66 (6H, m).



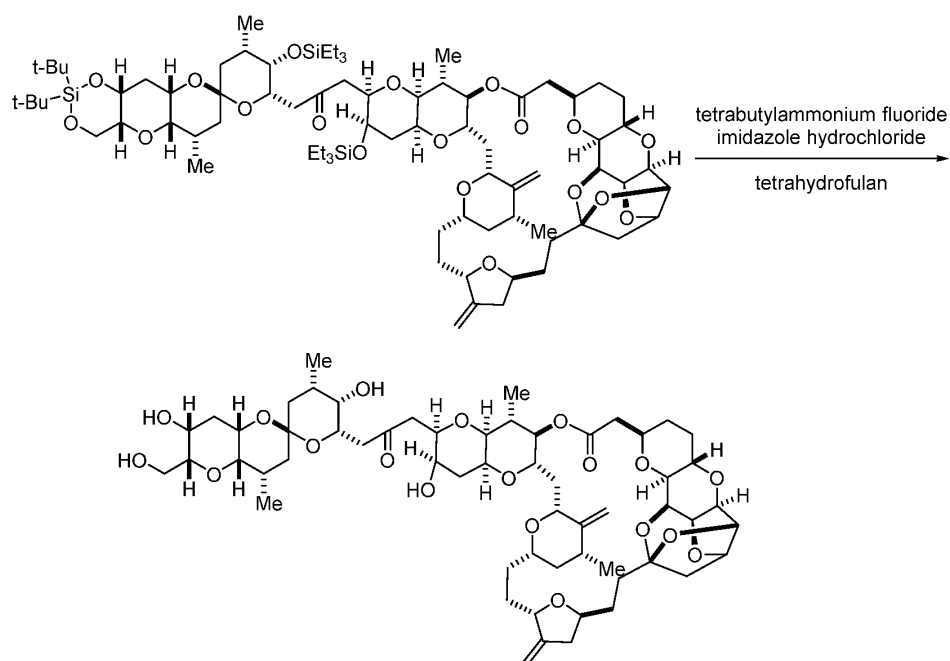
[00902] To a solution of starting material (35.1g, 57.1mmol) and triphenylphosphine (35.8g, 136mmol) in dehydrated acetonitrile (350mL) was added 2,2'-dipyridyldisulfide (15.1g, 68.5mmol) at 0°C (bath) under nitrogen atmosphere. After warmed up to 25°C, the mixture was stirred for 17.5h and the reaction was quenched with *n*-heptane (1050mL) and water (350mL) to give a biphasic mixture. The organic layer was separated and washed with 5 times 75vol% acetonitrile aqueous solution (351mL) and once with saturated sodium chloride aqueous solution (175mL). Then, the organic layer was concentrated to dryness. The residue was purified by column chromatography on silica-gel (1900g, eluent: 0%, 10% ethyl acetate in *n*-heptane) to give desired compound (38.6g, 54.5mmol, 96%).



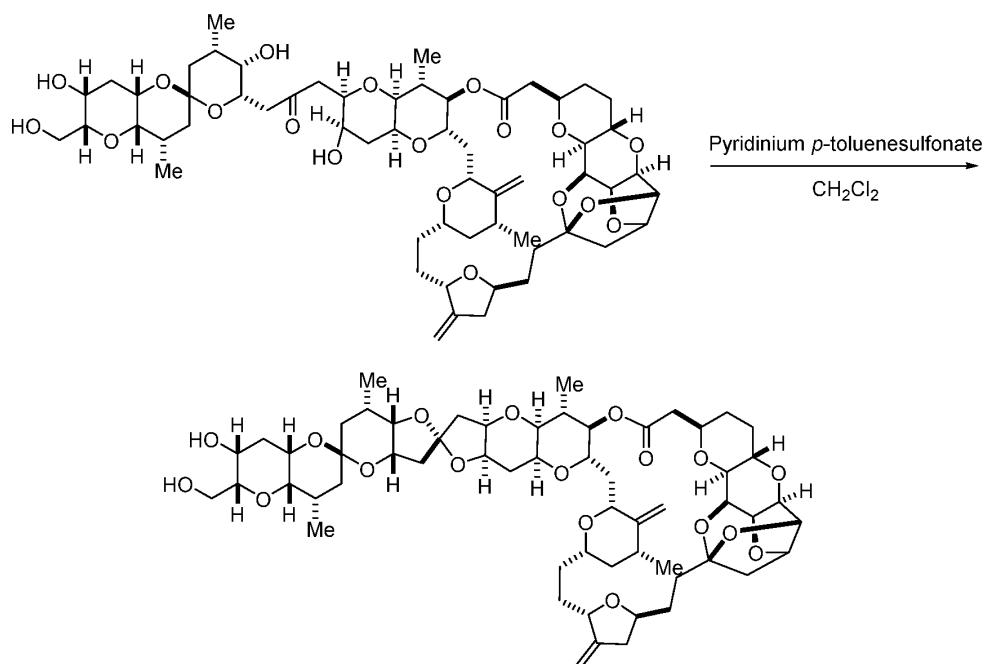
[00903] *Preparation of nickel catalyst solution.* A solution of nickel(II) bromide (5.00g, 22.9mmol) and 4,4'-di-*tert*-butyl-2,2'-dipyridyl (6.15g, 22.9mmol) in dehydrated ethanol

(100mL) was stirred at 80°C for 18h under argon atmosphere. After the reaction mixture was cooled to room temperature, the reaction mixture was concentrated to dryness. The residue was dried in vacuum at 150°C (bath) for 70h. Then, dehydrated DMI (62mL) was added to give nickel catalyst solution (used as 0.37mol/L). See, *e.g.*, Dr. Daniel J. Weix et.al., Chem. Eur.J.2016, 22,11564 –11567)

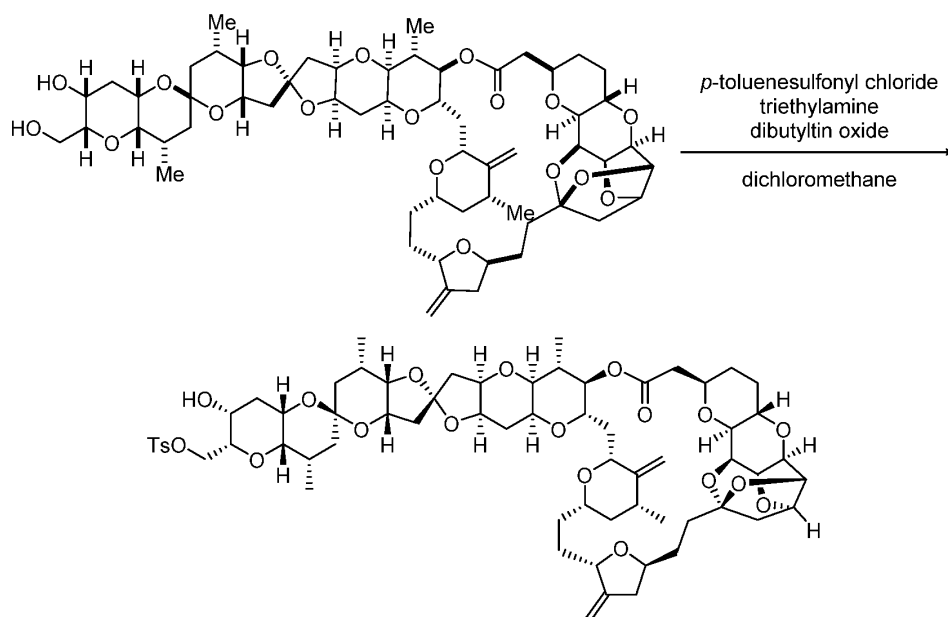
[00904] *Coupling Reaction.* To a solution of iodide (30.1g, 30.6mmol), thioester (30.4g, 42.9mmol), bis(cyclopentadienyl)zirconium(IV) dichloride (31.3g, 107mmol) and manganese (11.7g, 214mmol) in dehydrated DMI (270mL) was added nickel catalyst solution (30mL, 0.37mol/L in DMI, 11.1mmol) prepared above under argon atmosphere. After being stirred at 30°C (bath) for 4h under argon atmosphere, the reaction was quenched with heptane (600mL), 10wt% citric acid aqueous solution (600mL) below 22°C (0°C bath). Then Celite®545 (15.0g) was added and the mixture was stirred at 0°C for 1h. Then the mixture was passed through short pad of Celite®545 (60.0g) and the Celite®545 pad was washed with heptane (1500mL) to give biphasic mixture. The aqueous layer was separated and extracted with heptane (90mL). The combined organic layer was washed with a mixture of 5wt% sodium bicarbonate and 10wt% sodium sulfate aqueous solution (150mL) and then concentrated to dryness. The residue was purified by column chromatography on silica-gel (1500g, eluent: 10%, then 30% ethyl acetate in heptane) to give desired compound (36.6g, 25.2mmol, 82%).



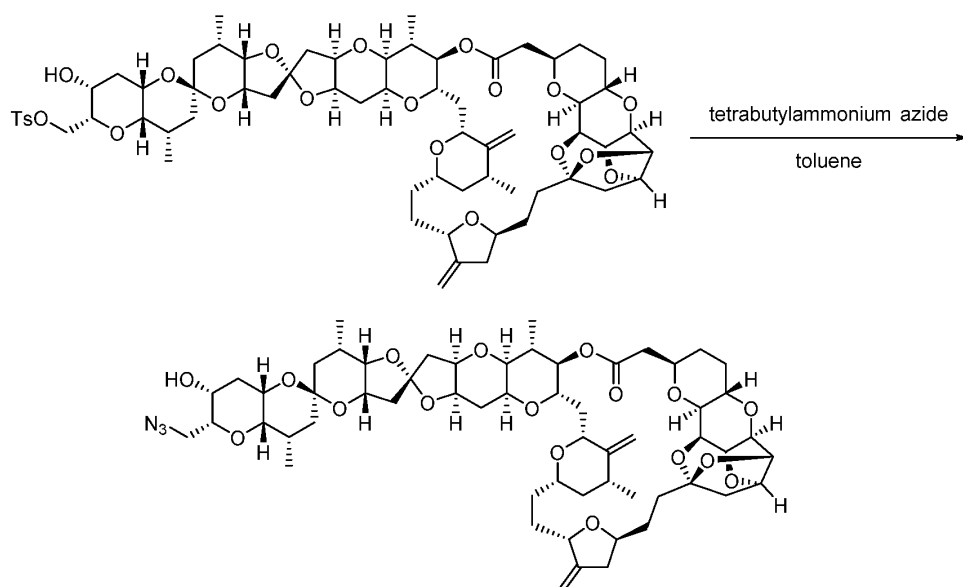
[00905] To a solution of starting material (35.5g, 24.4mmol) in THF (365mL) was added a mixture of imidazole hydrochloride (12.9g, 123mmol) and tetrabutylammonium fluoride (244mL, 1.0mol/L in THF, 244mmol) at 25°C (bath) under nitrogen atmosphere. After being stirred at 25°C for 19h, the reaction was quenched with toluene (710mL) and 10wt% sodium chloride aqueous solution (710mL) to give a biphasic mixture. The aqueous layer was separated and extracted with toluene (178mL). The combined organic layer was washed with water (71mL) and then concentrated to dryness. The residue was purified by column chromatography on silica gel (1500g, eluent: 10% to 100% ethyl acetate in heptane, then, 10% methanol in ethyl acetate) to give desired compound (25.5g, 23.5mmol, 96%).



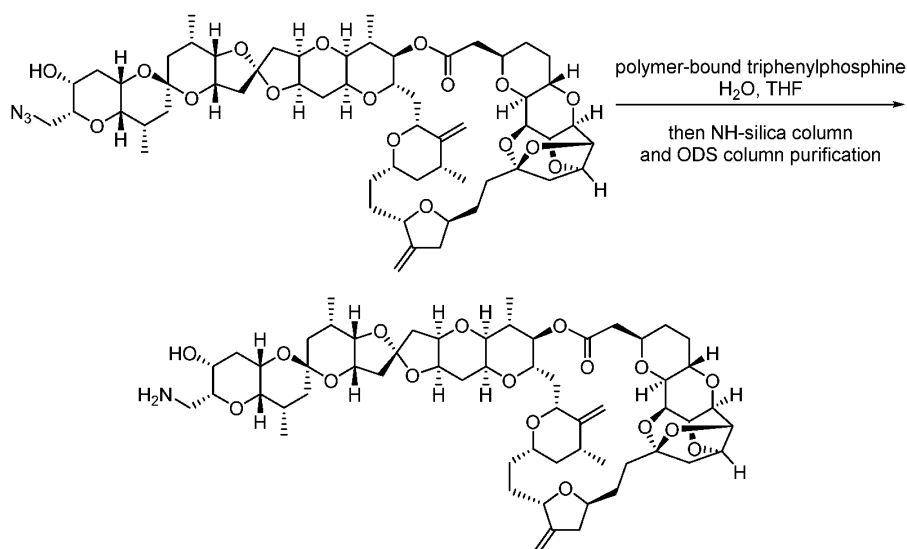
[00906] To a solution of desired compound (25.4g, 23.4mmol) in dichloromethane (765mL) was added pyridinium *p*-toluenesulfonate (29.5g, 117mmol) at 10°C under nitrogen atmosphere. After being stirred at 9-11°C for 3h, the reaction was quenched with water (508mL) to give a biphasic mixture. The aqueous layer was re-extracted with dichloromethane (25mL). The combined organic layer was washed with water (508mL) followed by a mixture of 5wt% sodium bicarbonate and 10wt% sodium sulfate aqueous solution (127mL). Then, the organic layer was concentrated to dryness. The residue was purified by column chromatography on silica gel (1500g, eluent: 10%, 95%, 100% ethyl acetate in heptane) to give the desired compound (23.8g). This compound was purified again by column chromatography on ODS (950g, eluent: 30%, 65%, 100% acetonitrile in water). The fractions containing desired compound were combined and acetonitrile was removed to some extent under reduced pressure. The resulting aqueous solution was extracted with ethyl acetate (1524mL + 762mL) and the combined organic layer was concentrated to dryness to give desired compound (18.1g, 17.0mmol, 73%).



[00907] To a solution of starting material (15.2g, 14.2mmol) in dichloromethane (30mL) was added triethylamine (12.3mL, 88.5mmol), dibutyltin oxide (2.13g, 8.56mmol) and *p*-toluenesulfonyl chloride (8.16g, 42.8mmol) successively at 25°C. After being stirred at 25°C for 3h, the reaction mixture was passed through glass-filter. The filtrate was directly purified by column chromatography on silica gel (750g, eluent: 10%, 30%, 50%, 70%, 85% ethyl acetate in heptane) to give desired compound (17.4g, 14.2mmol, quant.). ¹H NMR (600 MHz, BENZENE-*d*₆) δ ppm 0.86 (d, *J*=6.8 Hz, 3 H) 0.95 (d, *J*=6.8 Hz, 3 H) 1.00 - 1.05 (m, 4 H) 1.07 - 1.17 (m, 4 H) 1.19 (br td, *J*=14.7, 3.0 Hz, 1 H) 1.26 - 1.49 (m, 8 H) 1.54 (t, *J*=13.0 Hz, 1 H) 1.61 - 1.67 (m, 2 H) 1.75 - 2.39 (m, 25 H) 2.62 - 2.75 (m, 5 H) 3.25 (br s, 1 H) 3.27 (br s, 1 H) 3.35 (t, *J*=6.0 Hz, 1 H) 3.39 (dt, *J*=10.8, 5.6 Hz, 1 H) 3.44 (br d, *J*=11.0 Hz, 1 H) 3.50 (d, *J*=11.3 Hz, 1 H) 3.55 (br s, 1 H) 3.62 (dd, *J*=6.2, 4.3 Hz, 1 H) 3.78 (br t, *J*=9.6 Hz, 1 H) 3.83 (br t, *J*=6.0 Hz, 1 H) 3.86 - 3.96 (m, 3 H) 3.99 - 4.11 (m, 3 H) 4.14 (m, 1 H) 4.27 (br s, 1 H) 4.36 - 4.44 (m, 2 H) 4.49 (m, 1 H) 4.55 - 4.61 (m, 2 H) 4.80 (s, 1 H) 4.86 - 4.91 (m, 2 H) 4.95 (br s, 1 H) 5.02 (br s, 1 H) 6.65 (d, *J*=7.9 Hz, 2 H) 7.80 (d, *J*=7.9 Hz, 2 H).



[00908] To a solution of starting material (17.3g, 14.2mmol) in toluene (121mL) was added tetrabutylammonium azide (32.2g, 113mmol) at room temperature under nitrogen atmosphere. After being stirred at 100°C (bath) for 5h, the reaction mixture was cooled to room temperature and tetrahydrofuran (52mL) was added to give a homogenous solution. The solution was directly purified by column chromatography on silica gel (750g, eluent: 10%, 50%, 70%, 80% ethyl acetate in heptane) to give desired compound (15.5g, 14.2mmol, quant.). ¹H NMR (600 MHz, BENZENE-d₆) δ ppm 0.97 - 1.05 (m, 9 H) 1.08 - 1.37 (m, 8 H) 1.39 - 1.55 (m, 5 H) 1.58 - 1.71 (m, 3 H) 1.78 - 1.88 (m, 2 H) 1.91 - 2.08 (m, 8 H) 2.09 - 2.40 (m, 13 H) 2.62 - 2.75 (m, 4 H) 2.78 (d, J=2.3 Hz, 1 H) 2.97 (dd, J=13.0, 4.0 Hz, 1 H) 3.09 (dd, J=8.3, 3.8 Hz, 1 H) 3.26 (br d, J=11.3 Hz, 1 H) 3.30 (t, J=2.6 Hz, 1 H) 3.32 (br s, 1 H) 3.40 (dt, J=10.8, 5.6 Hz, 1 H) 3.56 (d, J=11.0 Hz, 1 H) 3.58 - 3.65 (m, 3 H) 3.78 (br dd, J=10.8, 8.9 Hz, 1 H) 3.83 (br dd, J=7.2, 5.7 Hz, 1 H) 3.86 - 3.99 (m, 3 H) 4.01 - 4.11 (m, 3 H) 4.14 (t, J=4.5 Hz, 1 H) 4.28 (dd, J=3.8, 1.5 Hz, 1 H) 4.50 (m, 1 H) 4.56 - 4.62 (m, 2 H) 4.80 (br s, 1 H) 4.86 - 4.92 (m, 2 H) 4.96 (br d, J=1.9 Hz, 1 H) 5.02 (br d, J=1.5 Hz, 1 H).



[00909] To a solution of starting material (15.4g, 14.1mmol) in a mixture of THF (92mL) and water (10mL) was added polymer-bound triphenylphosphine (20.4g, 2.07mmol/g, 42.2mmol) at 25°C. After being stirred at 25°C for 70h, the reaction mixture was passed through glass-filter. The filtrate was concentrated to dryness. The residue was purified by column chromatography on NH-silica-gel (900g, eluent: 50%, 100% MTBE in heptane, then 3%, 10% methanol MTBE) to give desired compound (14.1g, 13.2mmol, 94%) as a white solid. The desired compound was further purified by column chromatography on ODS (400g, eluent: 1%, 70%, 100% 0.05vol% acetic acid/acetonitrile in 0.05vol% acetic acid/water). The fractions containing desired compound were combined and dichloromethane (3304mL), 20wt% sodium chloride aqueous solution (661mL) and saturated sodium bicarbonate aqueous solution (1322mL) were added to give biphasic mixture. Then, the aqueous layer was extracted with dichloromethane (830mL). The combined organic layer was washed with water (165mL) and concentrated to dryness to give desired compound (12.4g, 11.6mmol, 88%). After the compound was dissolved in dichloromethane (25mL) and pentane (25mL), half of the solution was added to pentane (984mL) at 0°C with stirring to precipitate desired compound. After removal of approximately 271mL of solvent under slightly reduced pressure, pentane (246mL) was added. After removal of approximately 246mL of solvent under slightly reduced pressure, pentane (246mL) was added. To this mixture was added remaining half of the desired compound solution at 0°C with stirring to further precipitate the desired compound. After removal of approximately 271mL of solvent under slightly reduced pressure, pentane (246mL) was added. After removal of approximately 246mL of solvent under slightly reduced pressure, pentane (246mL) was added. The resulting slurry was

filtrated, dried at 40°C gave the desired compound (11.5g, 93%).

¹H NMR (600 MHz, METHANOL-*d*₄) δ ppm 0.98 (d, *J*=7.2 Hz, 3 H) 1.00 (d, *J*=6.8 Hz, 3 H) 1.02 (m, 1 H) 1.05 (d, *J*=6.8 Hz, 3 H) 1.09 (d, *J*=6.4 Hz, 3 H) 1.28 - 1.45 (m, 5 H) 1.46 - 1.59 (m, 4 H) 1.57 - 1.63 (m, 1 H) 1.65 - 1.71 (m, 1 H) 1.70 - 1.75 (m, 2 H) 1.79 - 1.86 (m, 2 H) 1.91 (dt, *J*=14.9, 3.1 Hz, 1 H) 1.94 - 2.11 (m, 8 H) 2.14 - 2.34 (m, 9 H) 2.39 (dd, *J*=13.2, 6.0 Hz, 1 H) 2.44 (dd, *J*=17.4, 1.9 Hz, 1 H) 2.56 (dd, *J*=17.6, 9.6 Hz, 1 H) 2.69 (dd, *J*=13.2, 4.2 Hz, 1 H) 2.79 (ddq, *J*=15.9, 7.6, 2.0 Hz, 1 H) 2.92 (dd, *J*=13.2, 8.3 Hz, 1 H) 2.97 (dd, *J*=9.6, 1.7 Hz, 1 H) 3.21 (dd, *J*=6.4, 4.9 Hz, 1 H) 3.29 (m, 1 H) 3.34 (dd, *J*=8.3, 4.15 Hz, 1 H) 3.58 (br. s., 1 H) 3.60 (br.d, *J*=11.3 Hz, 1 H) 3.68 - 3.73 (m, 2 H) 3.80 (br. s., 1 H) 3.84 - 3.90 (m, 2 H) 3.98 (d, *J*=2.3 Hz, 1 H) 4.03 - 4.13 (m, 4 H) 4.17 (dd, *J*=6.4, 4.9 Hz, 1 H) 4.24 (ddd, *J*=11.3, 4.5, 1.5 Hz, 1 H) 4.29 (dd, *J*=4.0, 1.9 Hz, 1 H) 4.32 (td, *J*=10.2, 4.2 Hz, 1 H) 4.44 (br. d, *J*=11.0 Hz, 1 H) 4.59 (t, *J*=4.5 Hz, 1 H) 4.62 (dd, *J*=7.4, 4.7 Hz, 1 H) 4.69 (t, *J*=4.7 Hz, 1 H) 4.80 (br. s., 1 H) 4.87 (s, 1 H) 5.00 (br. s., 1 H) 5.05 (br.d, *J*=1.1 Hz, 1 H).

EQUIVALENTS AND SCOPE

[0001] In the claims articles such as “a,” “an,” and “the” may mean one or more than one unless indicated to the contrary or otherwise evident from the context. Claims or descriptions that include “or” between one or more members of a group are considered satisfied if one, more than one, or all of the group members are present in, employed in, or otherwise relevant to a given product or process unless indicated to the contrary or otherwise evident from the context. The invention includes embodiments in which exactly one member of the group is present in, employed in, or otherwise relevant to a given product or process. The invention includes embodiments in which more than one, or all of the group members are present in, employed in, or otherwise relevant to a given product or process.

[0002] Furthermore, the invention encompasses all variations, combinations, and permutations in which one or more limitations, elements, clauses, and descriptive terms from one or more of the listed claims is introduced into another claim. For example, any claim that is dependent on another claim can be modified to include one or more limitations found in any other claim that is dependent on the same base claim. Where elements are presented as lists, *e.g.*, in Markush group format, each subgroup of the elements is also disclosed, and any element(s) can be removed from the group. It should be understood that, in general, where the invention, or aspects of the invention, is/are referred to as comprising particular elements and/or features, certain embodiments of the invention or aspects of the invention consist, or

consist essentially of, such elements and/or features. For purposes of simplicity, those embodiments have not been specifically set forth *in haec verba* herein.

[0003] It is also noted that the terms “comprising” and “containing” are intended to be open and permits the inclusion of additional elements or steps. Where ranges are given, endpoints are included. Furthermore, unless otherwise indicated or otherwise evident from the context and understanding of one of ordinary skill in the art, values that are expressed as ranges can assume any specific value or sub-range within the stated ranges in different embodiments of the invention, to the tenth of the unit of the lower limit of the range, unless the context clearly dictates otherwise.

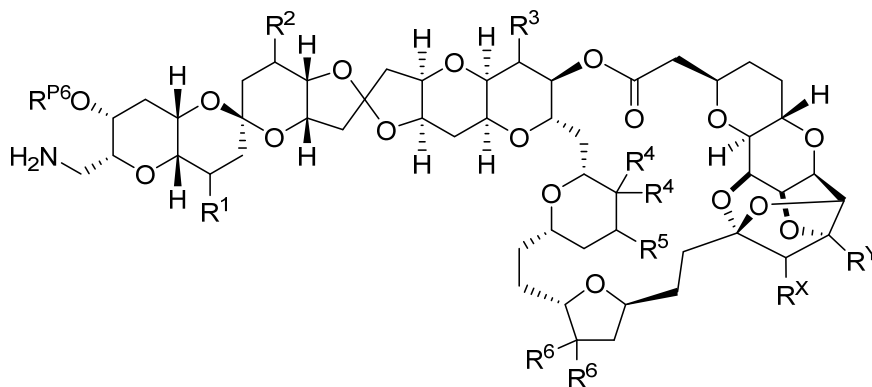
[0004] This application refers to various issued patents, published patent applications, journal articles, and other publications, all of which are incorporated herein by reference. If there is a conflict between any of the incorporated references and the instant specification, the specification shall control. In addition, any particular embodiment of the present invention that falls within the prior art may be explicitly excluded from any one or more of the claims. Because such embodiments are deemed to be known to one of ordinary skill in the art, they may be excluded even if the exclusion is not set forth explicitly herein. Any particular embodiment of the invention can be excluded from any claim, for any reason, whether or not related to the existence of prior art.

[0005] Those skilled in the art will recognize or be able to ascertain using no more than routine experimentation many equivalents to the specific embodiments described herein. The scope of the present embodiments described herein is not intended to be limited to the above Description, but rather is as set forth in the appended claims. Those of ordinary skill in the art will appreciate that various changes and modifications to this description may be made without departing from the spirit or scope of the present invention, as defined in the following claims.

CLAIMS

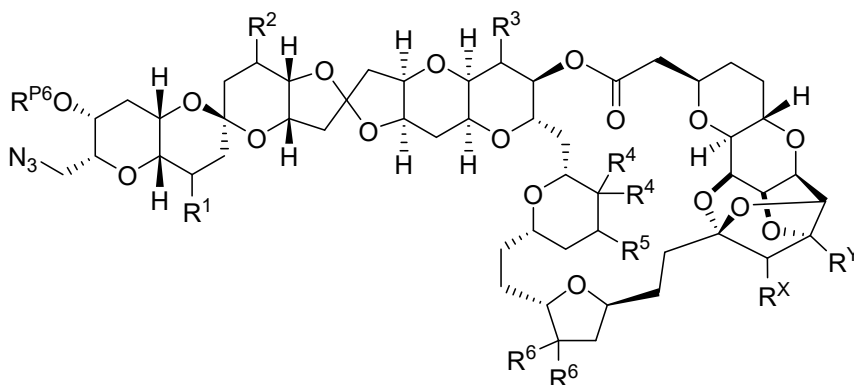
What is claimed is:

1. A method of preparing a compound of Formula (H3-A):



(H3-A),

or a salt thereof, the method comprising reducing a compound of Formula (H3-N3):



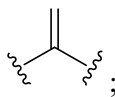
(H3-N3),

or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

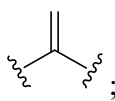
each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^4 groups are taken together to form:



each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^6 groups are taken together to form:



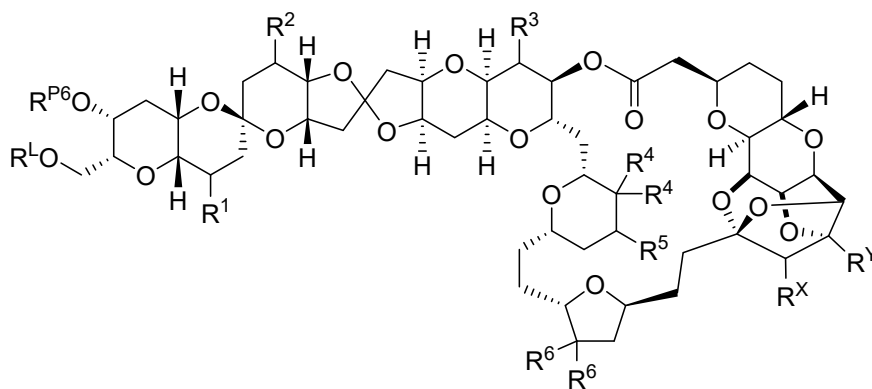
R^{P6} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

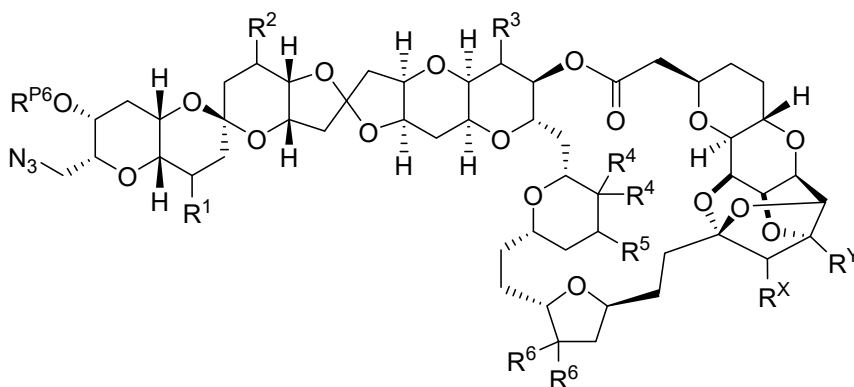
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocycl.

2. The method of claim 1 further comprising a step of reacting a compound of Formula (H3-L):



(H3-L),

or a salt thereof, in the presence of an azide, to yield a compound of Formula (H3-N3):

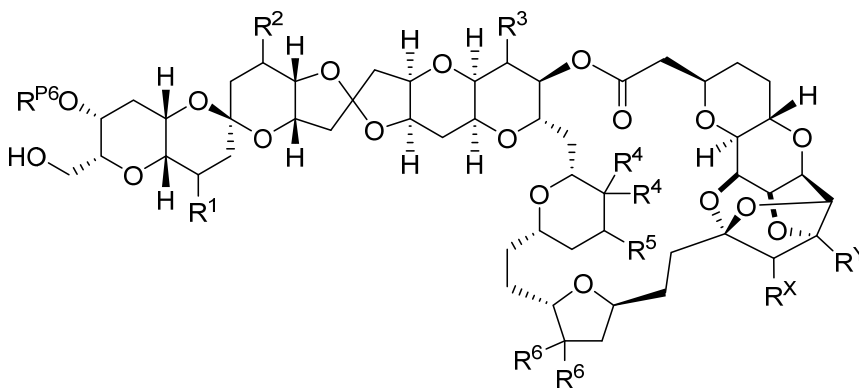


(H3-N3),

or a salt thereof, wherein:

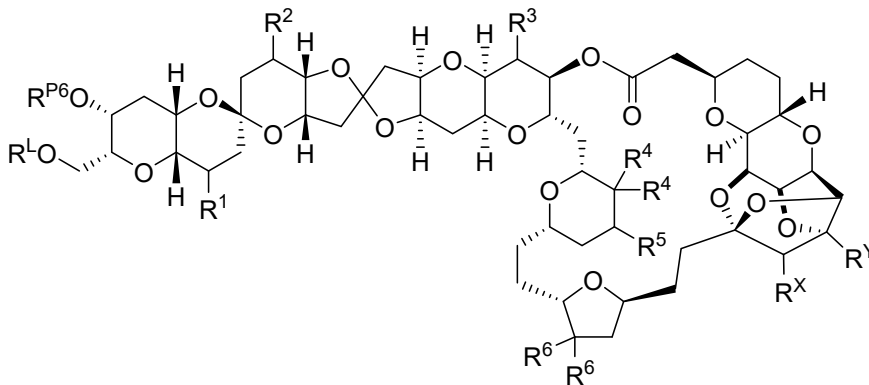
R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

3. The method of claim 2 further comprising a step of reacting a compound of Formula **(H3-OH)**:



(H3-OH),

or a salt thereof, in the presence of a reagent of the formula X^L-R^L , to yield a compound of Formula **(H3-L)**:

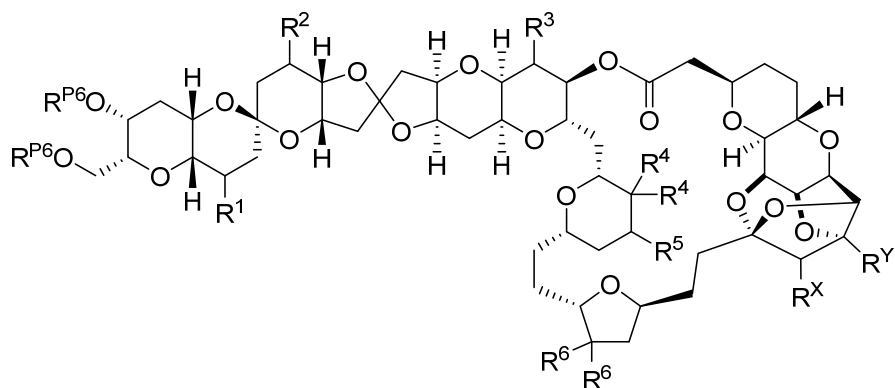


(H3-L),

or a salt thereof, wherein:

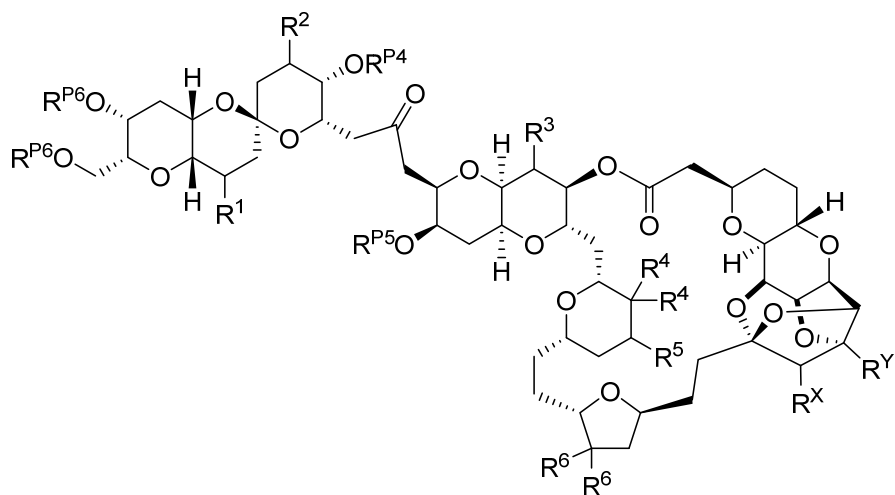
X^L is halogen or a leaving group.

4. A method of preparing a compound of Formula (H3-2-I):



(H3-2-I),

or a salt thereof, the method comprising cyclizing a compound of Formula (H3-2-II):



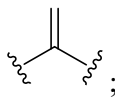
(H3-2-II),

or a salt thereof, wherein:

R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

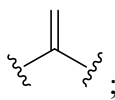
each instance of R⁴ is independently hydrogen, halogen, or optionally substituted alkyl,

or two R⁴ groups are taken together to form:



each instance of R⁶ is independently hydrogen, halogen, or optionally substituted alkyl,

or two R⁶ groups are taken together to form:



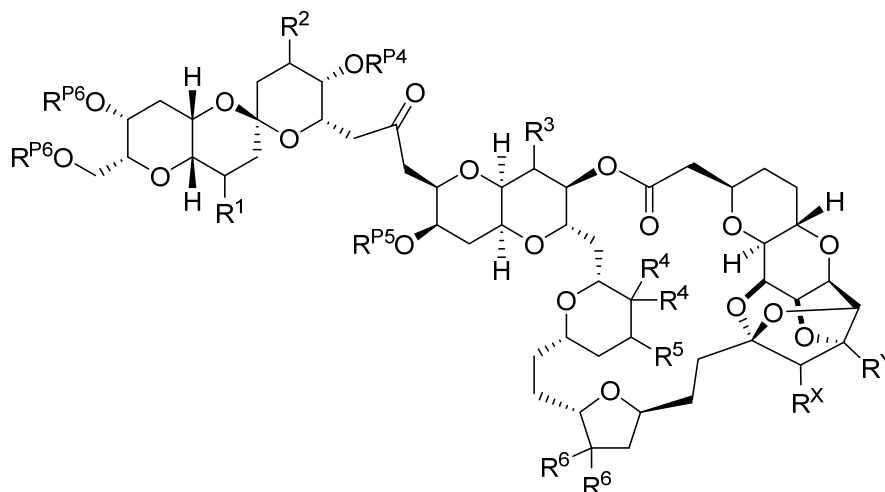
R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

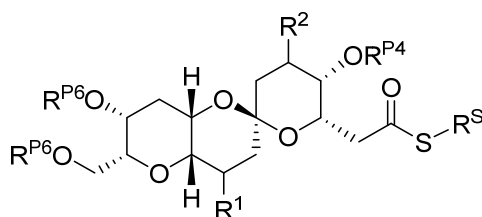
optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

5. A method of preparing a compound of Formula (H3-2-II):



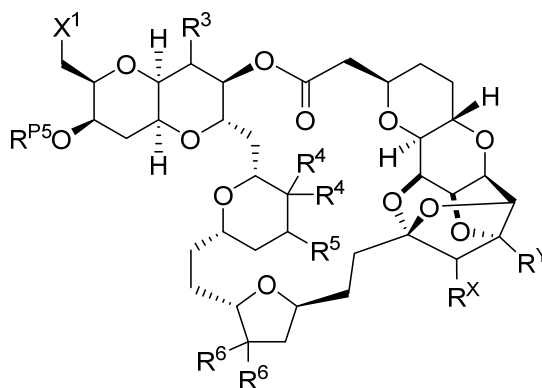
(H3-2-II),

or a salt thereof, the method comprising coupling a compound of Formula (L-2-6):



(L-2-6),

or a salt thereof, with a compound of Formula (**R-2-I**):



(**R-2-I**),

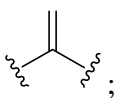
or a salt thereof, to yield a compound of Formula (**H3-2-II**), or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

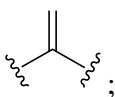
X¹ is halogen or a leaving group;

R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R⁴ is independently hydrogen, halogen, or optionally substituted alkyl,

or two R⁴ groups are taken together to form: ;

each instance of R⁶ is independently hydrogen, halogen, or optionally substituted alkyl,

or two R⁶ groups are taken together to form: ;

R^{P4}, R^{P5}, and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

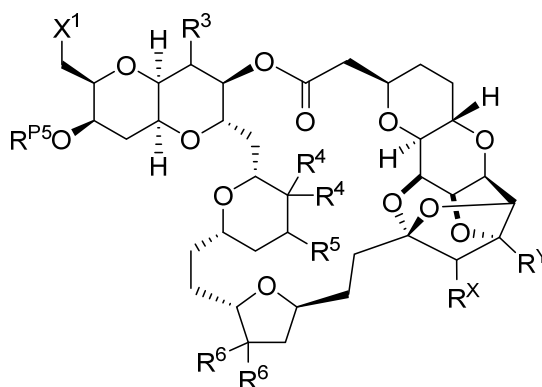
R^X is hydrogen or -OR^{Xa}, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or -OR^{Ya}, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl.

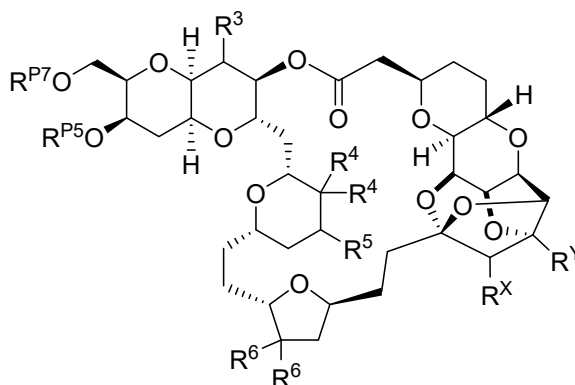
6. The method of claim 5 further comprising a step of deprotecting one or more oxygen atoms of the compound of Formula (H3-2-II), or a salt thereof.

7. The method of claim 5 further comprising preparing a compound of Formula (R-2-I):



(R-2-I),

or a salt thereof, by reacting a compound of Formula (R-4-11B):

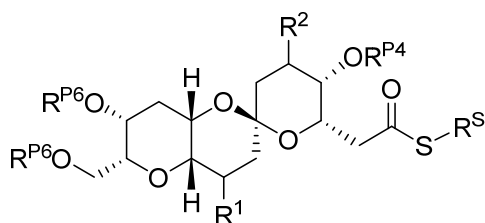


(R-4-11B),

or a salt thereof, with a nucleophile, thereby substituting the group $-OR^{P7}$ with the group $-X^1$; wherein:

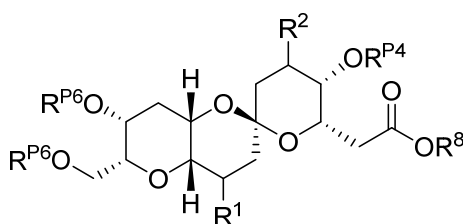
R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group.

8. The method of claim 5 further comprising preparing a compound of Formula (L-2-6):



(L-2-6),

or a salt thereof, by reacting a compound of Formula (L-5-7B):

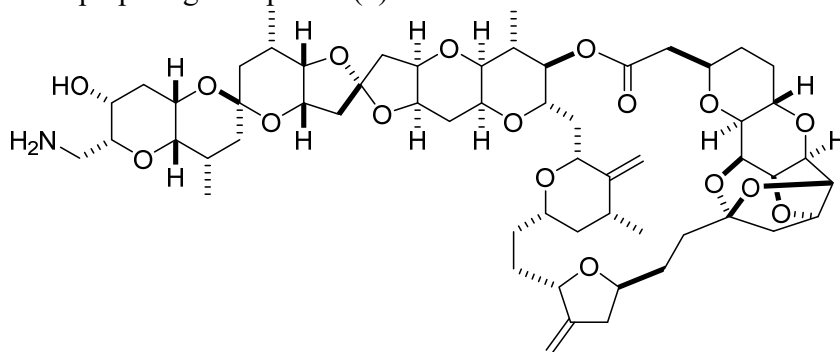


(L-5-7B),

or a salt thereof, with a thiolating agent; wherein:

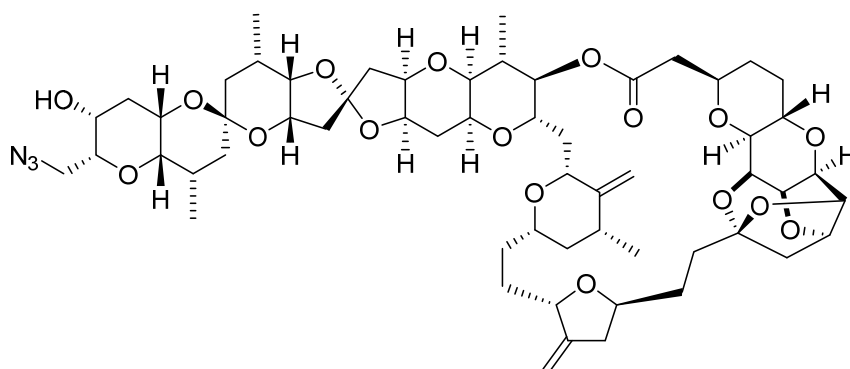
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

9. A method of preparing Compound (1):



(1),

or a salt thereof, the method comprising reducing Compound (B):



(B),

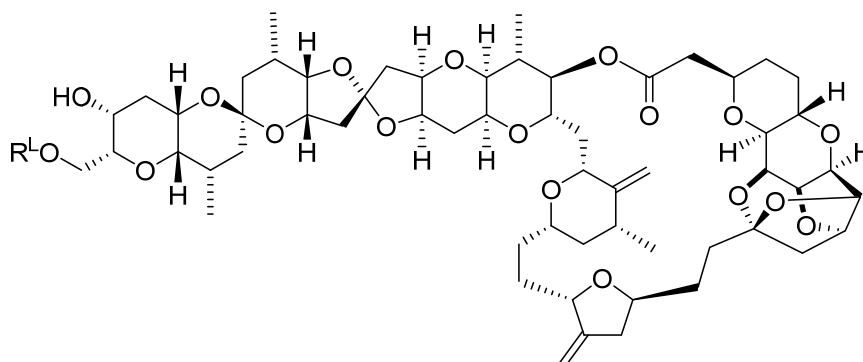
or a salt thereof.

10. The method of claim 1 or 9, wherein the step of reducing is carried out in the presence of a phosphine reagent; optionally wherein the phosphine reagent is triphenylphosphine (Ph_3P); optionally wherein the phosphine reagent is polymer-bound triphenylphosphine; and optionally wherein at least one equivalent of the phosphine reagent is present.

11. The method of any one of claims 1, 9, and 10, wherein the step of reducing is carried out in the presence of THF and water.

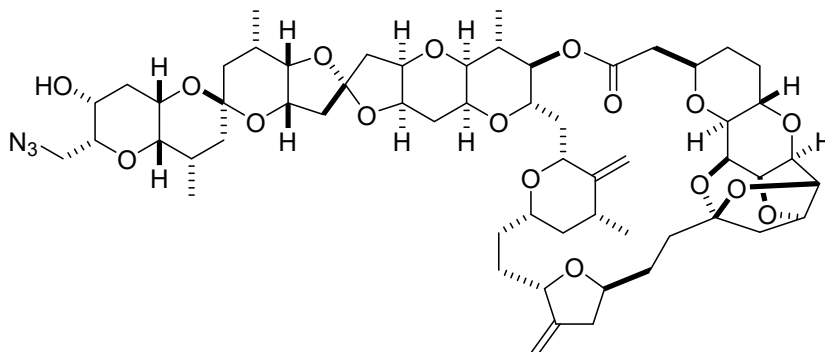
12. The method of any one of claims 1 and 9-11, wherein the compound of Formula (**H3-A**), Compound (1), or salt thereof is purified and isolated by precipitation.

13. The method of any one of claims 9-12 further comprising a step of reacting a compound of Formula (A):



(A),

or a salt thereof, in the presence of an azide, to yield Compound (B):



(B),

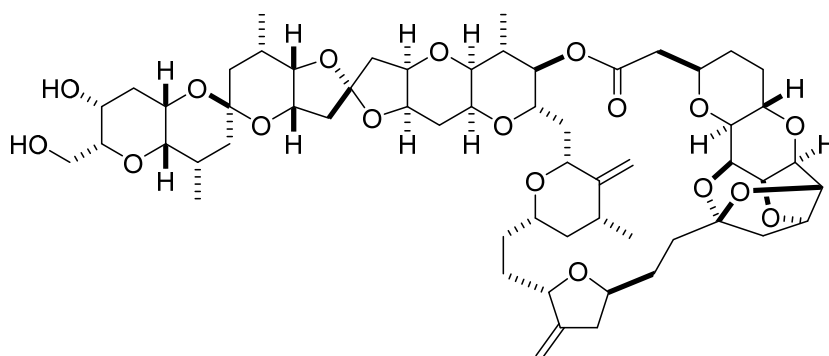
or a salt thereof, wherein:

R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

14. The method of claim 2 or 13, wherein the azide is sodium azide, potassium azide, or a tetraalkylammonium azide salt; optionally wherein the azide is a tetraalkylammonium azide; and optionally wherein the azide is tetrabutylammonium azide ([*n*-Bu₄N]N₃).

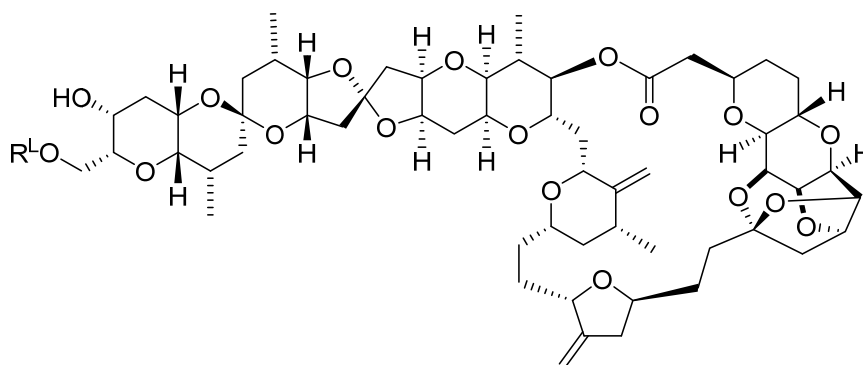
15. The method of any one of claims 2, 13, or 14, wherein the reaction is carried out in toluene.

16. The method of any one of claims 13-15 further comprising a step of reacting Compound (2):



(2),

or a salt thereof, in the presence of a reagent of the formula X^L-R^L , to yield a compound of Formula (A):



(A),

or a salt thereof, wherein:

X^L is halogen or a leaving group.

17. The method of claim 3 or 16, wherein the reagent of the formula X^L-R^L is a sulfonyl halide; R^L is optionally substituted sulfonyl; and X^L is halogen; optionally wherein reagent of the formula X^L-R^L is a sulfonyl chloride; and R^L is optionally substituted sulfonyl; and optionally wherein reagent of the formula X^L-R^L is tosyl chloride (TsCl).

18. The method of any one of claims 3, 16, or 17, wherein the reaction is carried out in the presence of a base; optionally wherein the base is a trialkylamine base, a pyridine base, or an imidazole base; optionally wherein the base is a trialkylamine base; and optionally

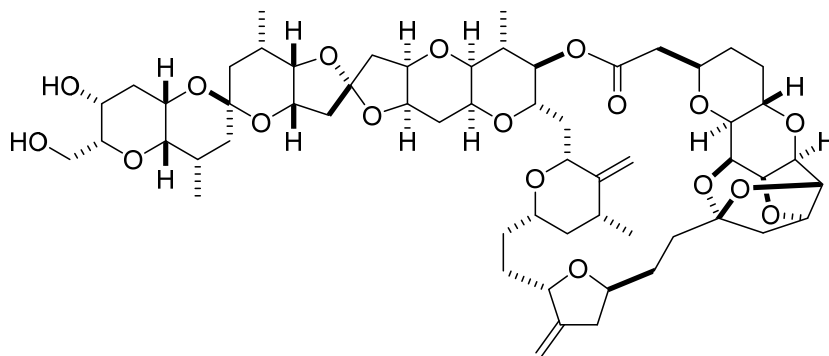
wherein the base is triethylamine (TEA).

19. The method of any one of claims 3 and 16-18, wherein the reaction is carried out in the presence of a Lewis acid; optionally wherein the reaction is carried out in the presence of dibutyltin oxide.

20. The method of any one of claims 3 and 16-19, wherein the reaction is carried out in the presence of TsCl, TEA, and dibutyltin oxide.

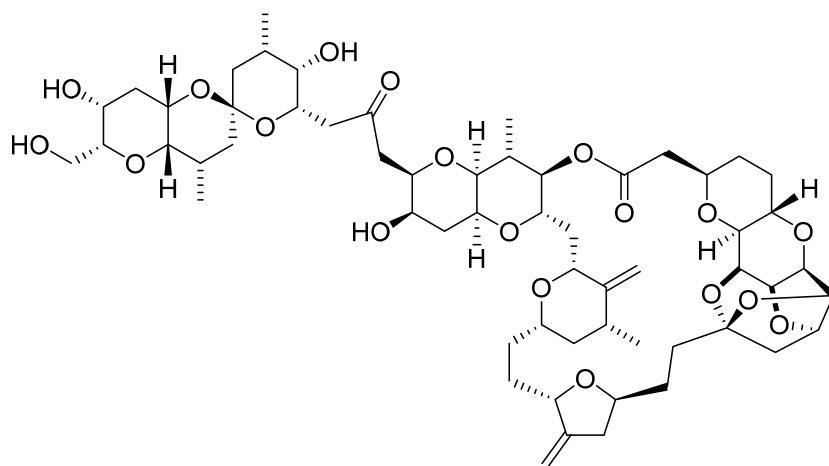
21. The method of any one of claims 3 and 16-20, wherein the reaction is carried out in dichloromethane (DCM).

22. A method of preparing Compound (2):



(2),

or a salt thereof, the method comprising cyclizing Compound (C):



(C),

or a salt thereof.

23. The method of claim 4 or 22, wherein the step of cyclizing is carried out in the presence of an acid; optionally

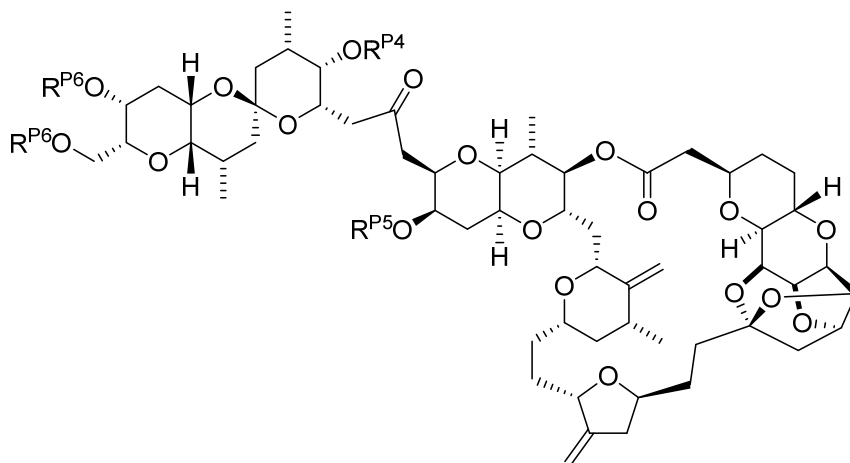
wherein the acid is a pyridinium salt; and optionally

wherein the acid is pyridinium *p*-toluenesulfonate (PPTS).

24. The method of any one of claims 4, 22, and 23, wherein the step of cyclizing is carried out in dichloromethane (DCM).

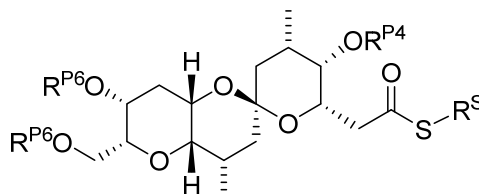
25. The method of any one of claims 22-24, optionally wherein one or more free hydroxyl groups is substituted with an oxygen protecting group; and optionally wherein the one or more oxygen protecting groups are silyl protecting groups.

26. A method of preparing a compound of Formula (E-1):



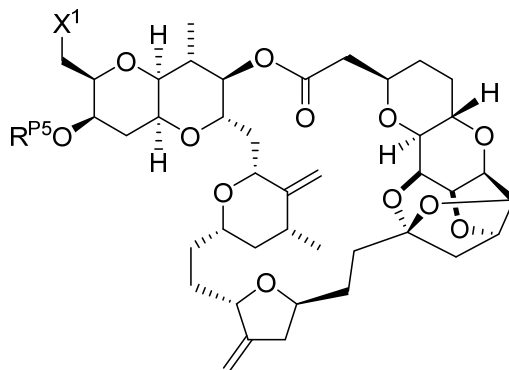
(E-1),

or a salt thereof, the method comprising coupling a compound of the Formula (E-L):



(E-L),

or a salt thereof, with a compound of the Formula (E-R):



(E-R),

or a salt thereof, to yield a compound of the Formula (E-1), or a salt thereof, wherein:

R^S is optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, or optionally substituted heteroaryl;

X¹ is halogen or a leaving group; and

R^{P4}, R^{P5}, and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl.

27. The method of claim 5 or 26, wherein the step of coupling is carried out in the presence of nickel and zirconium.

28. The method of claim 27, wherein the nickel is a nickel complex; optionally wherein the nickel is a nickel(II) or nickel(0) complex; optionally wherein the nickel complex is of the formula: NiX₂•(ligand), wherein X is halogen, and “ligand” is a bidentate ligand; optionally wherein the nickel complex is used after complexation of a nickel source and the “ligand” in solution; optionally wherein the nickel source is NiCl₂, the “ligand” is 4,4'-di-tert-butyl-2,2'-dipyridyl (tbbpy), and the nickel complex is of the formula NiCl₂•(tbbpy); optionally wherein the nickel source is NiBr₂, the “ligand” is 4,4'-di-tert-butyl-2,2'-dipyridyl (tbbpy), and the nickel complex is of the formula NiBr₂•(tbbpy); optionally

wherein the nickel is present in a catalytic amount; optionally
wherein the nickel is present in from 1-50 mol%; and optionally
wherein the nickel is present in from 1-10 mol%.

29. The method of claim 27 or 28, wherein the zirconium is a zirconium complex; optionally
wherein the zirconium complex is of the formula: (ligand)_nZrX₂, wherein n 0, 1, 2, 3, or 4, and X
is halogen; optionally
wherein the zirconium complex is Cp₂ZrCl₂; optionally
wherein the zirconium is present in stoichiometric or excess amount; and optionally
wherein the zirconium is present in 1 to 4 equivalents.

30. The method of any one of claims 5 and 26-29, wherein the step of coupling is carried out
in the presence of zinc or manganese; optionally
wherein the reaction is carried out in the presence of manganese metal; and optionally
wherein the manganese is present in an excess amount.

31. The method of any one of claims 5 and 26-30, wherein the step of coupling is carried out
in the presence of triethylsilyl chloride (TESCl).

32. The method of any one of claims 5 and 26-31, wherein the step of coupling is carried out
in the presence of a base or proton scavenger.

33. The method of any one of claims 5 and 26-32, wherein the step of coupling is carried out
in the presence of NiBr₂•(tbbpy), Cp₂ZrCl₂, and manganese metal.

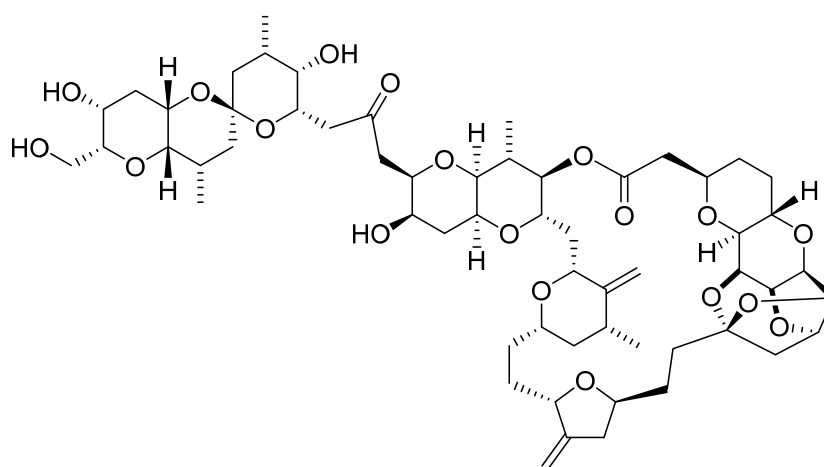
34. The method of claim 33, wherein the step of coupling is carried out by mixing a
compound of Formula (**L-2-6**), or a salt thereof, with a compound of Formula (**R-2-I**), or a salt
thereof, Cp₂ZrCl₂, and manganese metal in a solution; followed by addition of NiBr₂•(tbbpy).

35. The method of claim 33, wherein the step of coupling is carried out by mixing a compound of Formula (E-L), or a salt thereof, with a compound of Formula (E-R), or a salt thereof, Cp_2ZrCl_2 , and manganese metal in a solution; followed by addition of $\text{NiBr}_2 \cdot (\text{tbbpy})$.

36. The method of any one of claims 5 and 26-35, wherein the step of coupling is carried out in one or more solvents; and optionally wherein the step of coupling is carried out in 1,3-dimethyl-2-imidazolidinone (DMI).

37. The method of any one of claims 5 and 26-36, wherein the step of coupling is carried out at or around room temperature; or wherein the step of coupling is carried out at above room temperature; optionally wherein the step of coupling is carried out at approximately 30 °C.

38. The method of any one of claims 26-37 further comprising a step of deprotecting a compound of Formula (E-1), or a salt thereof, to yield Compound (C):



(C),

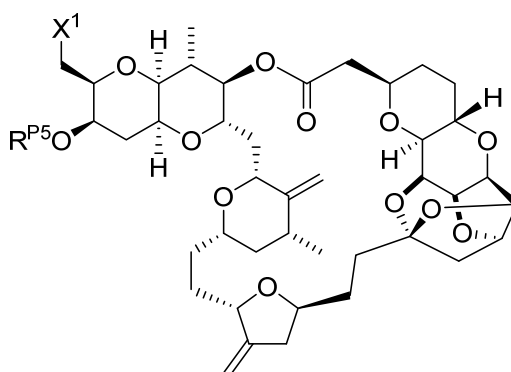
or a salt thereof.

39. The method of claim 6 or 38, wherein $\text{R}^{\text{P}6}$, $\text{R}^{\text{P}5}$, and $\text{R}^{\text{P}4}$ are silyl protecting groups; and the step of deprotecting is carried out in the presence of a fluoride source; optionally wherein the fluoride source is tetrabutylammonium fluoride (TBAF).

40. The method of any one of claims 6, 38, and 39, wherein the step of deprotecting is carried out in the presence of imidazole hydrochloride.

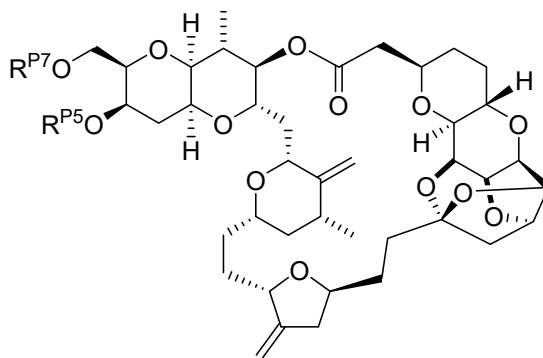
41. The method of any one of claims 6 and 38-40, wherein the step of deprotecting is carried out in THF.

42. The method of any one of claims 26-37, further comprising preparing a compound of Formula (E-R):



(E-R),

or a salt thereof, by reacting a compound of Formula (E-R-1):



(E-R-1),

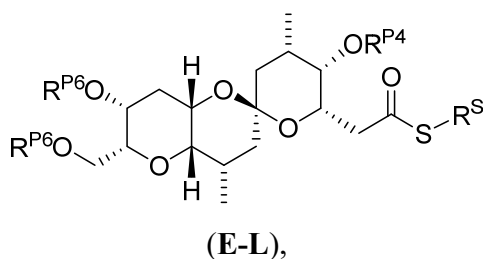
or a salt thereof, in the presence of a nucleophile, thereby substituting the group $-OR^{P7}$ with the group $-X^1$; wherein

R^{P7} is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, optionally substituted acyl, or an oxygen protecting group.

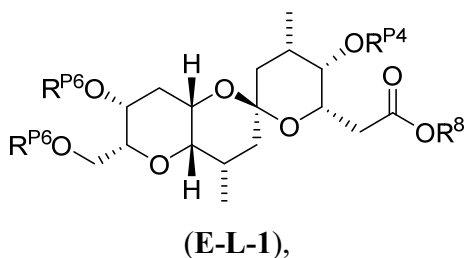
43. The method of claim 7 or 42, wherein X^1 is $-I$; and the reaction is carried out in the presence of an iodide salt; optionally wherein the iodide salt is sodium iodide (NaI).

44. The method of any one of claims 7, 42, and 43, wherein the reaction is carried out in DMI.

45. The method of any one of claims 26-37, further comprising preparing a compound of Formula (E-L):



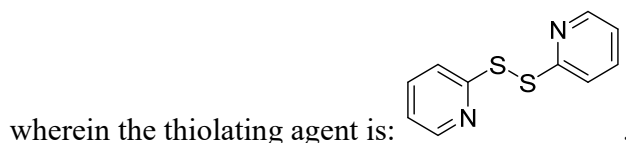
or a salt thereof, by thiolating a compound of Formula (E-L-1):



or a salt thereof, in the presence of a thiolating agent; wherein:

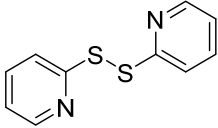
R^8 is hydrogen, optionally substituted alkyl, optionally substituted carbocyclyl, optionally substituted aryl, optionally substituted heterocyclyl, optionally substituted heteroaryl, optionally substituted acyl, or an oxygen protecting group.

46. The method of claim 8 or 45, wherein the thiolating agent is of the formula $(R^S S)_2$; optionally wherein the thiolating agent is of the formula (pyridine-S) $_2$; and optionally



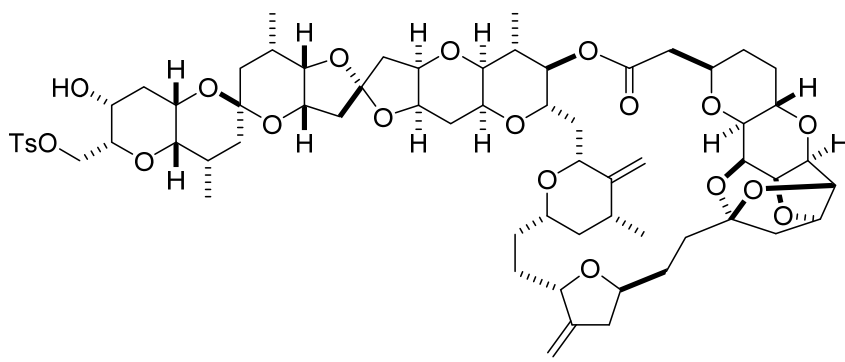
47. The method of any one of claims 8, 45, and 46, wherein the step of thiolating is carried out in the presence of a phosphine reagent; optionally wherein the phosphine reagent is triphenylphosphine (Ph_3P).

48. The method of any one of claims 8 and 45-47, wherein the step of thiolating is carried out

in the presence of the thiolating agent:  and PPh_3 .

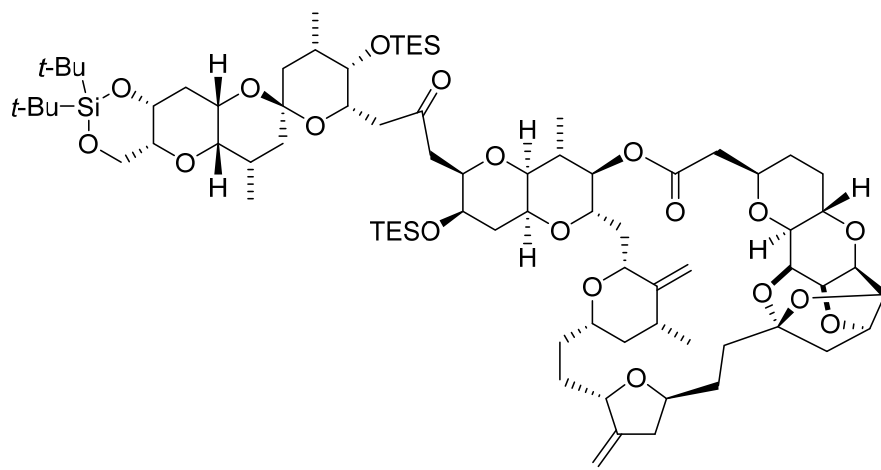
49. The method of any one of claims 8 and 45-48, wherein the step of thiolating is carried out in the presence of acetonitrile (MeCN).

50. The method of any one of claims 13-21, wherein the compound of Formula (A) is the following:



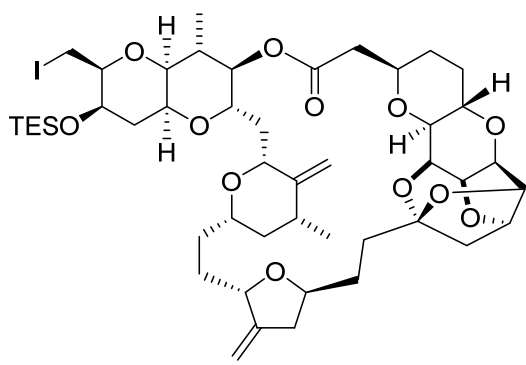
or a salt thereof.

51. The method of any one of claims 26-41, wherein the compound of Formula (E-1) is the following:



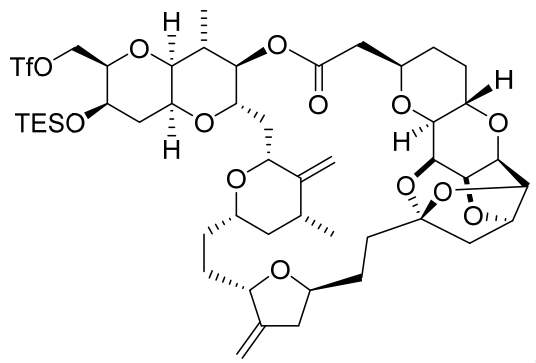
or a salt thereof.

52. The method of any one of claims 26-37, 42-44, and 51, wherein the compound of the Formula (E-R) is the following:



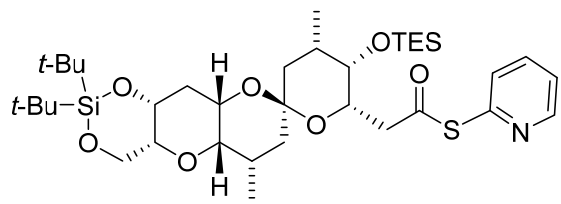
or a salt thereof.

53. The method of any one of claims 42-44 and 52, wherein the compound of Formula (E-R-1) is the following:



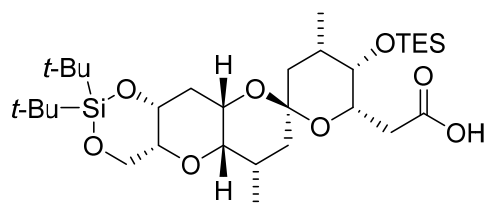
or a salt thereof.

54. The method of any one of claims 26-37, 45-49, 51, and 52, wherein the compound of Formula (E-L) is the following:



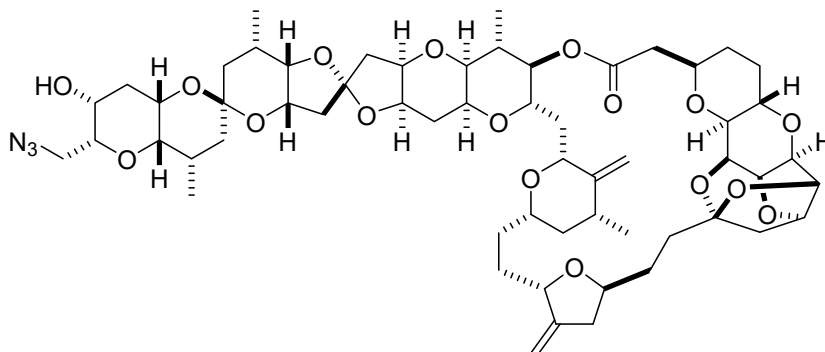
or a salt thereof.

55. The method of any one of claims 45-49 and 54, wherein the compound of Formula (E-L-1) is the following:



or a salt thereof.

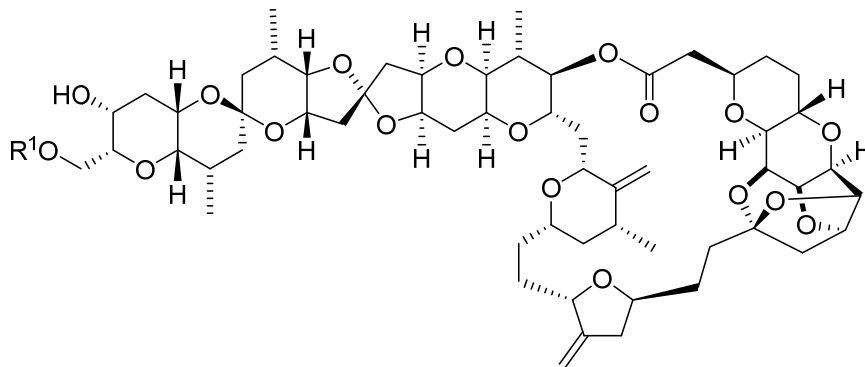
56. A compound of the formula:



(B),

or a salt thereof.

57. A compound of Formula (A):

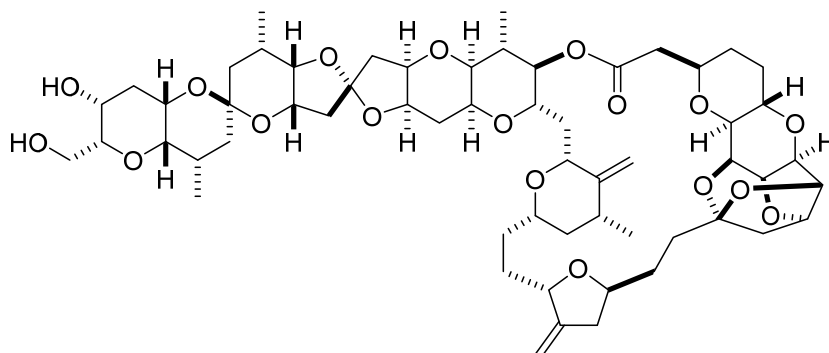


(A),

or a salt thereof, wherein:

R¹ is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl.

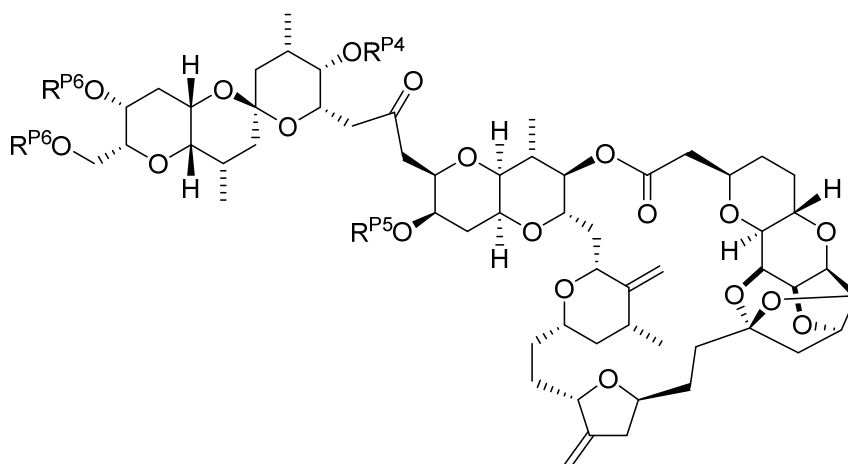
58. A compound of the formula:



(2),

or a salt thereof.

59. A compound of Formula (E-1):



(E-1),

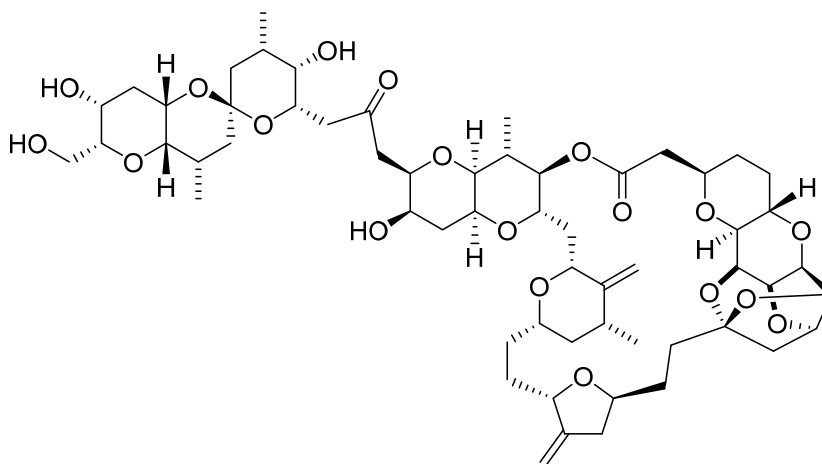
or a salt thereof, wherein:

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl; and

wherein each instance of an oxygen protecting group is independently selected from t-butyldiphenylsilyl (TBDPS), t-butyldimethylsilyl (TBDMS, TBS), triisopropylsilyl (TIPS), triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-

propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranlyl (THF), p-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), p-methoxybenzyl (PMB, MPM), t-butyl, benzyl (Bn), allyl, and pivaloyl (Piv).

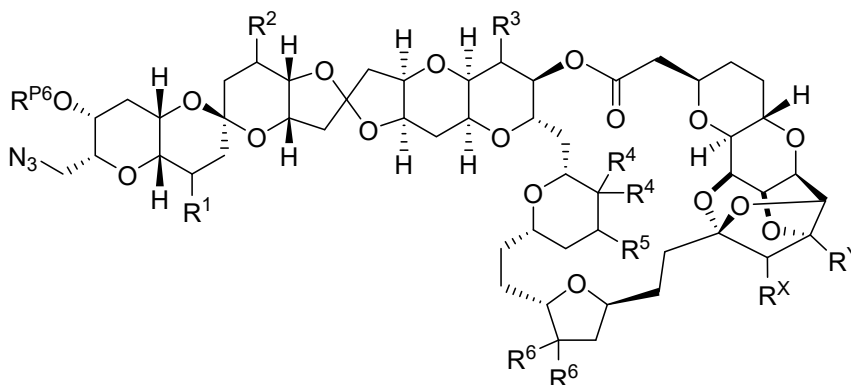
60. The compound of claim 59, wherein the compound is of the formula:



(C),

or a salt thereof.

61. A compound of Formula (H3-N3):

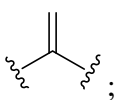


(H3-N3),

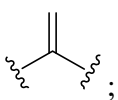
or a salt thereof, wherein:

R¹, R², R³, and R⁵ are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^4 groups are taken together to form:  ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^6 groups are taken together to form:  ;

R^{P6} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

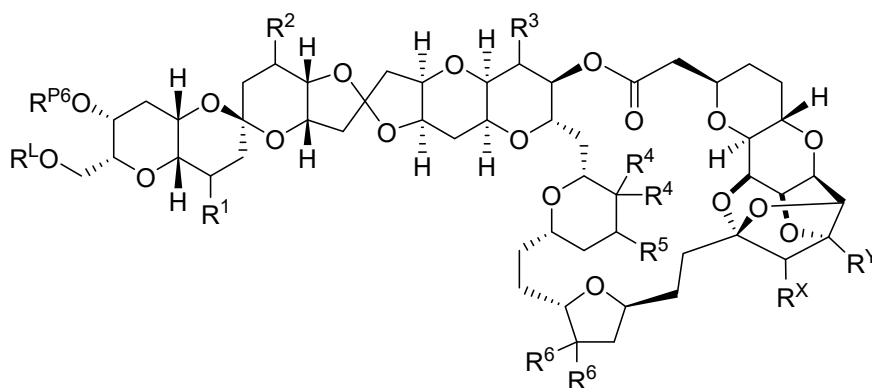
R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl; and

wherein each instance of an oxygen protecting group is independently selected from t-butyldiphenylsilyl (TBDPS), t-butyldimethylsilyl (TBDMS, TBS), triisopropylsilyl (TIPS), triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranyl (THF), p-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), p-methoxybenzyl (PMB, MPM), t-butyl, benzyl (Bn), allyl, and pivaloyl (Piv).

62. A compound of Formula (H3-L):



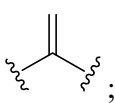
(H3-L),

or a salt thereof, wherein:

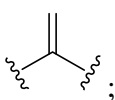
R^L is optionally substituted sulfonyl, optionally substituted sulfinyl, optionally substituted phosphoryl, or optionally substituted acyl;

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^6 groups are taken together to form: ;

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

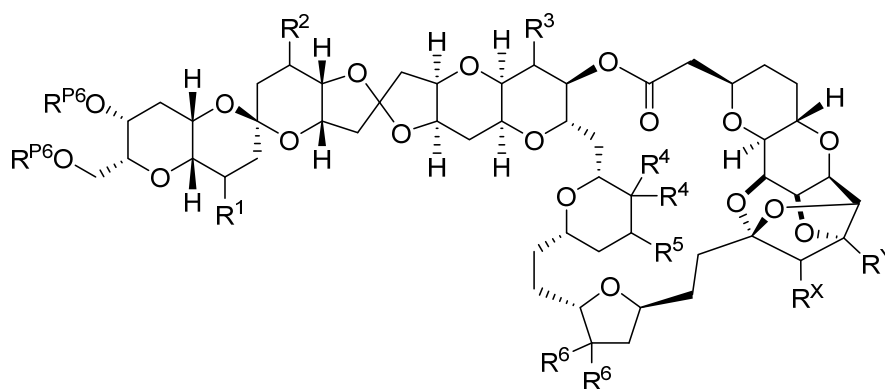
R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl; and

wherein each instance of an oxygen protecting group is independently selected from t-butyldiphenylsilyl (TBDPS), t-butyldimethylsilyl (TBDMS, TBS), triisopropylsilyl (TIPS),

triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranyl (THF), p-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), p-methoxybenzyl (PMB, MPM), t-butyl, benzyl (Bn), allyl, and pivaloyl (Piv).

63. A compound of Formula (H3-2-I):

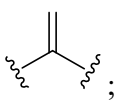


(H3-2-I),

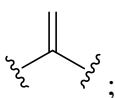
or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^4 groups are taken together to form: ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^6 groups are taken together to form: ;

each instance of R^{P6} is independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

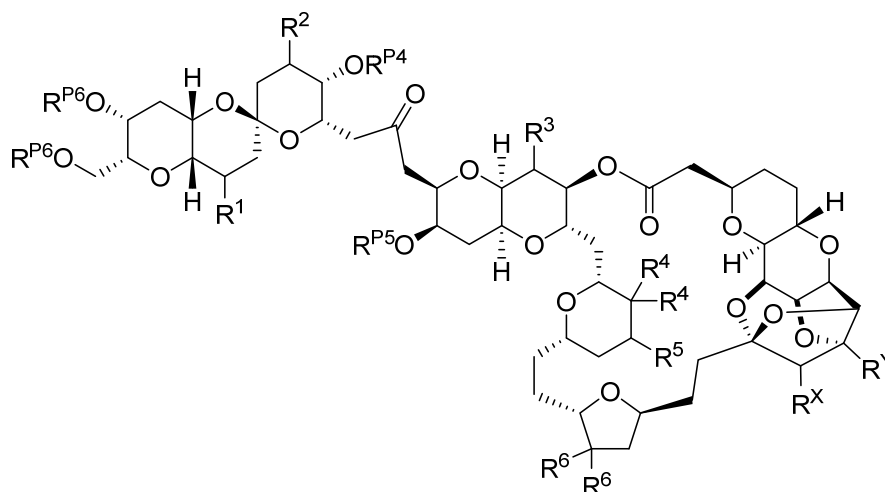
R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl; and

wherein each instance of an oxygen protecting group is independently selected from t-butylphenylsilyl (TBDPS), t-butyltrimethylsilyl (TBDMS, TBS), triisopropylsilyl (TIPS), triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranyl (THF), p-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), p-methoxybenzyl (PMB, MPM), t-butyl, benzyl (Bn), allyl, and pivaloyl (Piv).

64. A compound of Formula (H3-2-II):

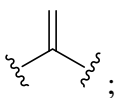


(H3-2-II),

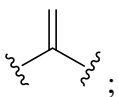
or a salt thereof, wherein:

R^1 , R^2 , R^3 , and R^5 are each independently hydrogen, halogen, or optionally substituted alkyl;

each instance of R^4 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^4 groups are taken together to form:  ;

each instance of R^6 is independently hydrogen, halogen, or optionally substituted alkyl,

or two R^6 groups are taken together to form:  ;

R^{P4} , R^{P5} , and R^{P6} are each independently hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; optionally wherein two R^{P6} are joined with the intervening atoms to form optionally substituted heterocyclyl;

R^X is hydrogen or $-OR^{Xa}$, wherein R^{Xa} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group; and

R^Y is hydrogen or $-OR^{Ya}$, wherein R^{Ya} is hydrogen, optionally substituted alkyl, optionally substituted acyl, or an oxygen protecting group;

optionally wherein R^{Xa} and R^{Ya} are joined together with their intervening atoms to form optionally substituted heterocyclyl; and

wherein each instance of an oxygen protecting group is independently selected from t-butyl diphenylsilyl (TBDPS), t-butyl dimethylsilyl (TBDMS, TBS), triisopropylsilyl (TIPS), triphenylsilyl (TPS), triethylsilyl (TES), trimethylsilyl (TMS), triisopropylsiloxymethyl (TOM), acetyl (Ac), benzoyl (Bz), allyl carbonate, 2,2,2-trichloroethyl carbonate (Troc), 2-trimethylsilylethyl carbonate, methoxymethyl (MOM), 1-ethoxyethyl (EE), 2-methoxy-2-propyl (MOP), 2,2,2-trichloroethoxyethyl, 2-methoxyethoxymethyl (MEM), 2-trimethylsilylethoxymethyl (SEM), methylthiomethyl (MTM), tetrahydropyranyl (THP), tetrahydrofuranyl (THF), p-methoxyphenyl (PMP), triphenylmethyl (Tr), methoxytrityl (MMT), dimethoxytrityl (DMT), p-methoxybenzyl (PMB, MPM), t-butyl, benzyl (Bn), allyl, and pivaloyl (Piv).