

BICYCLIC SULFONAMIDES

INCORPORATION BY REFERENCE TO ANY PRIORITY APPLICATIONS

[0001] Any and all applications for which a foreign or domestic priority claim is identified, for example, in the Application Data Sheet or Request as filed with the present application, are hereby incorporated by reference under 37 CFR 1.57, and Rules 4.18 and 20.6, including U.S. Provisional Application No. 62/805,725, filed February 14, 2019.

BACKGROUND

Field

[0002] The present application relates to the fields of chemistry, biochemistry and medicine. Disclosed herein are compounds of Formulae (I) and (II), or pharmaceutically acceptable salts of the foregoing, pharmaceutical compositions that include a compound described herein (including pharmaceutically acceptable salts of a compound described herein) and methods of synthesizing the same. Also disclosed herein are methods of treating diseases and/or conditions with a compound of Formulae (I) and/or (II), or a pharmaceutically acceptable salt of any of the foregoing.

Description

[0003] The hepatitis B virus (HBV) is a DNA virus and a member of the *Hepadnaviridae* family. HBV infects more than 300 million worldwide and is a causative agent of liver cancer and liver disease such as chronic hepatitis, cirrhosis, and hepatocellular carcinoma. Although there are approved drugs for treating HBV, by either boosting the immune system or slowing down the replication of the HBV virus, HBV continues to be a problem due to the drawbacks associated with each of the approved drugs.

SUMMARY

[0004] Some embodiments disclosed herein relate to a compound of Formula (I), or a pharmaceutically acceptable salt thereof. Other embodiments disclosed herein relate to a compound of Formula (II), or a pharmaceutically acceptable salt thereof.

[0005] Some embodiments disclosed herein relate to a pharmaceutical composition that can contain an effective amount of a compound of Formula (I), or a pharmaceutically acceptable salt thereof, and/or a compound of Formula (II), or a pharmaceutically acceptable salt thereof.

[0006] Some embodiments described herein relate to a method of treating a HBV and/or HDV infection that can include administering to a subject identified as suffering from the HBV and/or HDV infection an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for the use of treating a HBV and/or HDV infection.

[0007] Some embodiments disclosed herein relate to a method of inhibiting replication of HBV and/or HDV that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for the use of inhibiting the replication HBV and/or HDV.

[0008] These are other embodiments are described in greater detail below

DETAILED DESCRIPTION

[0009] HBV is a partially double-stranded circular DNA of about 3.2 kilobase (kb) pairs, and is classified into eight genotypes, A to H. The HBV replication pathway has been studied in great detail. T.J. Liang, *Heptaology* (2009) 49(5 Suppl):S13-S21. On part of replication includes the formation of the covalently closed circular (cccDNA) form. The presence of the cccDNA gives rise to the risk of viral reemergence throughout the life of the host organism. HBV carriers can transmit the disease for many years. An estimated 300

million people are living with hepatitis B virus infection, and it is estimated that over 750,000 people worldwide die of hepatitis B each year. In addition, immunosuppressed individuals or individuals undergoing chemotherapy are especially at risk for reactivation of a HBV infection. HBV can be acute and/or chronic. Acute HBV infection can be either asymptomatic or present with symptomatic acute hepatitis.

[0010] HBV can be transmitted by blood, semen, and/or another body fluid. This can occur through direct blood-to-blood contact, unprotected sex, sharing of needles, and from an infected mother to her baby during the delivery process. The HBV surface antigen (HBsAg) is most frequently used to screen for the presence of this infection. Currently available medications do not cure a HBV and/or HDV infection. Rather, the medications suppress replication of the virus.

[0011] The hepatitis D virus (HDV) is a DNA virus, also in the Hepadnaviridae family of viruses. HDV can propagate only in the presence of HBV. The routes of transmission of HDV are similar to those for HBV. Transmission of HDV can occur either via simultaneous infection with HBV (coinfection) or in addition to chronic hepatitis B or hepatitis B carrier state (superinfection). Both superinfection and coinfection with HDV results in more severe complications compared to infection with HBV alone. These complications include a greater likelihood of experiencing liver failure in acute infections and a rapid progression to liver cirrhosis, with an increased risk of developing liver cancer in chronic infections. In combination with hepatitis B, hepatitis D has the highest fatality rate of all the hepatitis infections, at 20%. There is currently no cure or vaccine for hepatitis D.

Definitions

[0012] Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of ordinary skill in the art. All patents, applications, published applications and other publications referenced herein are incorporated by reference in their entirety unless stated otherwise. In the event that there are a plurality of definitions for a term herein, those in this section prevail unless stated otherwise.

[0013] Whenever a group is described as being “optionally substituted” that group may be unsubstituted or substituted with one or more of the indicated substituents. Likewise, when a group is described as being “unsubstituted or substituted” if substituted,

the substituent(s) may be selected from one or more of the indicated substituents. If no substituents are indicated, it is meant that the indicated “optionally substituted” or “substituted” group may be substituted with one or more group(s) individually and independently selected from deuterium, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl), (heterocyclyl)alkyl, hydroxy, alkoxy, acyl, cyano, halogen, thiocarbonyl, O-carbamyl, N-carbamyl, O-thiocarbamyl, N-thiocarbamyl, C-amido, N-amido, S-sulfonamido, N-sulfonamido, C-carboxy, O-carboxy, isocyanato, thiocyanato, nitro, azido, silyl, sulfenyl, sulfinyl, sulfonyl, haloalkyl, haloalkoxy, trihalomethanesulfonyl, trihalomethanesulfonamido, an amino, a mono-substituted amino group and a di-substituted amino group.

[0014] As used herein, “C_a to C_b” in which “a” and “b” are integers refer to the number of carbon atoms in an alkyl, alkenyl or alkynyl group, or the number of carbon atoms in the ring of a cycloalkyl, cycloalkenyl, aryl, heteroaryl or heterocyclyl group. That is, the alkyl, alkenyl, alkynyl, ring of the cycloalkyl, ring of the cycloalkenyl, ring of the aryl, ring of the heteroaryl or ring of the heterocyclyl can contain from “a” to “b”, inclusive, carbon atoms. Thus, for example, a “C₁ to C₄ alkyl” group refers to all alkyl groups having from 1 to 4 carbons, that is, CH₃-, CH₃CH₂-, CH₃CH₂CH₂-, (CH₃)₂CH-, CH₃CH₂CH₂CH₂-, CH₃CH₂CH(CH₃)- and (CH₃)₃C-. If no “a” and “b” are designated with regard to an alkyl, alkenyl, alkynyl, cycloalkyl cycloalkenyl, aryl, heteroaryl or heterocyclyl group, the broadest range described in these definitions is to be assumed.

[0015] As used herein, “alkyl” refers to a straight or branched hydrocarbon chain that comprises a fully saturated (no double or triple bonds) hydrocarbon group. The alkyl group may have 1 to 20 carbon atoms (whenever it appears herein, a numerical range such as “1 to 20” refers to each integer in the given range; *e.g.*, “1 to 20 carbon atoms” means that the alkyl group may consist of 1 carbon atom, 2 carbon atoms, 3 carbon atoms, *etc.*, up to and including 20 carbon atoms, although the present definition also covers the occurrence of the term “alkyl” where no numerical range is designated). The alkyl group may also be a medium size alkyl having 1 to 10 carbon atoms. The alkyl group could also be a lower alkyl having 1 to 6 carbon atoms. The alkyl group of the compounds may be designated as “C₁-C₄ alkyl” or similar designations. By way of example only, “C₁-C₄ alkyl” indicates that there are one to four carbon atoms in the alkyl chain, *i.e.*, the alkyl chain is selected from methyl,

ethyl, propyl, iso-propyl, n-butyl, iso-butyl, sec-butyl and t-butyl. Typical alkyl groups include, but are in no way limited to, methyl, ethyl, propyl, isopropyl, butyl, isobutyl, tertiary butyl, pentyl and hexyl. The alkyl group may be substituted or unsubstituted.

[0016] As used herein, “alkenyl” refers to an alkyl group that contains in the straight or branched hydrocarbon chain one or more double bonds. The length of an alkenyl can vary. For example, the alkenyl can be a C₂₋₄ alkenyl, C₂₋₆ alkenyl or C₂₋₈ alkenyl. Examples of alkenyl groups include allenyl, vinylmethyl and ethenyl. An alkenyl group may be unsubstituted or substituted.

[0017] As used herein, “alkynyl” refers to an alkyl group that contains in the straight or branched hydrocarbon chain one or more triple bonds. The length of an alkynyl can vary. For example, the alkynyl can be a C₂₋₄ alkynyl, C₂₋₆ alkynyl or C₂₋₈ alkynyl. Examples of alkynyls include ethynyl and propynyl. An alkynyl group may be unsubstituted or substituted.

[0018] As used herein, “cycloalkyl” refers to a completely saturated (no double or triple bonds) mono- or multi- cyclic hydrocarbon ring system. When composed of two or more rings, the rings may be joined together in a fused fashion. Cycloalkyl groups can contain 3 to 10 atoms in the ring(s), 3 to 8 atoms in the ring(s) or 3 to 6 atoms in the ring(s). A cycloalkyl group may be unsubstituted or substituted. Typical cycloalkyl groups include, but are in no way limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl and cyclooctyl.

[0019] As used herein, “cycloalkenyl” refers to a mono- or multi- cyclic hydrocarbon ring system that contains one or more double bonds in at least one ring; although, if there is more than one, the double bonds cannot form a fully delocalized pi-electron system throughout all the rings (otherwise the group would be “aryl,” as defined herein). When composed of two or more rings, the rings may be connected together in a fused fashion. A cycloalkenyl can contain 3 to 10 atoms in the ring(s) or 3 to 8 atoms in the ring(s). A cycloalkenyl group may be unsubstituted or substituted.

[0020] As used herein, “aryl” refers to a carbocyclic (all carbon) monocyclic or polycyclic aromatic ring system (including fused ring systems where two carbocyclic rings share a chemical bond) that has a fully delocalized pi-electron system throughout all the rings. The number of carbon atoms in an aryl group can vary. For example, the aryl group

can be a C₆-C₁₄ aryl group, a C₆-C₁₀ aryl group, or a C₆ aryl group. Examples of aryl groups include, but are not limited to, benzene, naphthalene and azulene. An aryl group may be substituted or unsubstituted.

[0021] As used herein, “heteroaryl” refers to a monocyclic, bicyclic and tricyclic aromatic ring system (a ring system with fully delocalized pi-electron system) that contain(s) one or more heteroatoms (for example, 1 to 5 heteroatoms), that is, an element other than carbon, including but not limited to, nitrogen, oxygen and sulfur. The number of atoms in the ring(s) of a heteroaryl group can vary. For example, the heteroaryl group can contain 4 to 14 atoms in the ring(s), 5 to 10 atoms in the ring(s) or 5 to 6 atoms in the ring(s). Furthermore, the term “heteroaryl” includes fused ring systems where two rings, such as at least one aryl ring and at least one heteroaryl ring, or at least two heteroaryl rings, share at least one chemical bond. Examples of heteroaryl rings include, but are not limited to, furan, furazan, thiophene, benzothiophene, phthalazine, pyrrole, oxazole, benzoxazole, 1,2,3-oxadiazole, 1,2,4-oxadiazole, thiazole, 1,2,3-thiadiazole, 1,2,4-thiadiazole, benzothiazole, imidazole, benzimidazole, indole, indazole, pyrazole, benzopyrazole, isoxazole, benzoisoxazole, isothiazole, triazole, benzotriazole, thiadiazole, tetrazole, pyridine, pyridazine, pyrimidine, pyrazine, purine, pteridine, quinoline, isoquinoline, quinazoline, quinoxaline, cinnoline and triazine. A heteroaryl group may be substituted or unsubstituted.

[0022] As used herein, “heterocyclyl” refers to a monocyclic, bicyclic and tricyclic ring system wherein carbon atoms together with from 1 to 5 heteroatoms constitute said ring system. A heterocycle may optionally contain one or more unsaturated bonds situated in such a way, however, that a fully delocalized pi-electron system does not occur throughout all the rings. The number of atoms in the ring(s) of a heterocyclyl group can vary. For example, the heterocyclyl group can contain 4 to 14 atoms in the ring(s), 5 to 10 atoms in the ring(s) or 5 to 6 atoms in the ring(s). The heteroatom(s) is an element other than carbon including, but not limited to, oxygen, sulfur and nitrogen. A heterocycle may further contain one or more carbonyl or thiocarbonyl functionalities, so as to make the definition include oxo-systems and thio-systems such as lactams, lactones, cyclic imides, cyclic thioimides and cyclic carbamates. When composed of two or more rings, the rings may be joined together in a fused fashion. Additionally, any nitrogens in a heterocyclyl may be quaternized. Heterocyclyl groups may be unsubstituted or substituted. Examples of such

“heterocyclyl groups include but are not limited to, 1,3-dioxin, 1,3-dioxane, 1,4-dioxane, 1,2-dioxolane, 1,3-dioxolane, 1,4-dioxolane, 1,3-oxathiane, 1,4-oxathiin, 1,3-oxathiolane, 1,3-dithiole, 1,3-dithiolane, 1,4-oxathiane, tetrahydro-1,4-thiazine, 2H-1,2-oxazine, maleimide, succinimide, barbituric acid, thiobarbituric acid, dioxopiperazine, hydantoin, dihydrouracil, trioxane, hexahydro-1,3,5-triazine, imidazoline, imidazolidine, isoxazoline, isoxazolidine, oxazoline, oxazolidine, oxazolidinone, thiazoline, thiazolidine, morpholine, oxirane, piperidine *N*-Oxide, piperidine, piperazine, pyrrolidine, pyrrolidone, pyrroldione, 4-piperidone, pyrazoline, pyrazolidine, 2-oxopyrrolidine, tetrahydropyran, 4H-pyran, tetrahydrothiopyran, thiomorpholine, thiomorpholine sulfoxide, thiomorpholine sulfone and their benzo-fused analogs (e.g., benzimidazolidinone, tetrahydroquinoline and 3,4-methylenedioxyphenyl).

[0023] As used herein, “aryl(alkyl)” refer to an aryl group connected, as a substituent, via a lower alkylene group. The lower alkylene and aryl group of an aryl(alkyl) may be substituted or unsubstituted. Examples include but are not limited to benzyl, 2-phenyl(alkyl), 3-phenyl(alkyl), and naphthyl(alkyl).

[0024] As used herein, “heteroaryl(alkyl)” refer to a heteroaryl group connected, as a substituent, via a lower alkylene group. The lower alkylene and heteroaryl group of heteroaryl(alkyl) may be substituted or unsubstituted. Examples include but are not limited to 2-thienyl(alkyl), 3-thienyl(alkyl), furyl(alkyl), thienyl(alkyl), pyrrolyl(alkyl), pyridyl(alkyl), isoxazolyl(alkyl), imidazolyl(alkyl), and their benzo-fused analogs.

[0025] A “(heterocyclyl)alkyl” refer to a heterocyclic group connected, as a substituent, via a lower alkylene group. The lower alkylene and heterocyclyl of a heterocyclyl(alkyl) may be substituted or unsubstituted. Examples include but are not limited tetrahydro-2H-pyran-4-yl(methyl), piperidin-4-yl(ethyl), piperidin-4-yl(propyl), tetrahydro-2H-thiopyran-4-yl(methyl) and 1,3-thiazinan-4-yl(methyl).

[0026] “Lower alkylene groups” are straight-chained -CH₂- tethering groups, forming bonds to connect molecular fragments via their terminal carbon atoms. Examples include but are not limited to methylene (-CH₂-), ethylene (-CH₂CH₂-), propylene (-CH₂CH₂CH₂-) and butylene (-CH₂CH₂CH₂CH₂-). A lower alkylene group can be substituted by replacing one or more hydrogen of the lower alkylene group with a substituent(s) listed under the definition of “substituted.”

[0027] As used herein, “alkoxy” refers to the formula –OR wherein R is an alkyl, an alkenyl, an alkynyl, a cycloalkyl, a cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl) is defined herein. A non-limiting list of alkoxys are methoxy, ethoxy, n-propoxy, 1-methylethoxy (isopropoxy), n-butoxy, isobutoxy, sec-butoxy, tert-butoxy, phenoxy and benzoxy. An alkoxy may be substituted or unsubstituted.

[0028] As used herein, “acyl” refers to a hydrogen an alkyl, an alkenyl, an alkynyl, a cycloalkyl, a cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl) connected, as substituents, via a carbonyl group. Examples include formyl, acetyl, propanoyl, benzoyl, and acryl. An acyl may be substituted or unsubstituted.

[0029] As used herein, “hydroxyalkyl” refers to an alkyl group in which one or more of the hydrogen atoms are replaced by a hydroxy group. Exemplary hydroxyalkyl groups include but are not limited to, 2-hydroxyethyl, 3-hydroxypropyl, 2-hydroxypropyl and 2,2-dihydroxyethyl. A hydroxyalkyl may be substituted or unsubstituted.

[0030] As used herein, “haloalkyl” refers to an alkyl group in which one or more of the hydrogen atoms are replaced by a halogen (e.g., mono-haloalkyl, di-haloalkyl and tri-haloalkyl). Such groups include but are not limited to, chloromethyl, fluoromethyl, difluoromethyl, trifluoromethyl, 1-chloro-2-fluoromethyl and 2-fluoroisobutyl. A haloalkyl may be substituted or unsubstituted.

[0031] As used herein, “haloalkoxy” refers to a O-alkyl group in which one or more of the hydrogen atoms are replaced by a halogen (e.g., mono-haloalkoxy, di-haloalkoxy and tri- haloalkoxy). Such groups include but are not limited to, chloromethoxy, fluoromethoxy, difluoromethoxy, trifluoromethoxy, 1-chloro-2-fluoromethoxy and 2-fluoroisobutoxy. A haloalkoxy may be substituted or unsubstituted.

[0032] A “sulfenyl” group refers to an “-SR” group in which R can be hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). A sulfenyl may be substituted or unsubstituted.

[0033] A “sulfinyl” group refers to an “-S(=O)-R” group in which R can be the same as defined with respect to sulfenyl. A sulfinyl may be substituted or unsubstituted.

[0034] A “sulfonyl” group refers to an “SO₂R” group in which R can be the same as defined with respect to sulfenyl. A sulfonyl may be substituted or unsubstituted.

[0035] An “O-carboxy” group refers to a “RC(=O)O-” group in which R can be hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl), as defined herein. An O-carboxy may be substituted or unsubstituted.

[0036] The terms “ester” and “C-carboxy” refer to a “-C(=O)OR” group in which R can be the same as defined with respect to O-carboxy. An ester and C-carboxy may be substituted or unsubstituted.

[0037] A “thiocarbonyl” group refers to a “-C(=S)R” group in which R can be the same as defined with respect to O-carboxy. A thiocarbonyl may be substituted or unsubstituted.

[0038] A “trihalomethanesulfonyl” group refers to an “X₃CSO₂-” group wherein each X is a halogen.

[0039] A “trihalomethanesulfonamido” group refers to an “X₃CS(O)₂N(R_A)-” group wherein each X is a halogen, and R_A is hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl).

[0040] The term “amino” as used herein refers to a -NH₂ group.

[0041] As used herein, the term “hydroxy” refers to a -OH group.

[0042] A “cyano” group refers to a “-CN” group.

[0043] The term “azido” as used herein refers to a -N₃ group.

[0044] An “isocyanato” group refers to a “-NCO” group.

[0045] A “thiocyanato” group refers to a “-CNS” group.

[0046] An “isothiocyanato” group refers to an “-NCS” group.

[0047] A “mercapto” group refers to an “-SH” group.

[0048] A “carbonyl” group refers to a C=O group.

[0049] An “S-sulfonamido” group refers to a “-SO₂N(R_AR_B)” group in which R_A and R_B can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An S-sulfonamido may be substituted or unsubstituted.

[0050] An “N-sulfonamido” group refers to a “ $\text{RSO}_2\text{N}(\text{R}_\text{A})$ ” group in which R and R_A can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An N-sulfonamido may be substituted or unsubstituted.

[0051] An “O-carbamyl” group refers to a “ $-\text{OC}(=\text{O})\text{N}(\text{R}_\text{A}\text{R}_\text{B})$ ” group in which R_A and R_B can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An O-carbamyl may be substituted or unsubstituted.

[0052] An “N-carbamyl” group refers to an “ $\text{ROC}(=\text{O})\text{N}(\text{R}_\text{A})$ ” group in which R and R_A can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An N-carbamyl may be substituted or unsubstituted.

[0053] An “O-thiocarbamyl” group refers to a “ $-\text{OC}(=\text{S})\text{N}(\text{R}_\text{A}\text{R}_\text{B})$ ” group in which R_A and R_B can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An O-thiocarbamyl may be substituted or unsubstituted.

[0054] An “N-thiocarbamyl” group refers to an “ $\text{ROC}(=\text{S})\text{N}(\text{R}_\text{A})$ ” group in which R and R_A can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An N-thiocarbamyl may be substituted or unsubstituted.

[0055] A “C-amido” group refers to a “ $-\text{C}(=\text{O})\text{N}(\text{R}_\text{A}\text{R}_\text{B})$ ” group in which R_A and R_B can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). A C-amido may be substituted or unsubstituted.

[0056] An “N-amido” group refers to a “ $\text{RC}(=\text{O})\text{N}(\text{R}_\text{A})$ ” group in which R and R_A can be independently hydrogen, alkyl, alkenyl, alkynyl, cycloalkyl, cycloalkenyl, aryl, heteroaryl, heterocyclyl, aryl(alkyl), heteroaryl(alkyl) or heterocyclyl(alkyl). An N-amido may be substituted or unsubstituted.

[0057] The term “halogen atom” or “halogen” as used herein, means any one of the radio-stable atoms of column 7 of the Periodic Table of the Elements, such as, fluorine, chlorine, bromine and iodine.

[0058] The term “ α -amino acid” is used as understood by those skilled in the art. Examples of α -amino acids include, but are not limited to, alanine, asparagine, aspartate, cysteine, glutamate, glutamine, glycine, proline, serine, tyrosine, arginine, histidine, isoleucine, leucine, lysine, methionine, phenylalanine, threonine, tryptophan and valine.

[0059] The term “-O-linked α -amino acid” refers to an α -amino acid that is attached via the hydroxy from its main-chain carboxylic acid group. When the α -amino acid is attached in an -O-linked α -amino acid, the hydrogen that is part of the hydroxy from its main-chain carboxylic acid group is not present and the α -amino acid is attached via the oxygen. An -O-linked α -amino acid can be substituted or unsubstituted.

[0060] Where the numbers of substituents is not specified (e.g. haloalkyl), there may be one or more substituents present. For example, “haloalkyl” may include one or more of the same or different halogens. As another example, “C₁-C₃ alkoxyphenyl” may include one or more of the same or different alkoxy groups containing one, two or three atoms.

[0061] As used herein, the abbreviations for any protective groups, amino acids and other compounds, are, unless indicated otherwise, in accord with their common usage, recognized abbreviations, or the IUPAC-IUB Commission on Biochemical Nomenclature (See, Biochem. 11:942-944 (1972)).

[0062] The term “pharmaceutically acceptable salt” refers to a salt of a compound that does not cause significant irritation to an organism to which it is administered and does not abrogate the biological activity and properties of the compound. In some embodiments, the salt is an acid addition salt of the compound. Pharmaceutical salts can be obtained by reacting a compound with inorganic acids such as hydrohalic acid (e.g., hydrochloric acid or hydrobromic acid), sulfuric acid, nitric acid and phosphoric acid. Pharmaceutical salts can also be obtained by reacting a compound with an organic acid such as aliphatic or aromatic carboxylic or sulfonic acids, for example formic, acetic, succinic, lactic, malic, tartaric, citric, ascorbic, nicotinic, methanesulfonic, ethanesulfonic, p-toluenesulfonic, salicylic or naphthalenesulfonic acid. Pharmaceutical salts can also be obtained by reacting a compound with a base to form a salt such as an ammonium salt, an alkali metal salt, such as a sodium or a potassium salt, an alkaline earth metal salt, such as a calcium or a magnesium salt, a salt of organic bases such as dicyclohexylamine, N-methyl-D-glucamine,

tris(hydroxymethyl)methylamine, C₁-C₇ alkylamine, cyclohexylamine, triethanolamine, ethylenediamine, and salts with amino acids such as arginine and lysine.

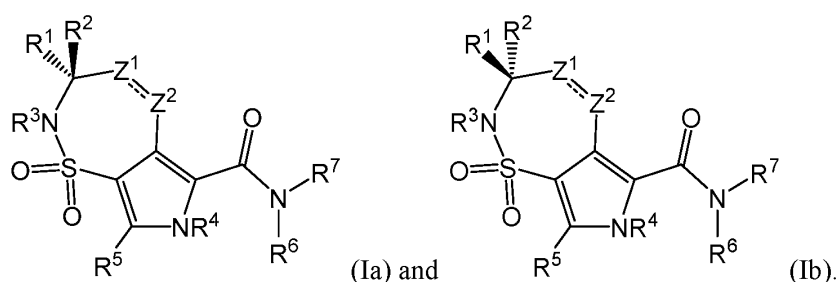
[0063] Terms and phrases used in this application, and variations thereof, especially in the appended claims, unless otherwise expressly stated, should be construed as open ended as opposed to limiting. As examples of the foregoing, the term 'including' should be read to mean 'including, without limitation,' 'including but not limited to,' or the like; the term 'comprising' as used herein is synonymous with 'including,' 'containing,' or 'characterized by,' and is inclusive or open-ended and does not exclude additional, unrecited elements or method steps; the term 'having' should be interpreted as 'having at least;' the term 'includes' should be interpreted as 'includes but is not limited to;' the term 'example' is used to provide exemplary instances of the item in discussion, not an exhaustive or limiting list thereof. In addition, the term "comprising" is to be interpreted synonymously with the phrases "having at least" or "including at least". When used in the context of a compound or composition, the term "comprising" means that the compound or composition includes at least the recited features or components but may also include additional features or components.

[0064] With respect to the use of substantially any plural and/or singular terms herein, those having skill in the art can translate from the plural to the singular and/or from the singular to the plural as is appropriate to the context and/or application. The various singular/plural permutations may be expressly set forth herein for sake of clarity. The indefinite article "a" or "an" does not exclude a plurality.

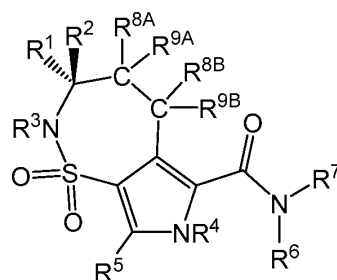
[0065] It is understood that, in any compound described herein having one or more chiral centers, if an absolute stereochemistry is not expressly indicated, then each center may independently be of (R)-configuration or (S)-configuration or a mixture thereof. Thus, the compounds provided herein may be enantiomerically pure, enantiomerically enriched, racemic mixture, diastereomerically pure, diastereomerically enriched, or a stereoisomeric mixture. In addition, it is understood that, in any compound described herein having one or more double bond(s) generating geometrical isomers that can be defined as E or Z, each double bond may independently be E or Z a mixture thereof. Likewise, it is understood that, in any compound described, all tautomeric forms are also intended to be included.

cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A}; R² can be hydrogen, deuterium or a substituted or an unsubstituted C₁₋₄ alkyl, wherein the substituted C₁₋₄ alkyl is substituted with one or more substituents selected from halogen, hydroxy and R^{11B}; R³ can be hydrogen, deuterium or an unsubstituted C₁₋₄ alkyl; R⁴ can be hydrogen, deuterium or an unsubstituted C₁₋₄ alkyl; R⁵ can be hydrogen, deuterium, halogen, an unsubstituted C₁₋₄ alkyl, cyano, an unsubstituted C₁₋₄ haloalkyl or an unsubstituted C₃₋₈ monocyclic cycloalkyl; R⁶ can be a substituted phenyl or a substituted pyridyl, wherein the substituted phenyl and the substituted pyridyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C₁₋₄ haloalkyl and an unsubstituted C₁₋₄ alkyl; R⁷ can be hydrogen, deuterium or an unsubstituted C₁₋₄ alkyl; R^{8A}, R^{8B}, R^{9A} and R^{9B} can be independently hydrogen, deuterium, halogen, an unsubstituted C₁₋₄ alkyl or hydroxy; each R¹⁰ can be independently hydrogen, deuterium, halogen or an unsubstituted C₁₋₄ alkyl; and R^{11A} and R^{11B} can be independently an optionally substituted -O-acyl, an unsubstituted O-linked α -amino acid, -O-P(=O)(OH)₂ or -CH₂-P(=O)(OH)₂.

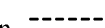
[0070] Compounds of Formula (I), or a pharmaceutically salt thereof, can include one or more chiral centers. As provided herein, if an absolute stereochemistry is not expressly indicated, then each center may independently be of (R)-configuration or (S)-configuration or a mixture thereof. Those skilled in the art recognize that the carbon to which R¹ and R² are attached can be a chiral center. In some embodiments, the stereochemistry of the carbon to which R¹ and R² are attached is (R). In other embodiments, the stereochemistry of the carbon to which R¹ and R² are attached is (S). A compound of Formula (I), or a pharmaceutically acceptable salt, can have a structure selected from:

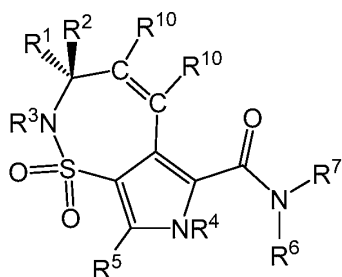


[0071] As shown in Formula (I), ----- can be a single or a double bond. When ----- is a single bond, Z¹ can be CR^{8A}R^{9A} and Z² can be CR^{8B}R^{9B}, such that Formula (I)



has the structure: . In some embodiments, Z^1 can be CH_2 . In some embodiments, Z^2 can be CH_2 . In some embodiments, R^{8A} and/or R^{8B} can be a deuterium. In some embodiments, R^{8A} and/or R^{8B} can be a halogen (such as F or Cl). In some embodiments, R^{8A} and/or R^{8B} can be an unsubstituted C_{1-4} alkyl. In some embodiments, R^{8A} and/or R^{8B} can be hydroxy. In some embodiments, R^{9A} and/or R^{9B} can be a deuterium. In some embodiments, R^{9A} and/or R^{9B} can be a halogen (such as F or Cl). In some embodiments, R^{9A} and/or R^{9B} can be an unsubstituted C_{1-4} alkyl. In some embodiments, R^{9A} and/or R^{9B} can be hydroxy. When, R^{8A} , R^{8B} , R^{9A} , and/or R^{9B} is an unsubstituted C_{1-4} alkyl, the C_{1-4} alkyl can be methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl or tert-butyl.

[0072] When  is a double bond, Z^1 and Z^2 can be each independently CR^{10} , and the compound of Formula (I), or a pharmaceutically acceptable salt, can be



, wherein each R^{10} can be independently hydrogen, deuterium, halogen and an unsubstituted C_{1-4} alkyl. In some embodiments, Z^1 can be CH . In other embodiments, Z^1 can be CD . In some embodiments, Z^2 can be CH . In other embodiments, Z^2 can be CD . The substituent attached to Z^1 and/or Z^2 can also be a halogen, such as F or Cl. In some embodiments, Z^1 can have an unsubstituted C_{1-4} alkyl attached. In some embodiments, Z^2 can have an unsubstituted C_{1-4} alkyl attached. Examples of unsubstituted C_{1-4} alkyls are described herein, including the previous paragraph. When two R^{10} groups are

present, in some embodiments, the R^{10} groups can be the same. In other embodiments, when two R^{10} groups are present, the R^{10} groups can be different.

[0073] Various alkenyls and alkynyls can be attached to the seven-membered ring of Formula (I) are provided herein. The alkenyl can have 2 to 8 carbons, 2 to 5 carbons or 3 to 4 carbons. In some embodiments, R^1 can be an unsubstituted C_{2-8} alkenyl. In other embodiments, R^1 can be an unsubstituted C_{2-8} alkynyl. In still other embodiments, R^1 can be a substituted C_{2-8} alkenyl, wherein the substituted C_{2-8} alkenyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A} . In yet still other embodiments, R^1 can be a substituted C_{3-4} alkenyl, wherein the substituted C_{3-4} alkenyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A} .

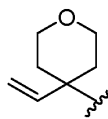
[0074] Examples of substituted alkenyls include a substituted alkenyl substituted with one or more halogens (such as F and/or Cl), a substituted alkenyl substituted with one or more hydroxys, a substituted alkenyl substituted with an unsubstituted monocyclic C_{3-6} cycloalkyl, a substituted alkenyl substituted with a substituted monocyclic C_{3-6} cycloalkyl, a substituted alkenyl substituted with an unsubstituted bicyclic C_{3-8} cycloalkyl selected from an unsubstituted fused bicyclic C_{3-8} cycloalkyl, an unsubstituted bridged bicyclic C_{3-8} cycloalkyl and an unsubstituted spiro bicyclic C_{3-8} cycloalkyl, a substituted alkenyl substituted with a substituted bicyclic C_{3-8} cycloalkyl selected from a substituted fused bicyclic C_{3-8} cycloalkyl, a substituted bridged bicyclic C_{3-8} cycloalkyl and a substituted spiro bicyclic C_{3-8} cycloalkyl, a substituted alkenyl substituted with an unsubstituted monocyclic heterocyclyl, a substituted alkenyl substituted with a substituted monocyclic heterocyclyl and a substituted alkenyl substituted with R^{11A} .

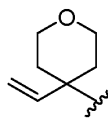
[0075] When R^1 is an alkenyl, the alkenyl can include a single double bond. The position of the double bond can vary. In some embodiments, the double bond can be located between the terminal carbon and a carbon adjacent to the terminal carbon. In some embodiments, the double bond can be located between the carbon adjacent to the seven-

membered ring of Formula (I) and the next carbon away from the seven-membered ring of Formula (I).

[0076] The alkynyl can have 2 to 8 carbons. In some embodiments, the alkynyl for R^1 can be a C_{2-8} alkynyl. In other embodiments, the alkynyl for R^1 can be a C_{2-5} alkynyl. In still embodiments, the alkynyl for R^1 can be a C_{3-5} alkynyl. In some embodiments, R^1 can be a substituted C_{2-8} alkynyl, wherein the substituted C_{2-8} alkynyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A} . In some embodiments, R^1 can be a substituted C_{3-5} alkynyl, wherein the substituted C_{3-5} alkynyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A} .

[0077] The alkynyl described herein for R^1 can be a substituted alkynyl substituted with one or more halogens (for example, F or Cl), a substituted alkynyl substituted with one or more hydroxys, a substituted alkynyl substituted with an unsubstituted monocyclic C_{3-6} cycloalkyl, a substituted alkynyl substituted with a substituted monocyclic C_{3-6} cycloalkyl, a substituted alkynyl substituted with an unsubstituted bicyclic C_{3-8} cycloalkyl selected from an unsubstituted fused bicyclic C_{3-8} cycloalkyl, an unsubstituted bridged bicyclic C_{3-8} cycloalkyl and an unsubstituted spiro bicyclic C_{3-8} cycloalkyl, a substituted alkynyl substituted with a substituted bicyclic C_{3-8} cycloalkyl selected from a substituted fused bicyclic C_{3-8} cycloalkyl, a substituted bridged bicyclic C_{3-8} cycloalkyl and a substituted spiro bicyclic C_{3-8} cycloalkyl, a substituted alkynyl substituted with an unsubstituted monocyclic heterocyclyl, a substituted alkynyl substituted with a substituted monocyclic heterocyclyl and a substituted alkynyl substituted with R^{11A} . Those skilled in the art will appreciate that when R^1 is substituted with an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl or an optionally substituted monocyclic heterocyclyl, the optionally substituted monocyclic C_{3-6} cycloalkyl, the optionally substituted bicyclic C_{3-8} cycloalkyl and the optionally substituted monocyclic heterocyclyl may substitute R^1 by replacing two hydrogens of R^1 . For example, when R^1 is a C_3 -alkenyl substituted with a tetrahydropyran (a monocyclic heterocyclyl) by replacing two



hydrogens of R^1 , R^1 may have the structure , wherein the tetrahydropyran moiety is connected in a spiro-fashion. In some embodiments, R^1 is substituted with an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl or an optionally substituted monocyclic heterocyclyl, wherein the optionally substituted monocyclic C_{3-6} cycloalkyl, the optionally substituted bicyclic C_{3-8} cycloalkyl or the optionally substituted monocyclic heterocyclyl replaces one hydrogen. In other embodiments, R^1 is substituted with an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl or an optionally substituted monocyclic heterocyclyl, wherein the optionally substituted monocyclic C_{3-6} cycloalkyl, the optionally substituted bicyclic C_{3-8} cycloalkyl or the optionally substituted monocyclic heterocyclyl replaces two hydrogen such that the aforementioned moieties are connected in a spiro-fashion.

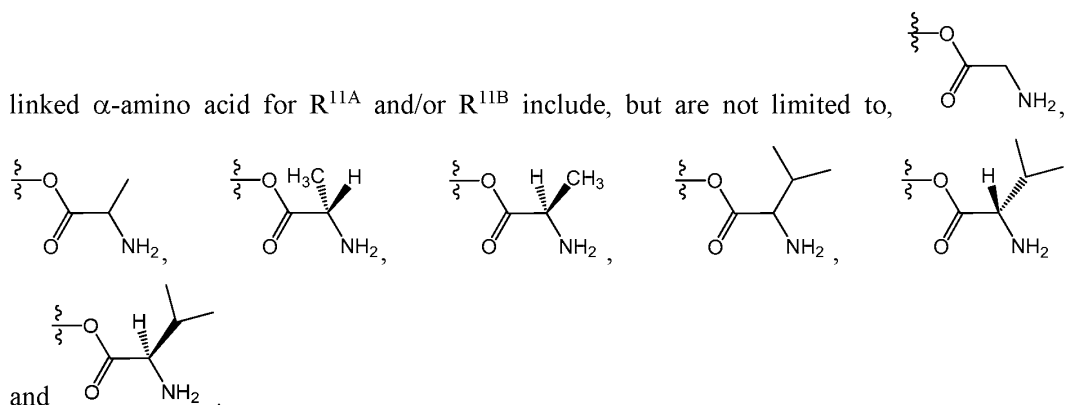
[0078] The alkynyl of R^1 can include a single triple bond. The position of the triple bond can vary. In some embodiments, the triple bond can be located between the terminal carbon and a carbon adjacent to the terminal carbon. In some embodiments, the triple bond can be located between the carbon adjacent to the seven-membered ring of Formula (I) and the next carbon away from the seven-membered ring of Formula (I).

[0079] When R^1 is a substituted alkenyl or a substituted alkynyl as described herein and substituted with R^{11A} , R^{11A} can be an optionally substituted $-O$ -acyl prodrug or an unsubstituted $-O$ -linked α -amino acid prodrug. An example of an optionally substituted $-O$ -acyl is $-O-C(=O)R^{11A1}$, wherein R^{11A1} can be an optionally substituted C_{1-6} alkyl or an optionally substituted C_6 or C_{14} aryl. In some embodiments, R^{11A} can be $-O-C(=O)R^{11A1}$, wherein R^{11A1} can be an unsubstituted C_{1-6} alkyl.

[0080] Alpha-amino acids are known to those skilled in the art, and include those described herein. In some embodiments, R^{11A} can be $-O$ -linked glycine, $-O$ -linked valine, $-O$ -linked leucine or $-O$ -linked isoleucine. As provided herein, R^{11A} can be a phosphate or a phosphonate. In some embodiments, R^{11A} can be $-O-P(=O)(OH)_2$. In other embodiments, R^{11A} can be $-CH_2-P(=O)(OH)_2$.

[0081] For R^2 , in some embodiments, R^2 can be hydrogen. In other embodiments, R^2 can be deuterium. In still other embodiments, R^2 can be an unsubstituted C_{1-4} alkyl, for example, methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl and tert-butyl. In some embodiments, R^2 can be a substituted C_{1-4} alkyl, wherein the substituted C_{1-4} alkyl can be substituted with one or more substituents selected from halogen (for example F or Cl), hydroxy and R^{11B} . Examples of R^{13} begin a substituted C_{1-4} alkyl include $-CF_3$, $-CHF_2$, $-CH_2OH$ and $-CH(OH)CH_3$.

[0082] As with R^{11A} , R^{11B} can be an optionally substituted $-O$ -acyl prodrug or an unsubstituted $-O$ -linked α -amino acid prodrug. The optionally substituted $-O$ -acyl of R^{11B} can be $-O-C(=O)R^{11B1}$, wherein R^{11B1} can be an optionally substituted C_{1-6} alkyl or an optionally substituted C_6 or C_{14} aryl. In some embodiments, R^{11B} can be $-O-C(=O)R^{11B1}$, wherein R^{11B1} can be an unsubstituted C_{1-6} alkyl. In some embodiments, R^{11B} can be $-O$ -linked glycine, $-O$ -linked valine, $-O$ -linked leucine or $-O$ -linked isoleucine. Exemplary $-O$ -



[0083] A phosphate or a phosphonate can be present at R^{11B} . In some embodiments, R^{11B} can be $-O-P(=O)(OH)_2$. In other embodiments, R^{11B} can be $-CH_2-P(=O)(OH)_2$.

[0084] In some embodiments, R^3 can be hydrogen. In other embodiments, R^3 can be deuterium. In still other embodiments, R^3 can be an unsubstituted C_{1-4} alkyl, such as those described herein. In some embodiments, R^3 can be methyl.

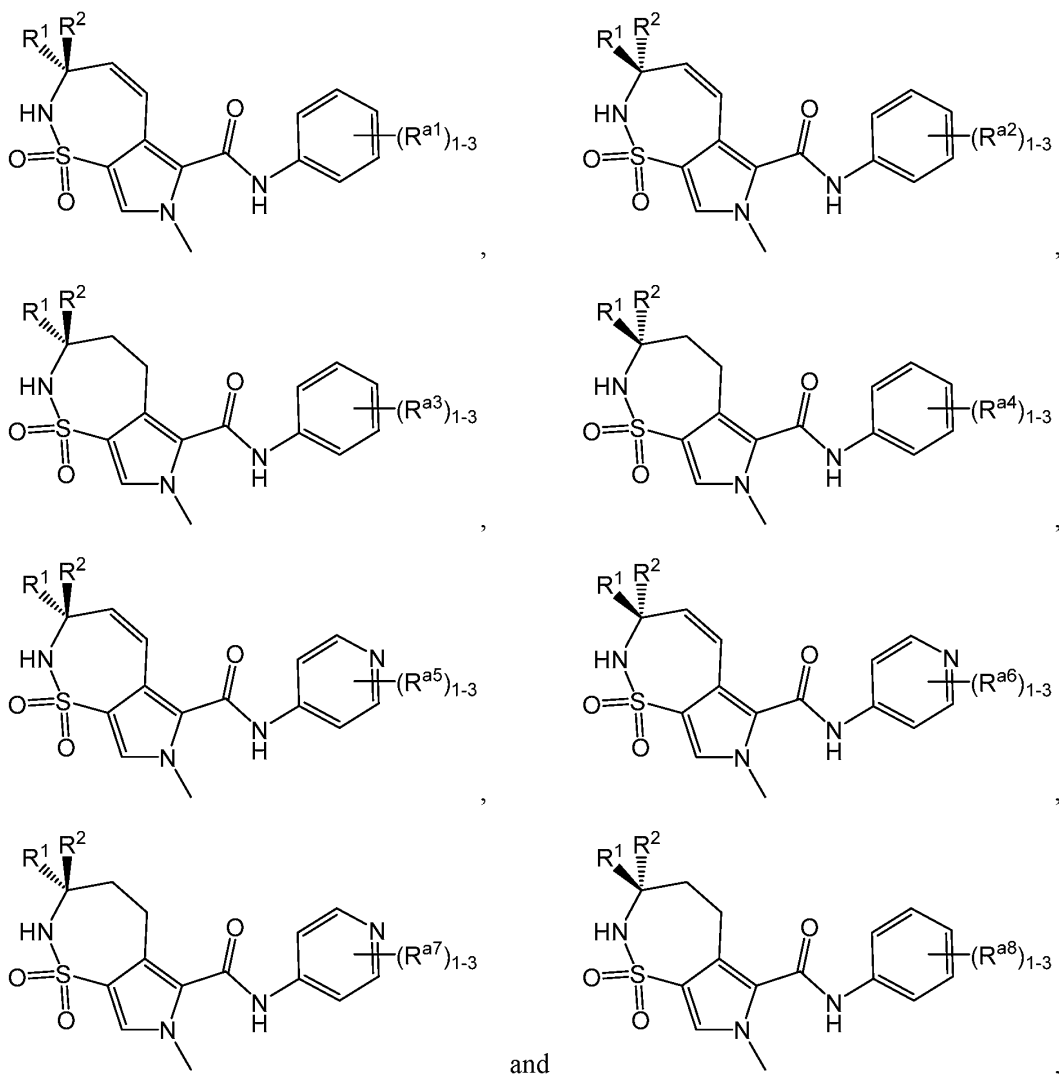
[0085] The 5-membered ring of Formula (I) can be unsubstituted or substituted with an unsubstituted C_{1-4} alkyl, cyano and/or an unsubstituted C_{1-4} haloalkyl. In some embodiments, R^4 can be hydrogen. In other embodiments, R^4 can be deuterium. In still

other embodiments, R^4 can be an unsubstituted C_{1-4} alkyl. In some embodiments, R^5 can be hydrogen. In other embodiments, R^5 can be deuterium. In still other embodiments, R^5 can be halogen (for example, F or Cl). In yet still other embodiments, R^5 can be an unsubstituted C_{1-4} alkyl such as those described herein. In some embodiments, R^5 can be cyano. In other embodiments, R^5 can be an unsubstituted C_{1-4} haloalkyl. An example of a suitable C_{1-4} haloalkyl is CF_3 . In still other embodiments, R^5 can be an unsubstituted C_{3-8} monocyclic cycloalkyl. In some embodiments, R^4 can be methyl; and R^5 can be hydrogen.

[0086] The C-amide of Formula (I) can include a substituted phenyl or a substituted pyridyl as described herein. In some embodiments, R^6 can be a substituted phenyl, wherein the phenyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C_{1-4} haloalkyl and an unsubstituted C_{1-4} alkyl. In other embodiments, R^6 can be a substituted pyridyl, wherein the pyridyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C_{1-4} haloalkyl and an unsubstituted C_{1-4} alkyl. The phenyl and pyridyl of R^6 can be substituted with one or more substituents as described herein. In some embodiments, R^6 can be a mono-substituted phenyl. In other embodiments, R^6 can be a di-substituted phenyl. In some embodiments, the phenyl of R^6 can be substituted at the para-position and/or meta-position. In some embodiments, the phenyl of R^6 can be a 3,4-disubstituted phenyl. In still other embodiments, R^6 can be a mono-substituted pyridyl. In yet still other embodiments, R^6 can be a di-substituted pyridyl. In some embodiments, the pyridyl can be substituted on a carbon adjacent to the nitrogen of the pyridyl. The unsubstituted C_{1-4} alkyl(s) that can be substituted on R^6 can be methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl or tert-butyl. Examples of unsubstituted C_{1-4} haloalkyls are CF_3 , CHF_2 and CH_2F . In some embodiments, R^6 can be substituted with F and/or Cl. In some embodiments, R^6 can be substituted with F, Cl and/or Br. In some embodiments, R^6 can be substituted with CF_3 . In some embodiments, R^6 can be substituted with CH_3 . When R^6 is di-substituted, the two groups can be the same or different.

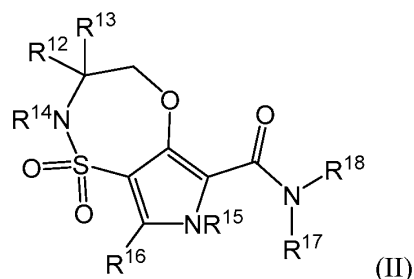
[0087] The other group of the C-amide of Formula (I), R^7 , can be hydrogen, deuterium or an unsubstituted C_{1-4} alkyl. In some embodiments, R^7 can be hydrogen. In other embodiments, R^7 can be deuterium. In still other embodiments, R^7 can be an unsubstituted C_{1-4} alkyl.

[0088] Examples of compounds of Formula (I), including pharmaceutically acceptable salts thereof, include the following:



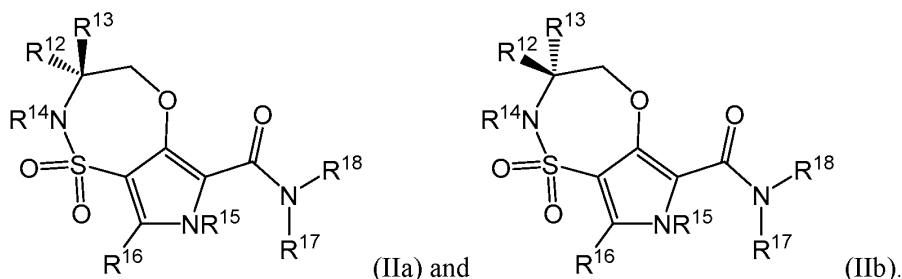
wherein each R^{a1} , each R^{a2} , each R^{a3} , each R^{a4} , each R^{a5} , each R^{a6} , each R^{a7} and each R^{a8} are independently selected from methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, tert-butyl, cyano, CF_3 , CHF_2 , CH_2F , F, Cl and Br.

[0089] Some embodiments disclosed herein relate to a compound of Formula (II), or a pharmaceutically acceptable salt thereof:



wherein: R^{12} can be a substituted or an unsubstituted C_{2-8} alkenyl or a substituted or an unsubstituted C_{2-8} alkynyl, wherein the substituted C_{2-8} alkenyl and the substituted C_{2-8} alkynyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A} ; R^{13} can be hydrogen, deuterium or a substituted or an unsubstituted C_{1-4} alkyl, wherein the substituted C_{1-4} alkyl can be substituted with one or more substituents selected from halogen, hydroxy and R^{19B} ; R^{14} can be hydrogen, deuterium or an unsubstituted C_{1-4} alkyl; R^{15} can be hydrogen, deuterium or an unsubstituted C_{1-4} alkyl; R^{16} can be hydrogen, deuterium, halogen, an unsubstituted C_{1-4} alkyl, cyano, an unsubstituted C_{1-4} haloalkyl or an unsubstituted C_{3-8} monocyclic cycloalkyl; R^{17} can be a substituted phenyl or a substituted pyridyl, wherein the substituted phenyl and the substituted pyridyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C_{1-4} haloalkyl and an unsubstituted C_{1-4} alkyl; R^{18} can be hydrogen, deuterium or an unsubstituted C_{1-4} alkyl; and R^{19A} and R^{19B} can be independently an optionally substituted $-O$ -acyl, an unsubstituted O -linked α -amino acid, $-O-P(=O)(OH)_2$ or $-CH_2-P(=O)(OH)_2$.

[0090] A compound of Formula (II), or a pharmaceutically acceptable salt thereof, can have the following structures:



[0091] The groups present on R¹² can be a C₂₋₈ alkenyl or a C₂₋₈ alkynyl. In some embodiments, R¹² can be a C₂₋₈ alkenyl. In other embodiments, R¹² can be a C₂₋₅ alkenyl. In still other embodiments, R¹² can be a C₃₋₄ alkenyl. In some embodiments, R¹² can be an unsubstituted C₂₋₈ alkenyl. As provided herein, in some embodiments, R¹² can be a substituted C₂₋₈ alkenyl, wherein the substituted C₂₋₈ alkenyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A}. In some embodiments, R¹² can be a substituted C₃₋₄ alkenyl, wherein the substituted C₃₋₄ alkenyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A}.

[0092] Examples of substituted alkenyls for R¹² include a substituted alkenyl substituted with one or more halogens (for example, F and/or Cl), a substituted alkenyl substituted with one or more hydroxys, a substituted alkenyl substituted with an unsubstituted monocyclic C₃₋₆ cycloalkyl, a substituted alkenyl substituted with a substituted monocyclic C₃₋₆ cycloalkyl, a substituted alkenyl substituted with an unsubstituted bicyclic C₃₋₈ cycloalkyl selected from an unsubstituted fused bicyclic C₃₋₈ cycloalkyl, an unsubstituted bridged bicyclic C₃₋₈ cycloalkyl and an unsubstituted spiro bicyclic C₃₋₈ cycloalkyl, a substituted alkenyl substituted with a substituted bicyclic C₃₋₈ cycloalkyl selected from a substituted fused bicyclic C₃₋₈ cycloalkyl, a substituted bridged bicyclic C₃₋₈ cycloalkyl and a substituted spiro bicyclic C₃₋₈ cycloalkyl, a substituted alkenyl substituted

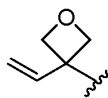
with an unsubstituted monocyclic heterocyclyl, a substituted alkenyl substituted with a substituted monocyclic heterocyclyl and a substituted alkenyl substituted with R^{19A}.

[0093] When R¹² is an alkenyl, the alkenyl can include a single double bond. The position of the double bond can vary. In some embodiments, the double bond can be located between the terminal carbon and a carbon adjacent to the terminal carbon. In some embodiments, the double bond can be located between the carbon adjacent to the seven-membered ring of Formula (I) and the next carbon away from the seven-membered ring of Formula (I).

[0094] In some embodiments R¹² can be an unsubstituted C₂₋₈ alkynyl. In other embodiments, R¹² can be a substituted C₂₋₈ alkynyl, wherein the substituted C₂₋₈ alkynyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A}. The alkynyl can have 2 to 8 carbons, 3 to 6 carbons or 3 to 5 carbons. In some embodiments, R¹² is a substituted C₃₋₅ alkynyl, wherein the substituted C₃₋₅ alkynyl can be substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A}.

[0095] Examples of substituted C₂₋₈ alkynyl for R¹² include a substituted alkynyl substituted with one or more halogens (for example, F or Cl), a substituted alkynyl substituted with one or more hydroxys, a substituted alkynyl substituted with an unsubstituted monocyclic C₃₋₆ cycloalkyl, a substituted alkynyl substituted with a substituted monocyclic C₃₋₆ cycloalkyl, a substituted alkynyl substituted with an unsubstituted bicyclic C₃₋₈ cycloalkyl selected from an unsubstituted fused bicyclic C₃₋₈ cycloalkyl, an unsubstituted bridged bicyclic C₃₋₈ cycloalkyl and an unsubstituted spiro bicyclic C₃₋₈ cycloalkyl, a substituted alkynyl substituted with a substituted bicyclic C₃₋₈ cycloalkyl selected from a substituted fused bicyclic C₃₋₈ cycloalkyl, a substituted bridged bicyclic C₃₋₈ cycloalkyl and a substituted spiro bicyclic C₃₋₈ cycloalkyl, a substituted alkynyl substituted with an unsubstituted monocyclic heterocyclyl, a substituted alkynyl substituted with a substituted monocyclic heterocyclyl and a substituted alkynyl substituted with R^{19A}. When R¹² is substituted with an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally

substituted bicyclic C₃₋₈ cycloalkyl or an optionally substituted monocyclic heterocyclyl, the optionally substituted monocyclic C₃₋₆ cycloalkyl, the optionally substituted bicyclic C₃₋₈ cycloalkyl and the optionally substituted monocyclic heterocyclyl may substitute R¹² by replacing two hydrogens of R¹². For example, when R¹² is a C₃-alkenyl substituted with an oxetane (a monocyclic heterocyclyl) by replacing two hydrogens of R¹², R¹² may have the



structure. In some embodiments, R¹² is substituted with an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl or an optionally substituted monocyclic heterocyclyl, wherein the optionally substituted monocyclic C₃₋₆ cycloalkyl, the optionally substituted bicyclic C₃₋₈ cycloalkyl or the optionally substituted monocyclic heterocyclyl replaces one hydrogen. In other embodiments, R¹² is substituted with an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl or an optionally substituted monocyclic heterocyclyl, wherein the optionally substituted monocyclic C₃₋₆ cycloalkyl, the optionally substituted bicyclic C₃₋₈ cycloalkyl or the optionally substituted monocyclic heterocyclyl replaces two hydrogen such that the aforementioned moieties are connected in a spiro-fashion.

[0096] The alkynyl of R¹² can include a single triple bond. The position of the triple bond can vary. In some embodiments, the triple bond can be located between the terminal carbon and a carbon adjacent to the terminal carbon. In some embodiments, the triple bond can be located between the carbon adjacent to the seven-membered ring of Formula (I) and the next carbon away from the seven-membered ring of Formula (I).

[0097] When R¹² is an alkenyl substituted with R^{19A} or an alkynyl substituted with R^{19A}, R^{19A} can be an optionally substituted -O-acyl prodrug or an unsubstituted -O-linked α -amino acid prodrug. When R¹² is a substituted alkenyl or a substituted alkynyl substituted with an optionally substituted -O-acyl, the optionally substituted -O-acyl can have the structure -O-C(=O)R^{19A1}, wherein R^{19A1} can be an optionally substituted C₁₋₆ alkyl or an optionally substituted C₆ or C₁₄ aryl. In some embodiments, R^{19A} can be -O-C(=O)R^{19A1}, wherein R^{19A1} can be an unsubstituted C₁₋₆ alkyl.

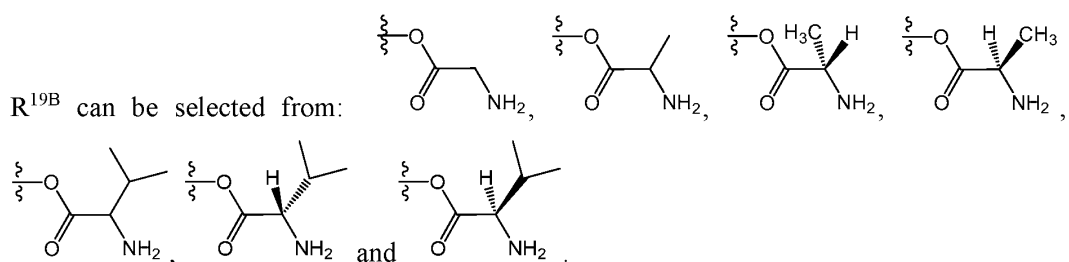
[0098] When the alkenyl and/or alkynyl of R¹² is substituted, each of the aforementioned can be substituted with an unsubstituted -O-linked α -amino acid, such as

those known in the art and described herein. In some embodiments, R^{19A} can be -O-linked glycine, -O-linked valine, -O-linked leucine or -O-linked isoleucine. A phosphate or a phosphonate can be present at R^{19A} . In some embodiments, R^{19A} can be $-O-P(=O)(OH)_2$. In other embodiments, R^{19A} can be $-CH_2-P(=O)(OH)_2$.

[0099] As provided herein, R^{13} can be hydrogen, deuterium or a substituted or an unsubstituted C_{1-4} alkyl, wherein the substituted C_{1-4} alkyl can be substituted with one or more substituents selected from halogen, hydroxy and R^{19B} . In some embodiments, R^{13} can be hydrogen. In other embodiments, R^{13} can be deuterium. In still other embodiments, R^{13} can be an unsubstituted C_{1-4} alkyl. In yet still other embodiments, R^{13} can be a substituted C_{1-4} alkyl, substituted with one or more substituents selected from halogen (for example F or Cl), hydroxy and R^{19B} . Examples of R^{13} begin a substituted C_{1-4} alkyl include $-CF_3$, $-CHF_2$, $-CH_2OH$ and $-CH(OH)CH_3$. In some embodiments, R^{19B} can be $-O-P(=O)(OH)_2$. In other embodiments, R^{19B} can be $-CH_2-P(=O)(OH)_2$.

[0100] Suitable C_{1-4} alkyls for R^{12} and R^{13} are described herein, and include methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl or tert-butyl. When, R^{13} is a substituted C_{1-4} alkyl, substituted with R^{19B} , R^{19B} can be an optionally substituted -O-acyl prodrug or an unsubstituted -O-linked α -amino acid prodrug. The optionally substituted -O-acyl of R^{19B} can be $-O-C(=O)R^{19B1}$, wherein R^{19B1} can be an optionally substituted C_{1-6} alkyl or an optionally substituted C_6 or C_{14} aryl. In some embodiments, R^{19B} can be $-O-C(=O)R^{19B1}$, wherein R^{19B1} can be an unsubstituted C_{1-6} alkyl.

[0101] As described herein, R^{12} and R^{13} can include an -O-linked α -amino acid, such as -O-linked glycine, -O-linked valine, -O-linked leucine or -O-linked isoleucine. When R^{19A} and/or R^{19B} is an -O-linked α -amino acid, in some embodiments, R^{19A} and/or



[0102] In some embodiments, R¹⁴ can be hydrogen. In other embodiments, R¹⁴ can be deuterium. In still other embodiments, R¹⁴ can be an unsubstituted C₁₋₄ alkyl such as those described herein. In some embodiments, R¹⁴ can be methyl.

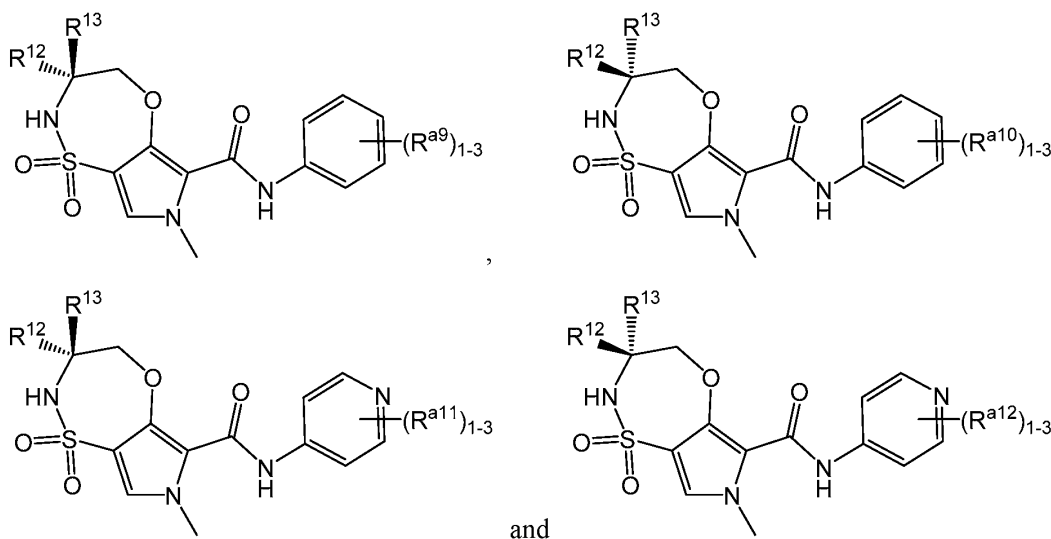
[0103] Formula (II) includes a 5-membered ring that can be unsubstituted or substituted with an unsubstituted C₁₋₄ alkyl, cyano and/or an unsubstituted C₁₋₄ haloalkyl. In some embodiments, R¹⁵ can be hydrogen. In other embodiments, R¹⁵ can be deuterium. In still other embodiments, R¹⁵ can be an unsubstituted C₁₋₄ alkyl (for example, methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl or tert-butyl). In some embodiments, R¹⁶ can be hydrogen. In other embodiments, R¹⁶ can be deuterium. In still other embodiments, R¹⁶ can be halogen (for example, F or Cl). In yet still other embodiments, R¹⁶ can be an unsubstituted C₁₋₄ alkyl such as those described herein. In some embodiments, R¹⁶ can be cyano. In other embodiments, R¹⁶ can be an unsubstituted C₁₋₄ haloalkyl, such as CF₃. In still other embodiments, R¹⁶ can be an unsubstituted C₃₋₈ monocyclic cycloalkyl. In some embodiments, R¹⁵ can be methyl; and R¹⁶ can be hydrogen.

[0104] The -C(=O)NR¹⁷R¹⁸ moiety of Formula (II) can include a substituted phenyl or a substituted pyridyl as described herein. In some embodiments, R¹⁷ can be a substituted phenyl, wherein the phenyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C₁₋₄ haloalkyl and an unsubstituted C₁₋₄ alkyl. In other embodiments, R¹⁷ can be a substituted pyridyl, wherein the pyridyl can be substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C₁₋₄ haloalkyl and an unsubstituted C₁₋₄ alkyl. The phenyl and pyridyl of R¹⁷ can be substituted with one or more substituents as described herein. In some embodiments, R¹⁷ can be a mono-substituted phenyl. In other embodiments, R¹⁷ can be a di-substituted phenyl. In some embodiments, the phenyl of R¹⁷ can be substituted at the para-position and/or meta-position. In some embodiments, the phenyl of R¹⁷ can be a 3,4-disubstituted phenyl. In still other embodiments, R¹⁷ can be a mono-substituted pyridyl. In yet still other embodiments, R¹⁷ can be a di-substituted pyridyl. In some embodiments, the pyridyl can be substituted on a carbon adjacent to the nitrogen of the pyridyl. The unsubstituted C₁₋₄ alkyl(s) that can be substituted on R¹⁷ can be methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl or tert-butyl. Examples of unsubstituted C₁₋₄ haloalkyls are CF₃, CHF₂ and CH₂F. In some embodiments, R¹⁷ can be substituted with F and/or Cl. In some

embodiments, R^{17} can be substituted with F, Cl and/or Br. In some embodiments, R^{17} can be substituted with CF_3 . In some embodiments, R^{17} can be substituted with CH_3 . When R^{17} is di-substituted, the two groups can be the same or different. In some embodiments, R^{17} cannot be substituted with 2 fluoros. For example, in some embodiments, R^{17} cannot be 3,4-difluorophenyl.

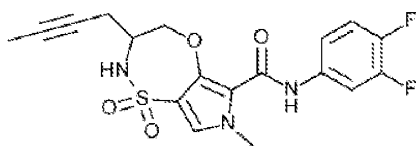
[0105] As described herein, R^{18} group of $-C(=O)NR^{17}R^{18}$ can be hydrogen, deuterium or an unsubstituted C_{1-4} alkyl. In some embodiments, R^{18} can be hydrogen. In other embodiments, R^{18} can be deuterium. In still other embodiments, R^{18} can be an unsubstituted C_{1-4} alkyl.

[0106] Some examples of compounds of Formula (II), including pharmaceutically acceptable salts thereof, include the following:

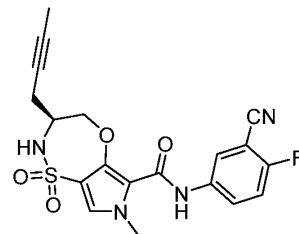


wherein each R^{a9} , each R^{a10} , each R^{a11} and each R^{a12} are independently selected from methyl, ethyl, n-propyl, iso-propyl, n-butyl, iso-butyl, tert-butyl, cyano, CF_3 , CHF_2 , CH_2F , F, Cl and Br.

[0107] In some embodiments, when R^{12} is an unsubstituted 2-butyryl, R^{13} is hydrogen, R^{14} and R^{18} are each hydrogen, R^{15} is methyl and R^{16} is hydrogen, then R^{17} cannot be 3,4-difluorophenyl. In some embodiments, a compound of Formula (II) cannot be



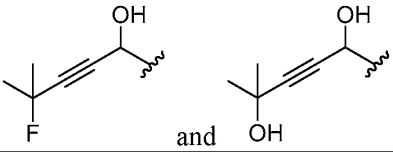
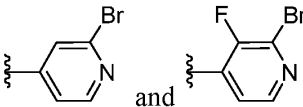
, or a pharmaceutically acceptable salt thereof. In



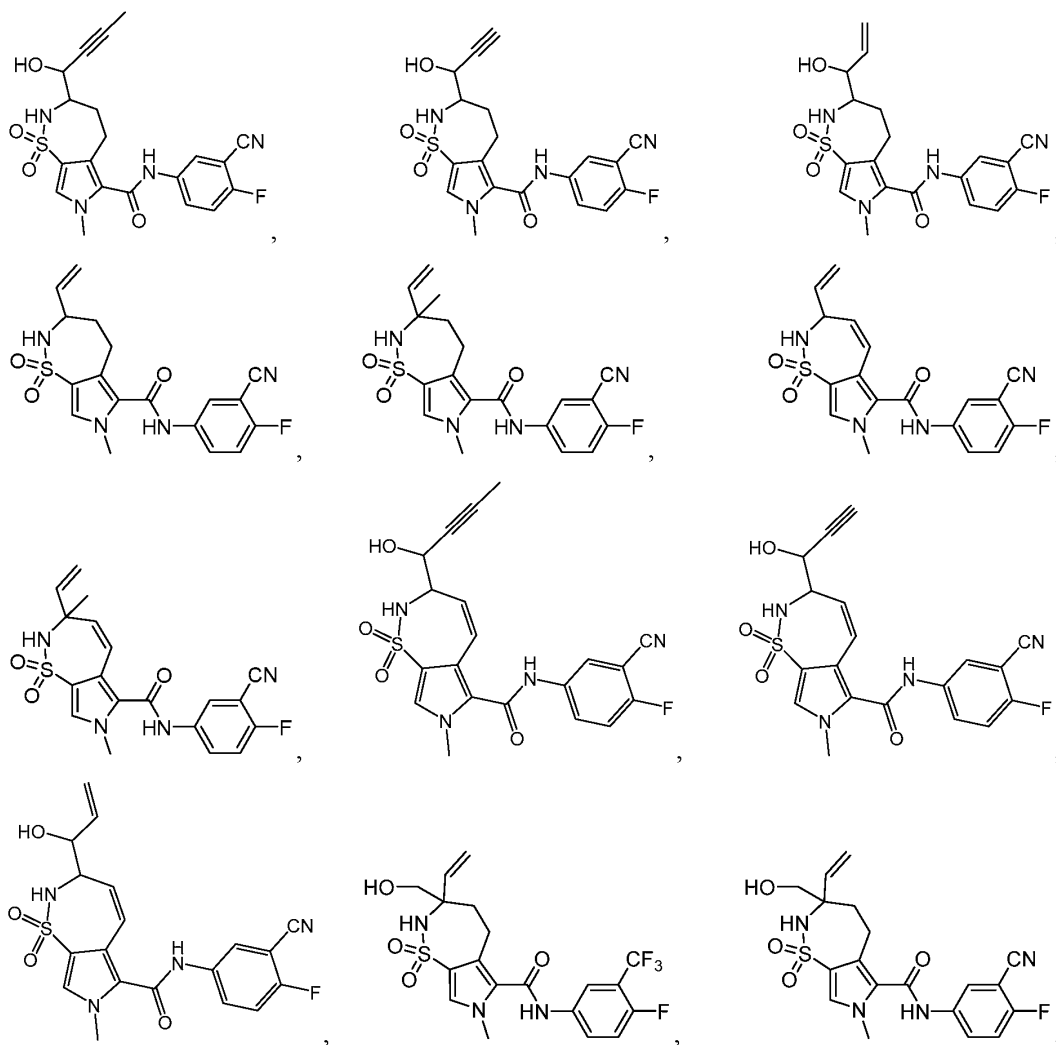
some embodiments, a compound of Formula (II) cannot be , or a pharmaceutically acceptable salt thereof.

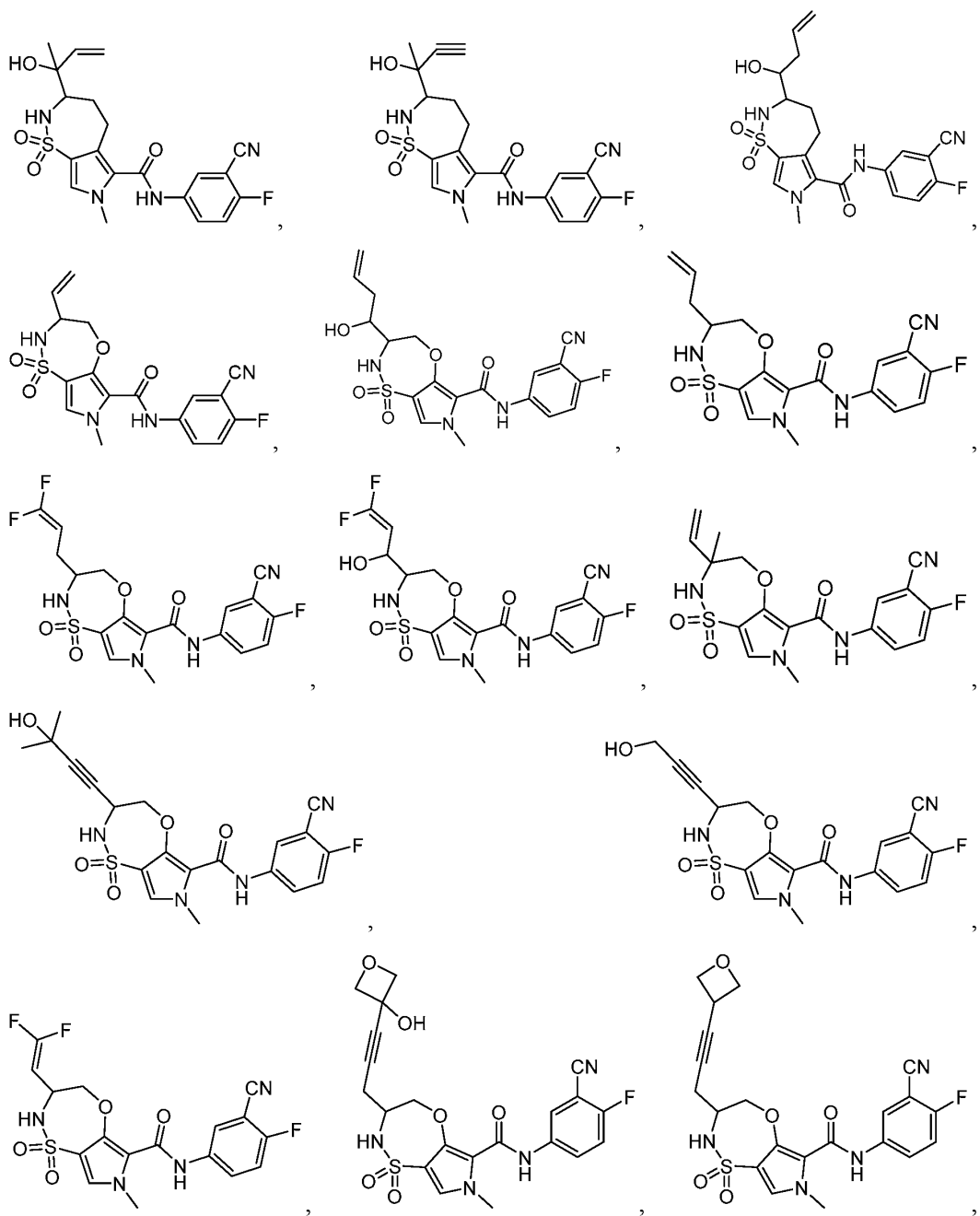
[0108] For Formula (I) and (II), exemplary R^1 , R^2 , R^{12} , R^{13} , R^6 and R^{17} moieties include, but are not limited to, the following:

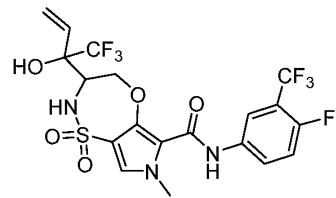
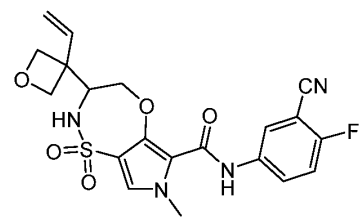
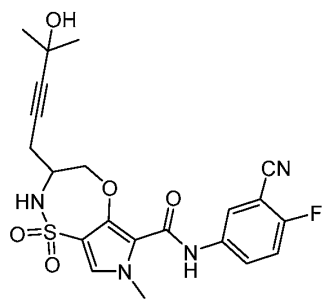
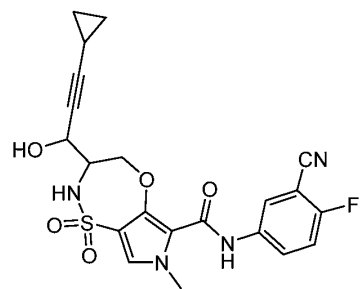
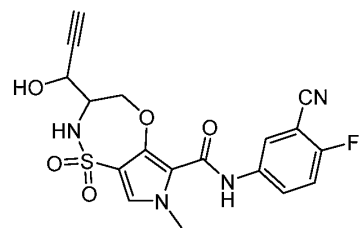
R^1/R^{12}	R^6/R^{17}

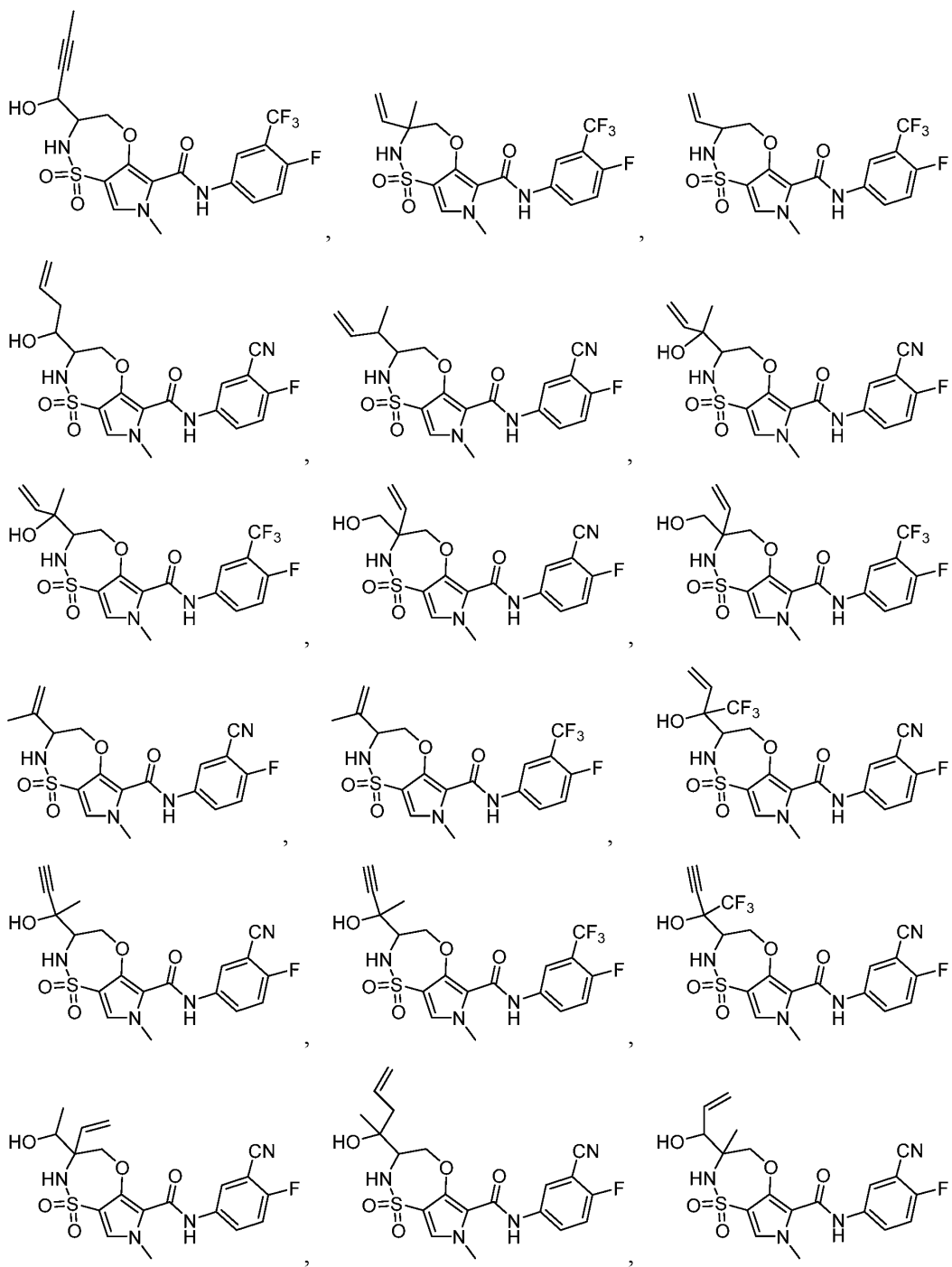
R^1/R^{12}	R^6/R^{17}
	
R^2/R^{13}	
-H, -CH ₃ , -CH ₂ OH and -CH(OH)CH ₃	

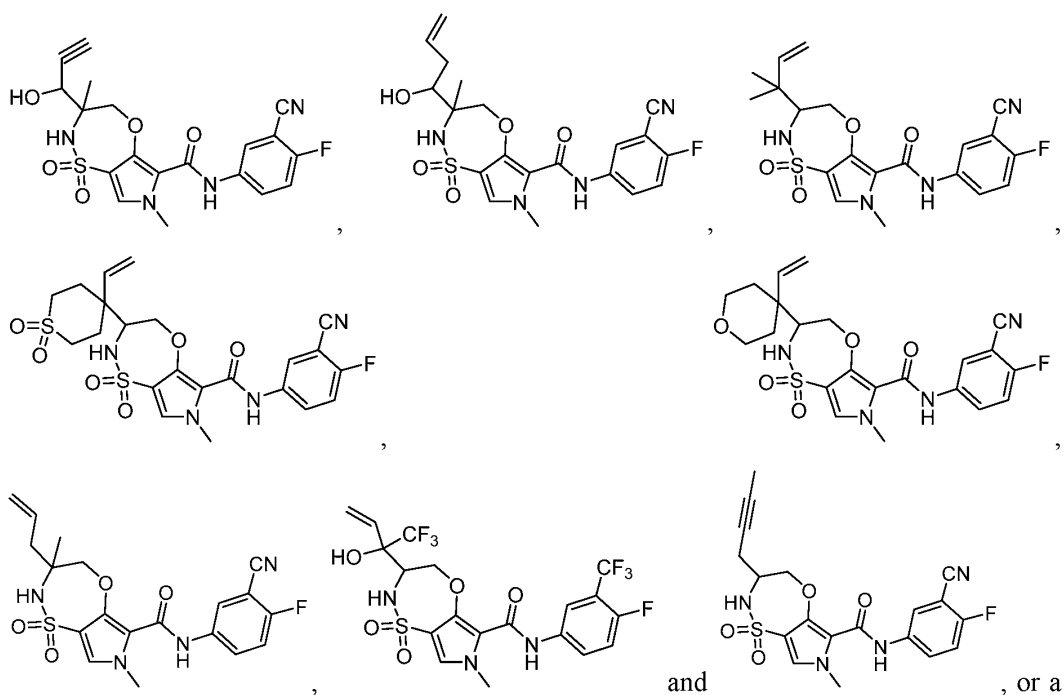
[0109] Examples of compounds of Formulae (I) and (II), or a pharmaceutically acceptable salt of any of the foregoing, include the following:





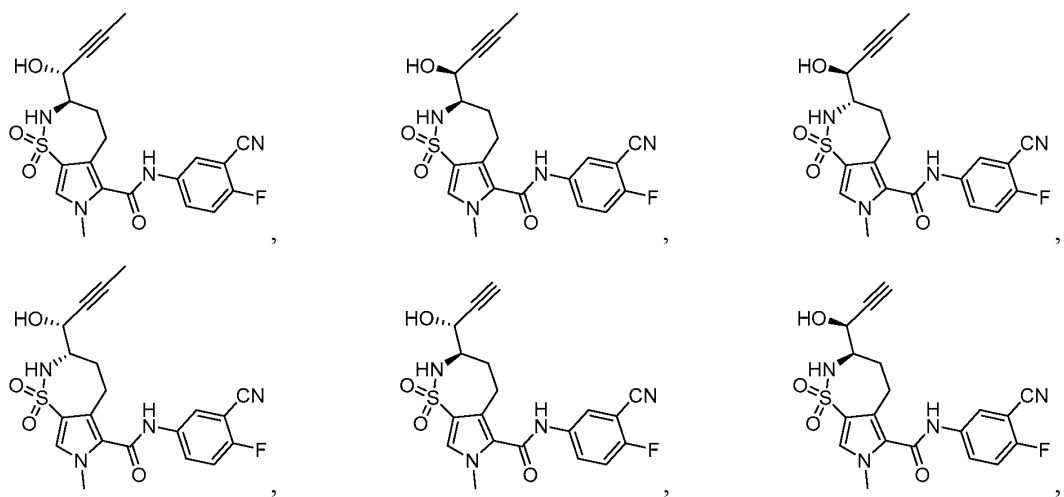


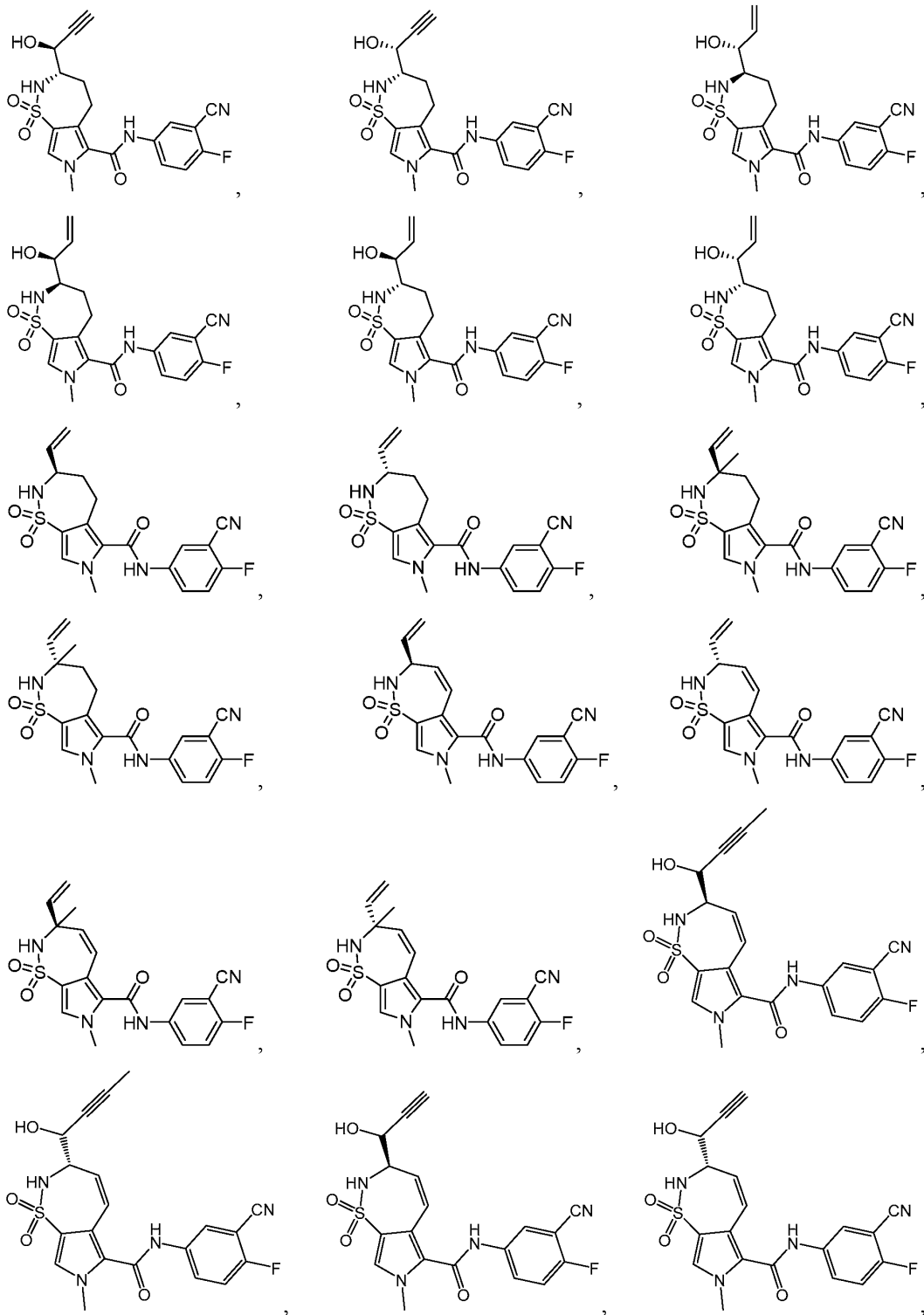


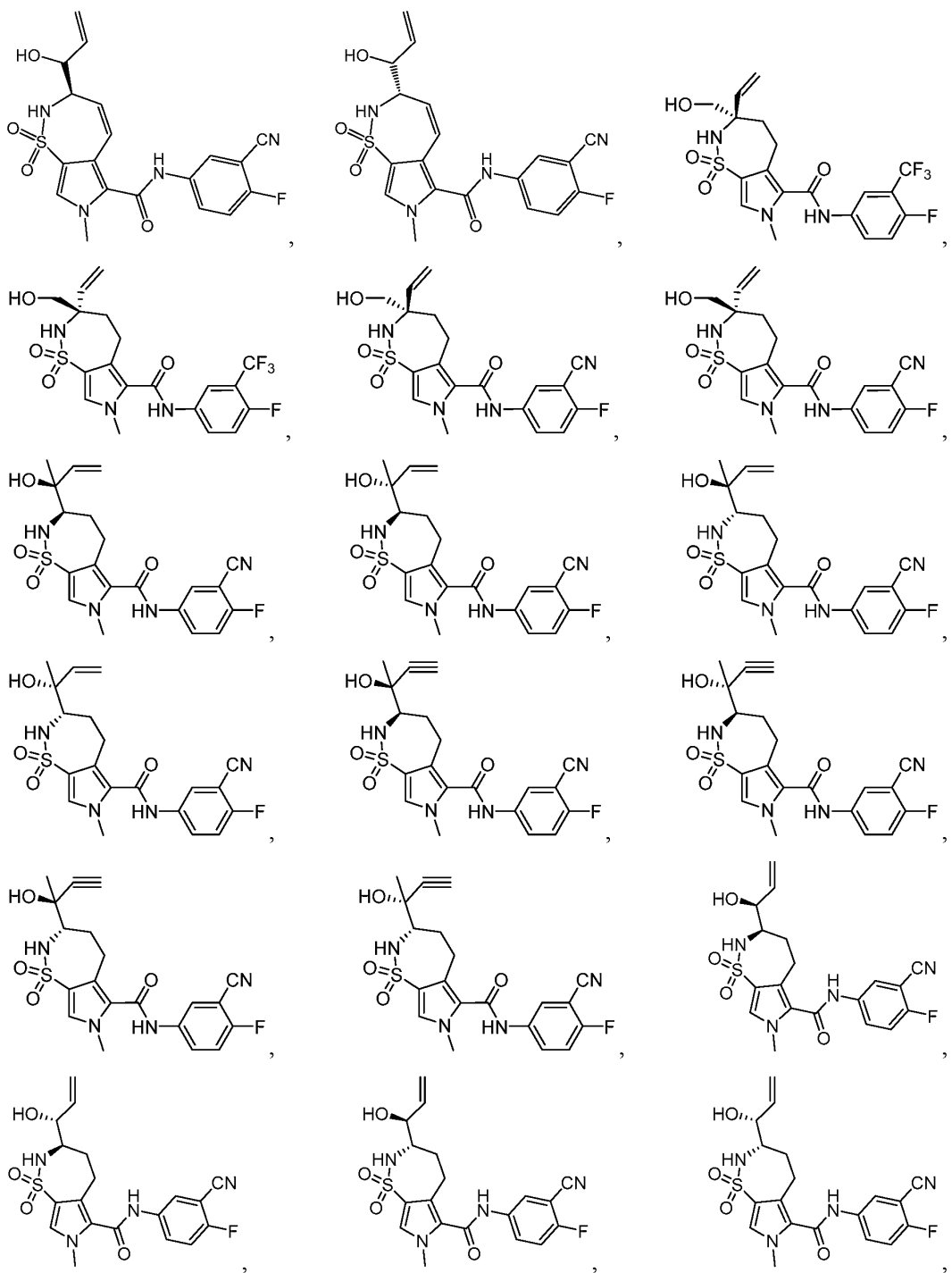


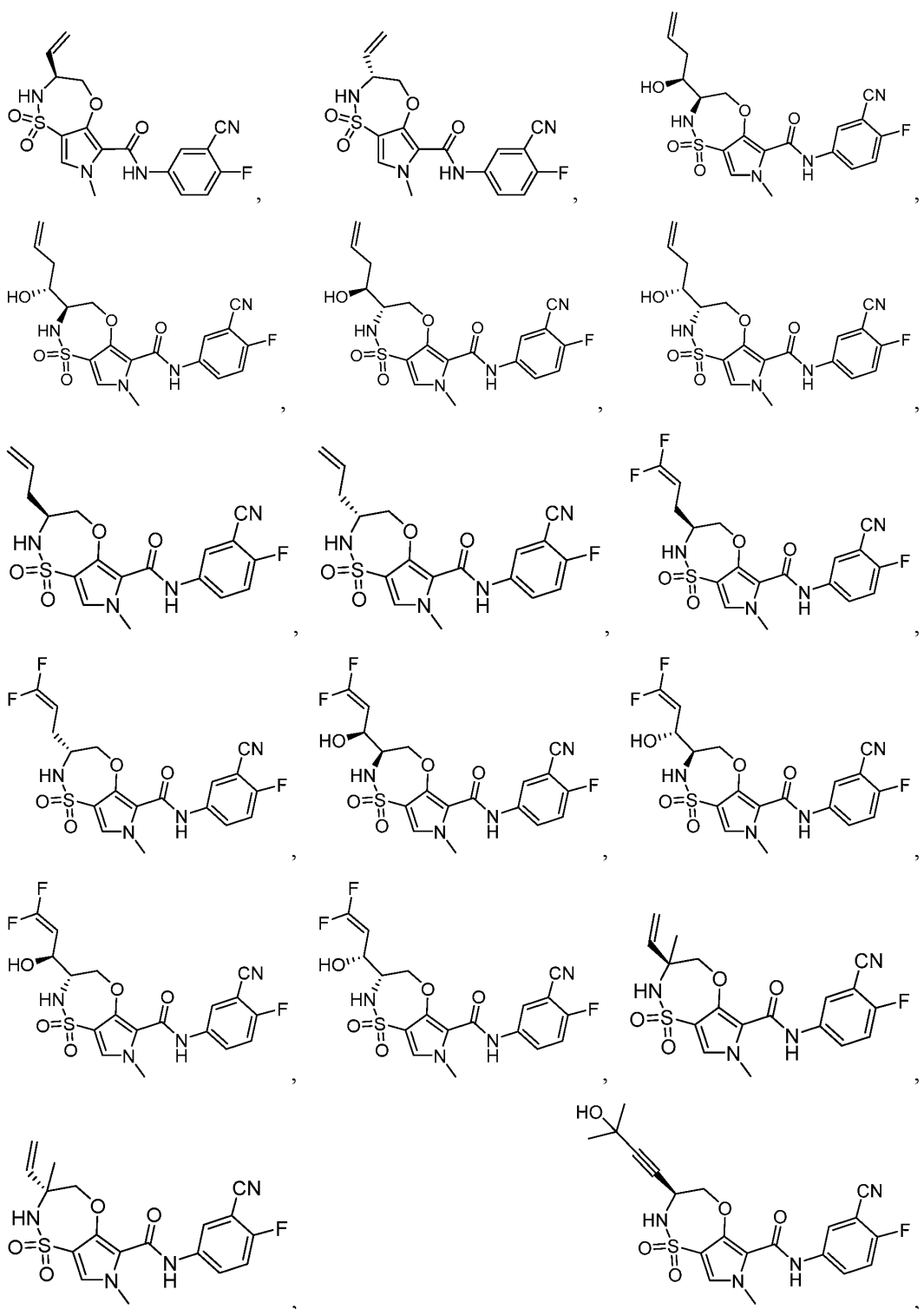
pharmaceutically acceptable salt of any of the foregoing.

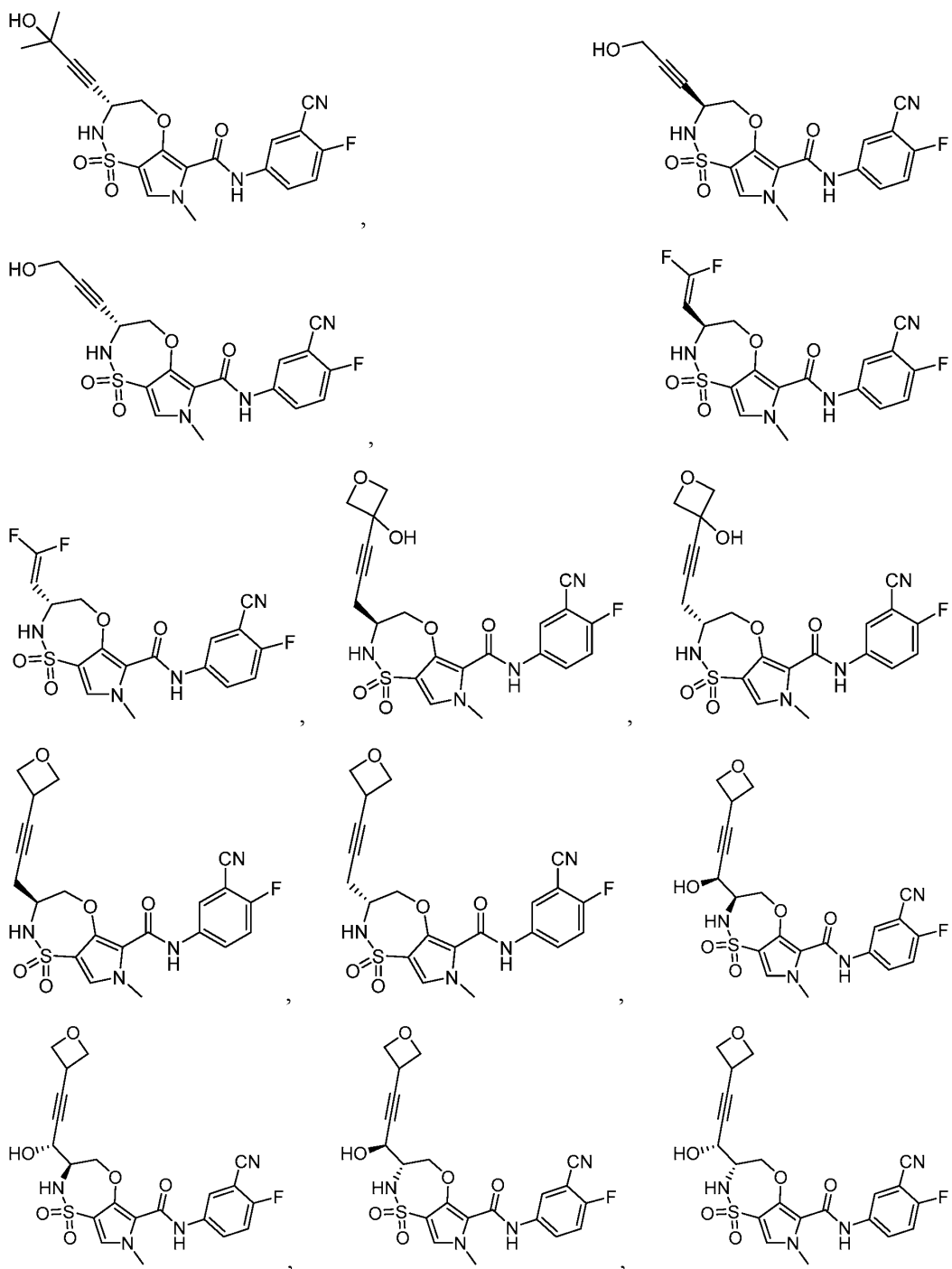
[0110] As provided herein, Compounds of Formulae (I) and (II) may include one or more chiral centers; and therefore, the compounds may exist as enantiomers and/or diastereomers.

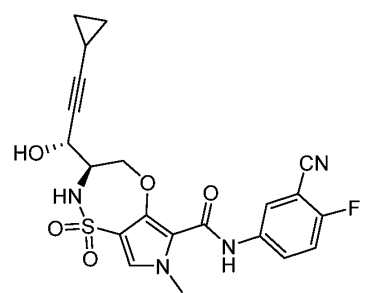
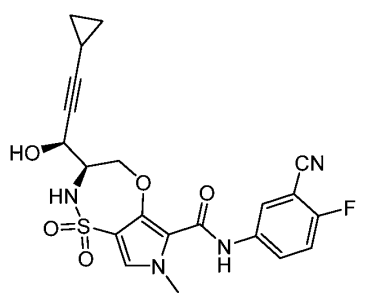
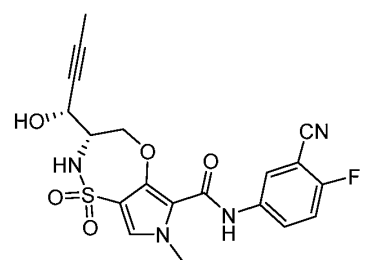
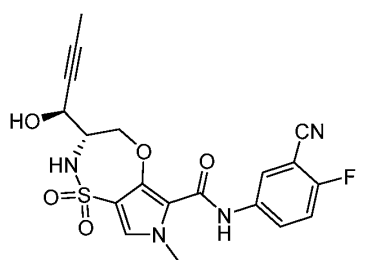
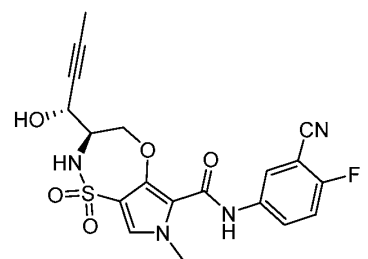
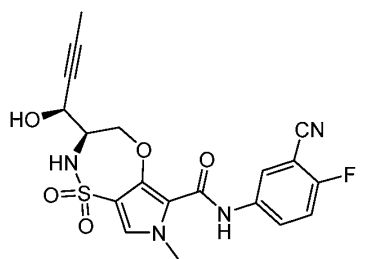
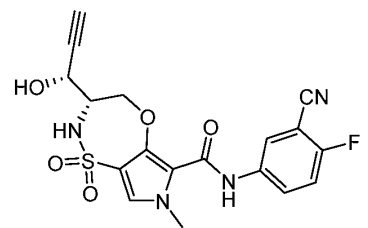
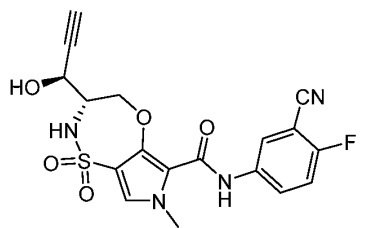
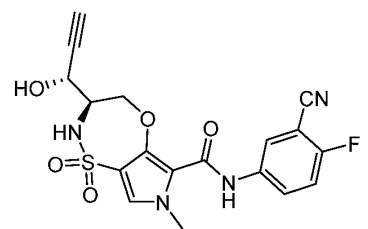
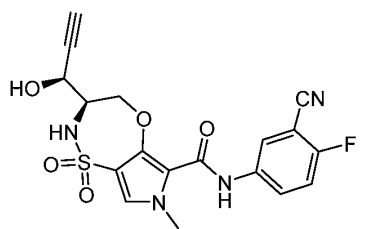


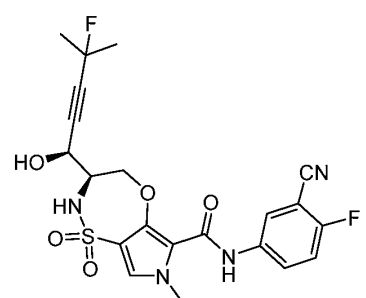
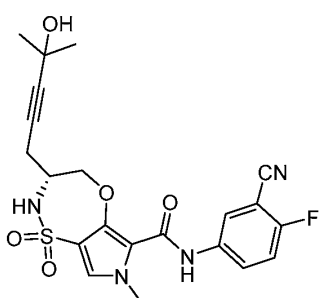
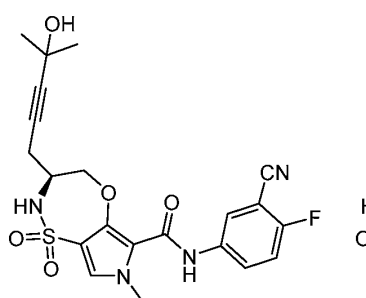
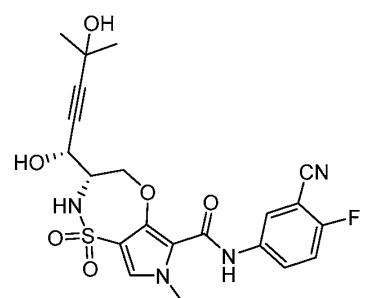
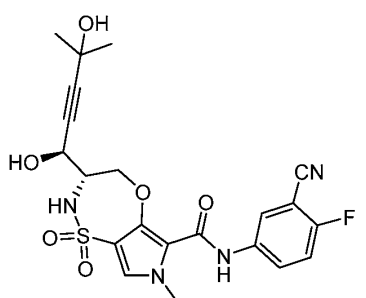
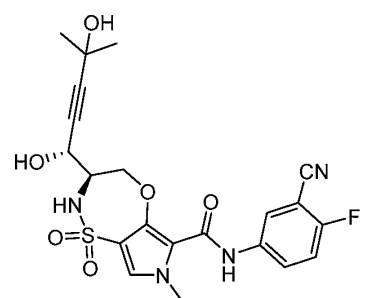
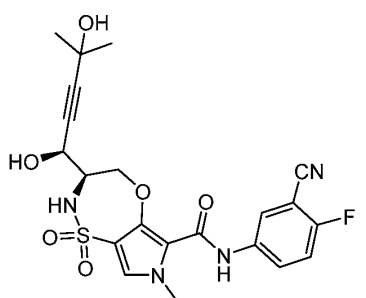
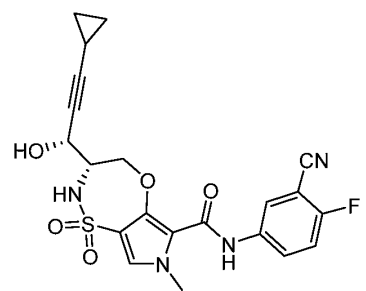
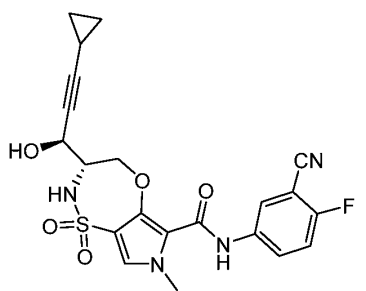


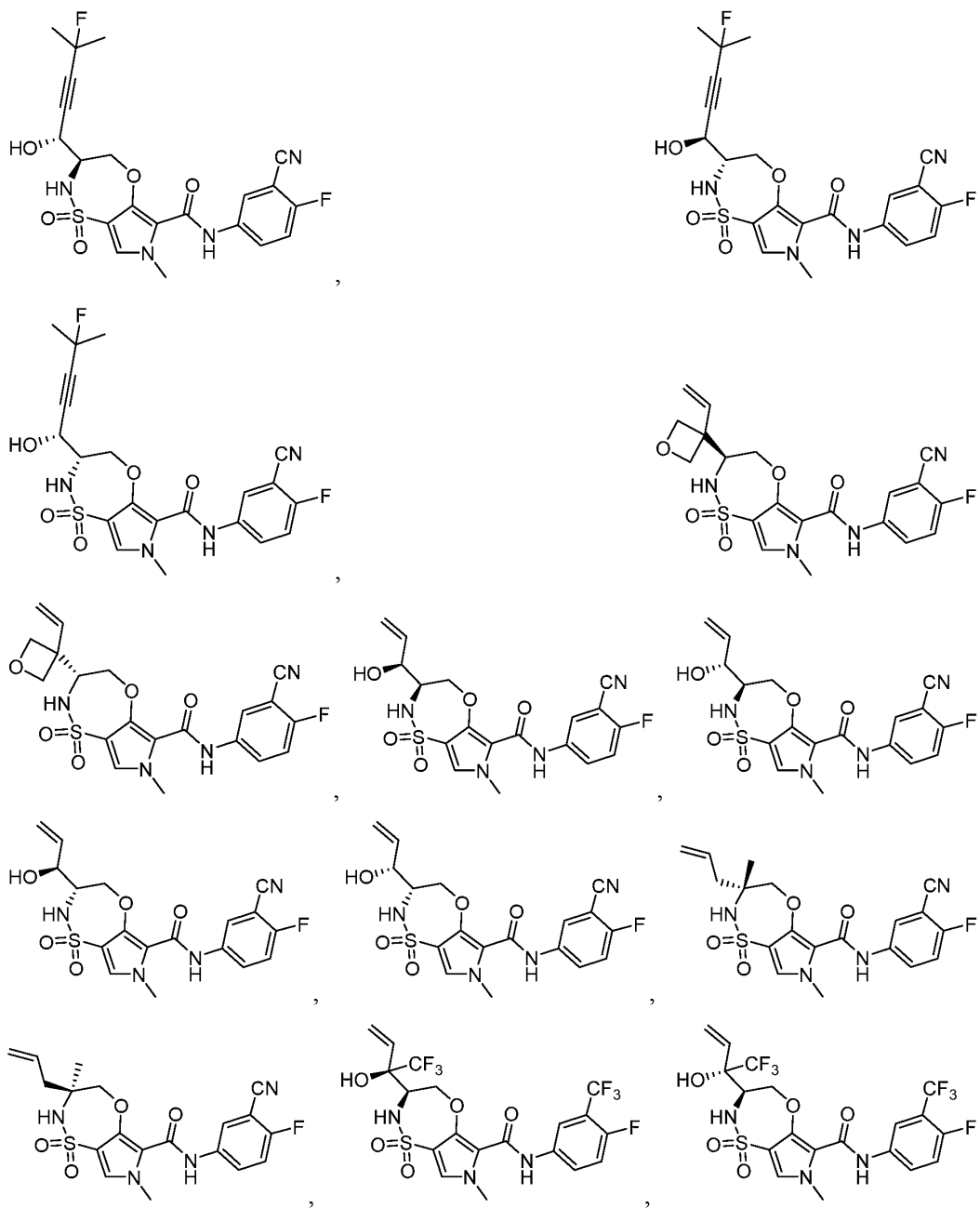


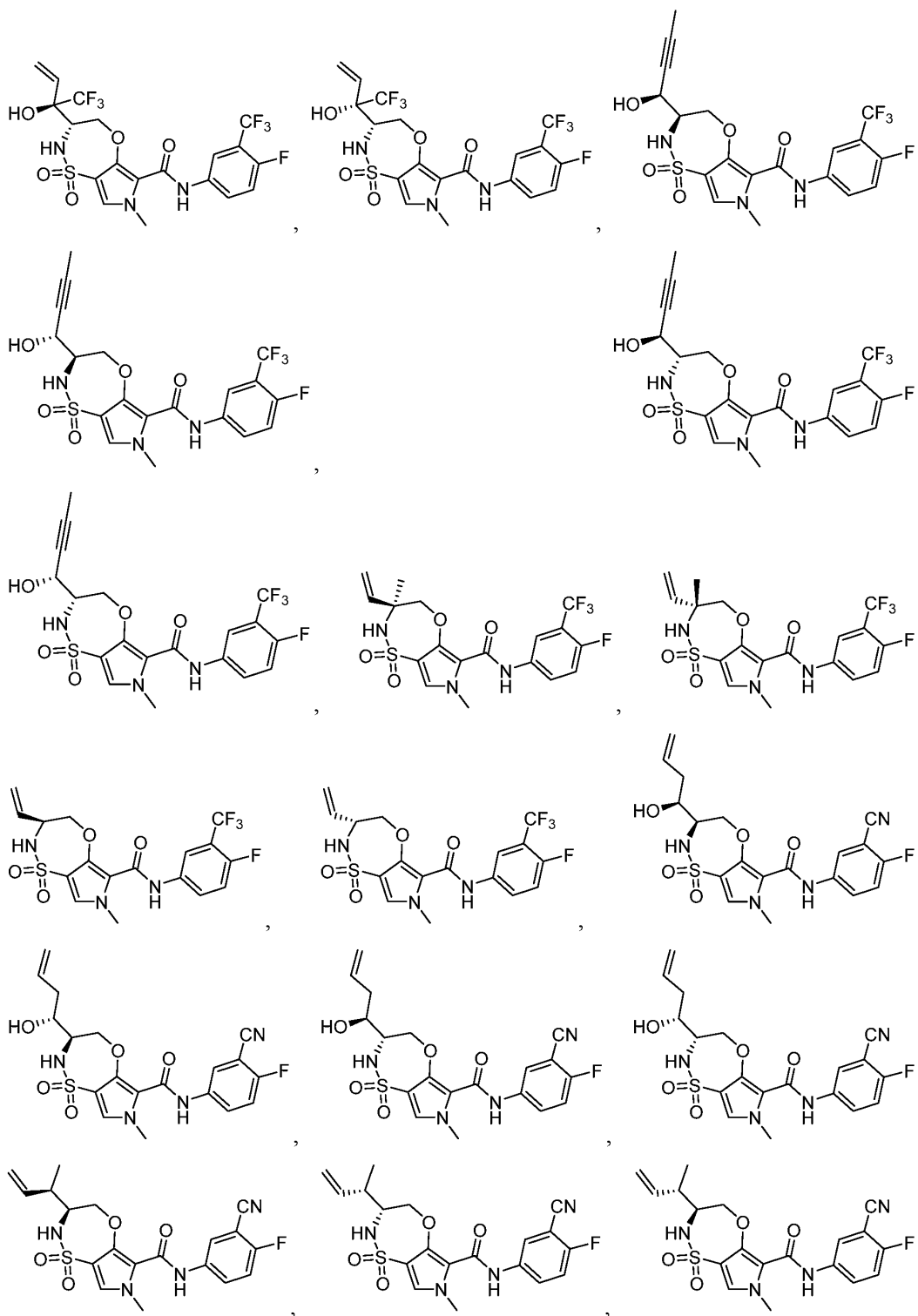


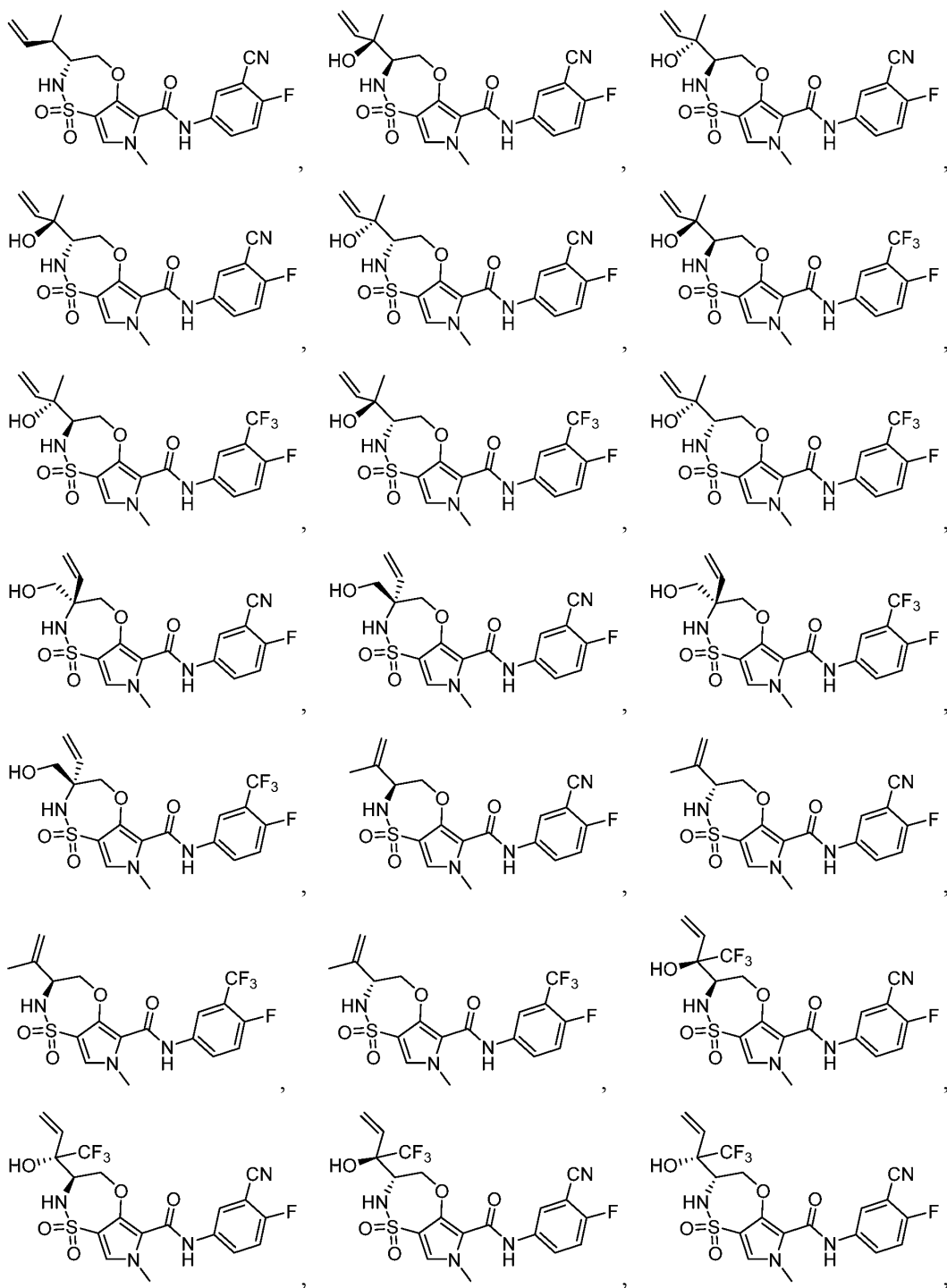


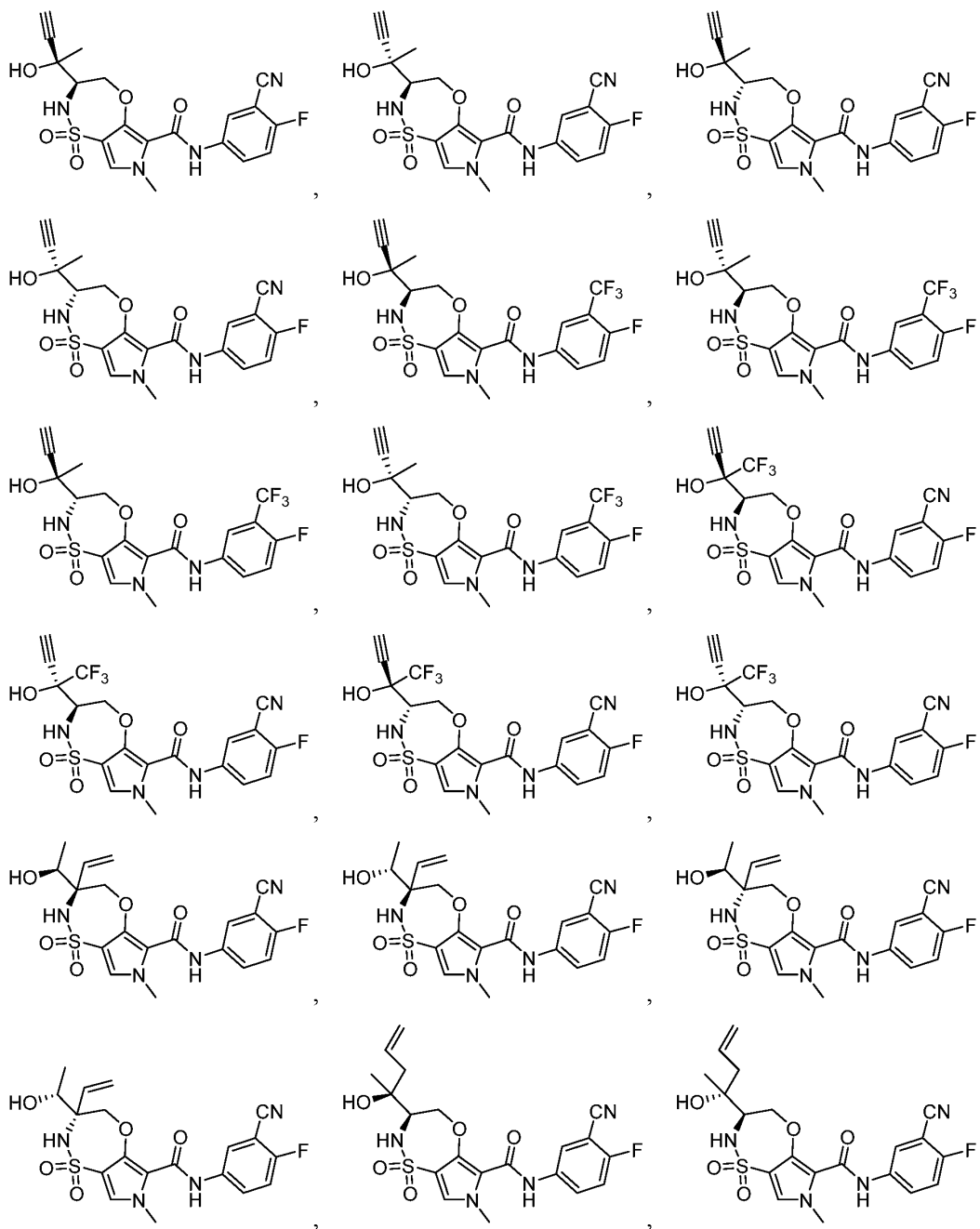


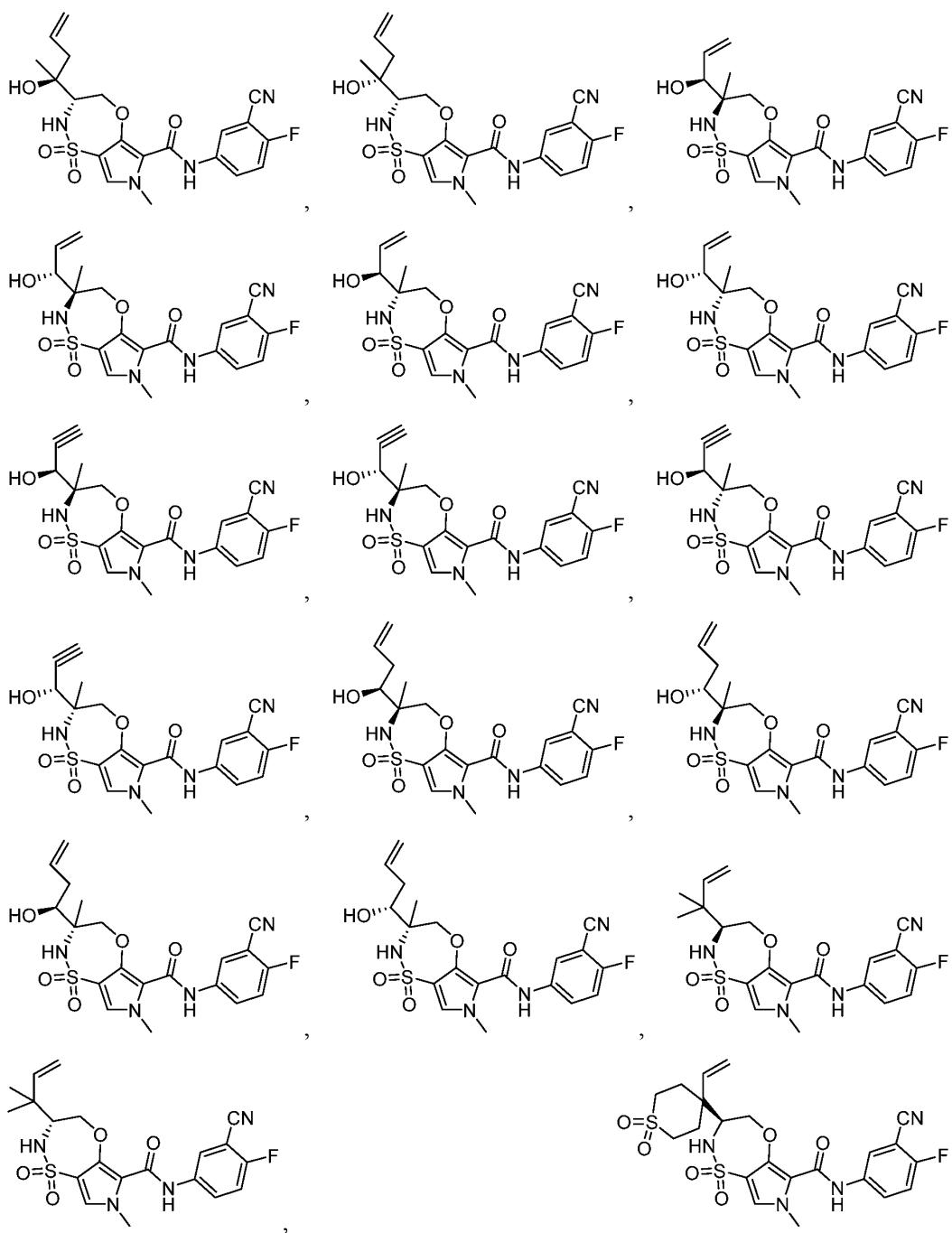


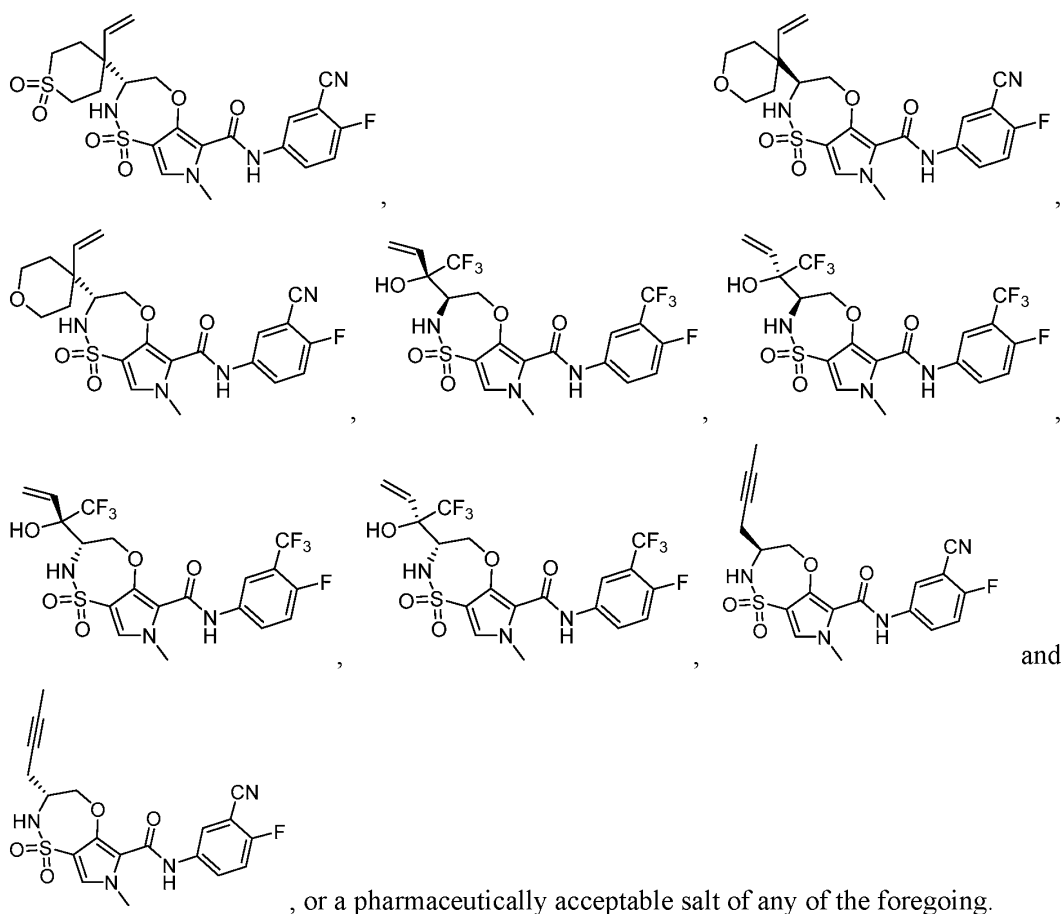








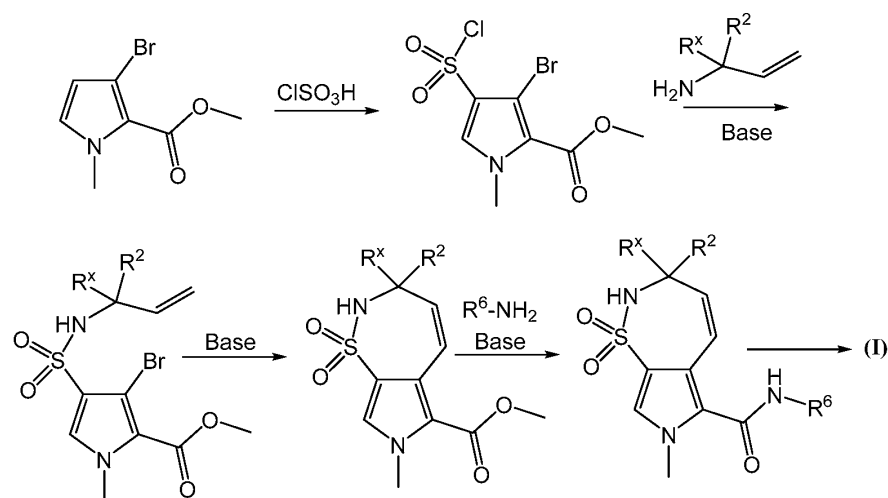




Synthesis

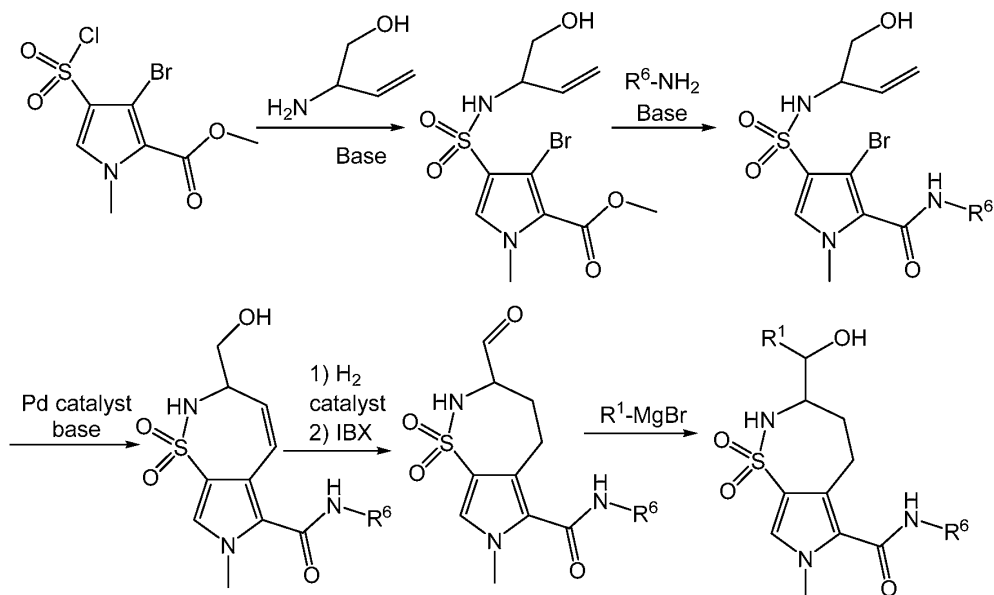
[0111] Compounds of Formulae (I) and (II) along with those described herein may be prepared in various ways. General synthetic routes for preparing compounds of Formulae (I) and (II) are shown and described herein along with some examples of starting materials used to synthesize compounds described herein. The routes shown and described herein are illustrative only and are not intended, nor are they to be construed, to limit the scope of the claims in any manner whatsoever. Those skilled in the art will be able to recognize modifications of the disclosed syntheses and to devise alternate routes based on the disclosures herein; all such modifications and alternate routes are within the scope of the claims.

Scheme 1



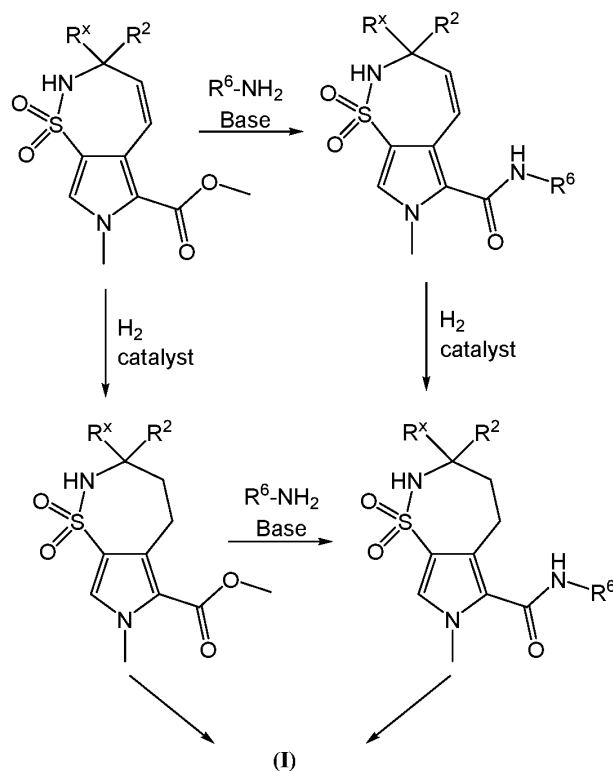
[0112] As shown in Scheme 1, sulfonyl chloride can be added to the pyrrolyl ring. The sulfonyl chloride can be then transformed to a sulfonamide utilizing an amine. Using a suitable base, the 7-membered ring can be formed, and the C-carboxy can be transformed to a C-amido using conditions known to those skilled in the art, such as a base and $R^6\text{-NH}_2$. In Scheme 1, R^x can be R^1 , or R^x can be transformed to R^1 during an appropriate step in the synthesis using methods known to those skilled in the art.

Scheme 2



[0113] Bicyclic sulfonamides described herein can be also prepared as shown in Scheme 2. After the addition of the sulfonyl chloride as shown in Scheme 1, a sulfonamide can be prepared by reacting the sulfonyl chloride attached to the pyrrolyl with an amine. The C-amido can be formed via an amidation reaction using a suitable base and R^6-NH_2 . Using a catalyst and base, for example a palladium catalyst, the saturated 7-membered ring can be formed. The hydroxyalkyl group attached to the saturated 7-membered ring can then be oxidized to an aldehyde using methods known to those skilled in the art (such as IBX). The R^1 group can then be added to the aldehyde to form the secondary alcohol using suitable conditions, such as a Grignard reaction. Using Wittig reaction conditions known to those skilled in the art, the aldehyde can be transformed to an alkenyl derivative.

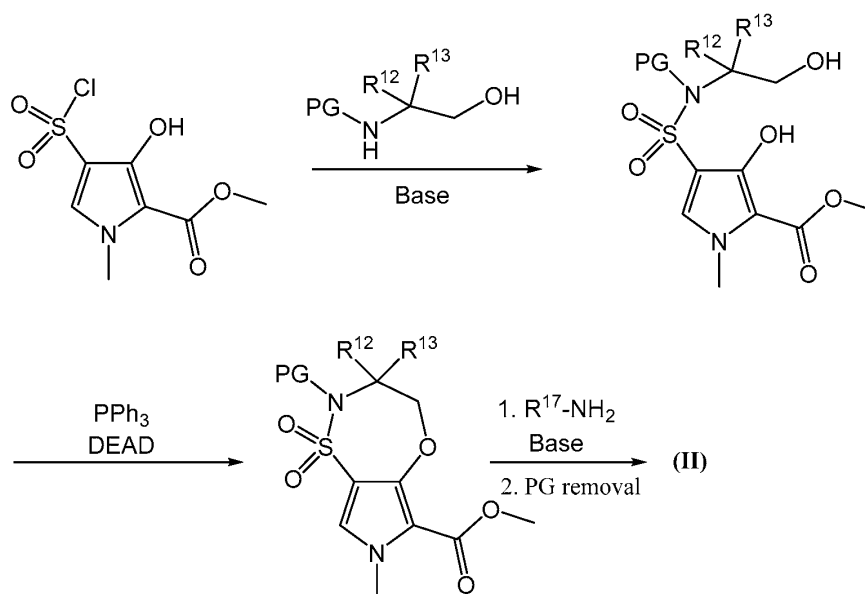
Scheme 3



[0114] As shown in Scheme 3, the compounds can be hydrogenated using a catalyst, such as nickel, palladium and/or platinum catalyst, and H_2 . The C-carboxy group can be transformed to C-amido via an amidation reaction prior to or after hydrogenation. In

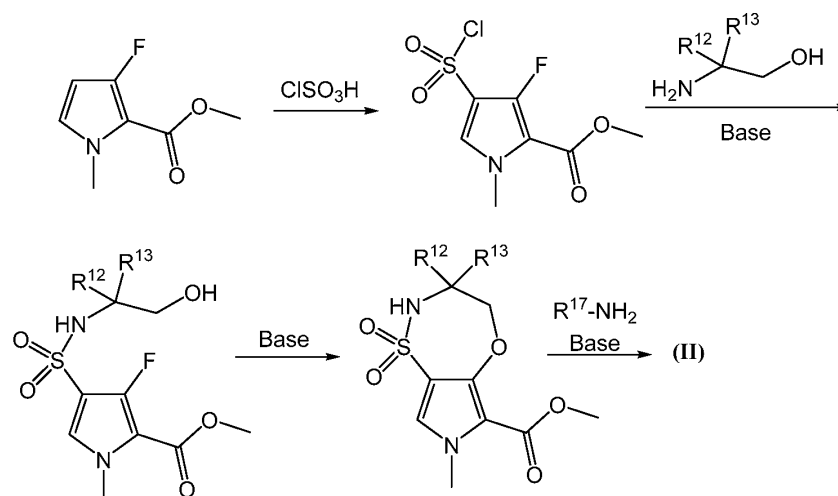
Scheme 3, R^x can be R^1 , or R^x can be transformed to R^1 during an appropriate step in the synthesis using methods known to those skilled in the art.

Scheme 4



[0115] In Scheme 4, “PG” represents a protecting group. The sulfonamide shown in Scheme 4 can be obtained as described herein, for example using a suitable amine and base. Utilizing Mitsunobu reaction conditions, the 7-membered oxygen-containing ring can be formed, using a protected aminoalcohol, for example, using a paramethoxybenzyl protecting group. Amidation of the C-carboxy forms the N-amido, and removal of the protecting group can provide the final compound.

Scheme 5



[0116] The shown bicyclic sulfonamides can be prepared in a similar manner as described for Scheme 1. The sulfonyl chloride can be added to the pyrrolyl. Formation of the sulfonamide can be accomplished using a suitable amine and base. Using a suitable base, the 7-membered ring can be formed. The C-carboxy can be converted utilizing a base and $R^{17}\text{-NH}_2$.

Pharmaceutical Compositions

[0117] Some embodiments described herein relate to a pharmaceutical composition, that can include an effective amount of a compound described herein (e.g., a compound, or a pharmaceutically acceptable salt thereof, as described herein) and a pharmaceutically acceptable carrier, excipient or combination thereof. A pharmaceutical composition described herein is suitable for human and/or veterinary applications.

[0118] As used herein, a “carrier” refers to a compound that facilitates the incorporation of a compound into cells or tissues. For example, without limitation, dimethyl sulfoxide (DMSO) is a commonly utilized carrier that facilitates the uptake of many organic compounds into cells or tissues of a subject.

[0119] As used herein, a “diluent” refers to an ingredient in a pharmaceutical composition that lacks pharmacological activity but may be pharmaceutically necessary or desirable. For example, a diluent may be used to increase the bulk of a potent drug whose

mass is too small for manufacture and/or administration. It may also be a liquid for the dissolution of a drug to be administered by injection, ingestion or inhalation. A common form of diluent in the art is a buffered aqueous solution such as, without limitation, phosphate buffered saline that mimics the composition of human blood.

[0120] As used herein, an “excipient” refers to an inert substance that is added to a pharmaceutical composition to provide, without limitation, bulk, consistency, stability, binding ability, lubrication, disintegrating ability etc., to the composition. A “diluent” is a type of excipient.

[0121] Pharmaceutical compositions may be formulated in a variety forms, such as tablets, capsules or solutions for oral administration; suppositories for rectal or vaginal administration; sterile solutions or suspensions for injectable administration. Injectables can be prepared in conventional forms, either as liquid solutions or suspensions, solid forms suitable for solution or suspension in liquid prior to injection, or as emulsions.

[0122] Proper formulation is dependent upon the route of administration chosen. Techniques for formulation and administration of the compounds described herein are known to those skilled in the art. Multiple techniques of administering a compound exist in the art including, but not limited to, oral, rectal, topical, aerosol, injection and parenteral delivery, including intramuscular, subcutaneous, intravenous, intramedullary injections, intrathecal, direct intraventricular, intraperitoneal, intranasal and intraocular injections. Pharmaceutical compositions will generally be tailored to the specific intended route of administration.

[0123] One may also administer the compound in a local rather than systemic manner, for example, via injection of the compound directly into the infected area, often in a depot or sustained release formulation. Furthermore, one may administer the compound in a targeted drug delivery system, for example, in a liposome coated with a tissue-specific antibody. The liposomes may be targeted to and taken up selectively by the organ.

[0124] The pharmaceutical compositions disclosed herein may be manufactured in a manner that is itself known, *e.g.*, by means of conventional mixing, dissolving, granulating, dragee-making, levigating, emulsifying, encapsulating, entrapping or tableting processes. As described herein, compounds used in a pharmaceutical composition may be provided as salts with pharmaceutically compatible counterions.

Methods of Use

[0125] Some embodiments described herein relate to a method of treating a HBV and/or HDV infection that can include administering to a subject identified as suffering from the HBV and/or HDV infection an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically acceptable salt thereof, as described herein in the manufacture of a medicament for treating a HBV and/or HDV infection. Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein or a pharmaceutical composition that includes a compound, or a pharmaceutically acceptable salt thereof, as described herein for treating a HBV and/or HDV infection.

[0126] Some embodiments disclosed herein relate to a method of treating a HBV and/or HDV infection that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically acceptable salt thereof, as described herein in the manufacture of a medicament for treating a HBV and/or HDV infection. Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for treating a HBV and/or HDV infection.

[0127] Some embodiments disclosed herein relate to a method of inhibiting replication of HBV and/or HDV that can include contacting a cell infected with the HBV and/or HDV with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically

acceptable salt thereof, as described herein in the manufacture of a medicament for inhibiting replication of HBV and/or HDV. Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, for inhibiting replication of HBV and/or HDV.

[0128] In some embodiments, the HBV infection can be an acute HBV infection. In some embodiments, the HBV infection can be a chronic HBV infection.

[0129] Some embodiments disclosed herein relate to a method of treating liver cirrhosis that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from liver cirrhosis and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from liver cirrhosis with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically acceptable salt thereof, as described herein in the manufacture of a medicament for treating liver cirrhosis with an effective amount of the compound, or a pharmaceutically acceptable salt thereof. Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for treating liver cirrhosis.

[0130] Some embodiments disclosed herein relate to a method of treating liver cancer (such as hepatocellular carcinoma) that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from the liver cancer and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from the liver cancer with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically acceptable salt thereof, as described herein in the manufacture of a medicament for treating

liver cancer (such as hepatocellular carcinoma). Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for treating liver cancer (such as hepatocellular carcinoma).

[0131] Some embodiments disclosed herein relate to a method of treating liver failure that is developed because of a HBV and/or HDV infection that can include administering to a subject suffering from liver failure and/or contacting a cell infected with the HBV and/or HDV in a subject suffering from liver failure with an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein. Other embodiments described herein relate to using a compound, or a pharmaceutically acceptable salt thereof, as described herein in the manufacture of a medicament for treating liver failure. Still other embodiments described herein relate to the use of a compound, or a pharmaceutically acceptable salt thereof, as described herein, or a pharmaceutical composition that includes an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein for treating liver failure.

[0132] Various indicators for determining the effectiveness of a method for treating an HBV and/or HDV infection are also known to those skilled in the art. Examples of suitable indicators include, but are not limited to, a reduction in viral load indicated by reduction in HBV DNA (or load) (e.g., reduction $<10^5$ copies/mL in serum), HBV surface antigen (HBsAg) and HBV e-antigen (HBeAg), a reduction in plasma viral load, a reduction in viral replication, a reduction in time to seroconversion (virus undetectable in patient serum), an increase in the rate of sustained viral response to therapy, an improvement in hepatic function, and/or a reduction of morbidity or mortality in clinical outcomes.

[0133] As used herein, the terms “treat,” “treating,” “treatment,” “therapeutic,” and “therapy” do not necessarily mean total cure or abolition of the disease or condition. Any alleviation of any undesired signs or symptoms of a disease or condition, to any extent can be considered treatment and/or therapy. Furthermore, treatment may include acts that may worsen the subject’s overall feeling of well-being or appearance.

[0134] As used herein, a “subject” refers to an animal that is the object of treatment, observation or experiment. “Animal” includes cold- and warm-blooded vertebrates and invertebrates such as fish, shellfish, reptiles and, in particular, mammals. “Mammal” includes, without limitation, mice, rats, rabbits, guinea pigs, dogs, cats, sheep, goats, cows, horses, primates, such as monkeys, chimpanzees, and apes, and, in particular, humans. In some embodiments, the subject is human.

[0135] The term “effective amount” is used to indicate an amount of an active compound, or pharmaceutical agent, that elicits the biological or medicinal response indicated. For example, an effective amount of compound can be the amount needed to alleviate or ameliorate symptoms of disease or prolong the survival of the subject being treated. This response may occur in a tissue, system, animal or human and includes alleviation of the signs or symptoms of the disease being treated. Determination of an effective amount is well within the capability of those skilled in the art, in view of the disclosure provided herein. The effective amount of the compounds disclosed herein required as a dose will depend on the route of administration, the type of animal, including human, being treated, and the physical characteristics of the specific animal under consideration. The dose can be tailored to achieve a desired effect, but will depend on such factors as weight, diet, concurrent medication and other factors which those skilled in the medical arts will recognize.

[0136] In some embodiments, an effective amount of a compound, or a pharmaceutically acceptable salt thereof, as described herein is an amount that is effective to achieve a sustained virologic response, for example, a sustained viral response 12 month after completion of treatment.

[0137] Subjects who are clinically diagnosed with a HBV and/or HDV infection include “naïve” subjects (e.g., subjects not previously treated for HBV and/or HDV) and subjects who have failed prior treatment for HBV and/or HDV (“treatment failure” subjects). Treatment failure subjects include “non-responders” (subjects who did not achieve sufficient reduction in ALT (alanine aminotransferase) levels, for example, subject who failed to achieve more than 1 log₁₀ decrease from base-line within 6 months of starting an anti-HBV and/or anti-HDV therapy) and “relapsers” (subjects who were previously treated for HBV and/or HDV whose ALT levels have increased, for example, ALT > twice the upper normal

limit and detectable serum HBV DNA by hybridization assays). Further examples of subjects include subjects with a HBV and/or HDV infection who are asymptomatic.

[0138] In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be provided to a treatment failure subject suffering from HBV and/or HDV. In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be provided to a non-responder subject suffering from HBV and/or HDV. In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be provided to a relapser subject suffering from HBV and/or HDV. In some embodiments, the subject can have HBeAg positive chronic hepatitis B. In some embodiments, the subject can have HBeAg negative chronic hepatitis B. In some embodiments, the subject can have liver cirrhosis. In some embodiments, the subject can be asymptomatic, for example, the subject can be infected with HBV and/or HDV but does not exhibit any symptoms of the viral infection. In some embodiments, the subject can be immunocompromised. In some embodiments, the subject can be undergoing chemotherapy.

[0139] Examples of agents that have been used to treat HBV and/or HDV include immunomodulating agents, and nucleosides/nucleotides. Examples of immunomodulating agents include interferons (such as IFN- α and pegylated interferons that include PEG-IFN- α -2a); and examples of nucleosides/nucleotides include lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide and tenofovir disoproxil. However, some of the drawbacks associated with interferon treatment are the adverse side effects, the need for subcutaneous administration and high cost. Potential advantages of a compound of Formula (I) and/or (II), or a pharmaceutically acceptable salt of any of the foregoing, can be less adverse side effects, delay in the onset of an adverse side effect and/or reduction in the severity of an adverse side effect. A drawback with nucleoside/nucleotide treatment can be the development of resistance, including cross-resistance.

[0140] Resistance can be a cause for treatment failure. The term “resistance” as used herein refers to a viral strain displaying a delayed, lessened and/or null response to an anti-viral agent. In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be provided to a subject infected with an HBV and/or HDV strain that is resistant to one or more anti-HBV and/or anti-HDV agents. Examples of anti-

viral agents wherein resistance can develop include lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir atafenamide and tenofovir disoproxil. In some embodiments, development of resistant HBV and/or HDV strains is delayed when a subject is treated with a compound, or a pharmaceutically acceptable salt thereof, as described herein compared to the development of HBV and/or HDV strains resistant to other HBV and/or HDV anti-viral agents, such as those described.

[0141] The dosage may range broadly, depending upon the desired effects and the therapeutic indication. Alternatively, dosages may be based and calculated upon the surface area of the patient, as understood by those of skill in the art. Although the exact dosage will be determined on a drug-by-drug basis, in most cases, some generalizations regarding the dosage can be made. The daily dosage regimen for an adult human patient may be, for example, an oral dose of between 0.01 mg and 3000 mg of each active ingredient, preferably between 1 mg and 700 mg, e.g. 5 to 200 mg. The dosage may be a single one or a series of two or more given in the course of one or more days, as is needed by the subject.

[0142] In instances where human dosages for compounds have been established for at least some condition, those same dosages may be used, or dosages that are between about 0.1% and 500%, more preferably between about 25% and 250% of the established human dosage. Where no human dosage is established, as will be the case for newly-discovered pharmaceutical compositions, a suitable human dosage can be inferred from ED₅₀ or ID₅₀ values, or other appropriate values derived from *in vitro* or *in vivo* studies, as qualified by toxicity studies and efficacy studies in animals.

[0143] In cases of administration of a pharmaceutically acceptable salt, dosages may be calculated as the free base. As will be understood by those of skill in the art, in certain situations it may be necessary to administer the compounds disclosed herein in amounts that exceed, or even far exceed, the above-stated, preferred dosage range in order to effectively and aggressively treat particularly aggressive diseases or infections.

[0144] Dosage amount and interval may be adjusted individually to provide plasma levels of the active moiety which are sufficient to maintain the modulating effects, or minimal effective concentration (MEC). The MEC will vary for each compound but can be estimated from *in vitro* data. Dosages necessary to achieve the MEC will depend on individual characteristics and route of administration. However, HPLC assays or bioassays

can be used to determine plasma concentrations. Dosage intervals can also be determined using MEC value. Compositions should be administered using a regimen which maintains plasma levels above the MEC for 10-90% of the time, preferably between 30-90% and most preferably between 50-90%. In cases of local administration or selective uptake, the effective local concentration of the drug may not be related to plasma concentration.

[0145] It should be noted that the attending physician would know how to and when to terminate, interrupt, or adjust administration due to toxicity or organ dysfunctions. Conversely, the attending physician would also know to adjust treatment to higher levels if the clinical response were not adequate (precluding toxicity). The magnitude of an administered dose in the management of the disorder of interest will vary with the severity of the condition to be treated and to the route of administration. The severity of the condition may, for example, be evaluated, in part, by standard prognostic evaluation methods. Further, the dose and perhaps dose frequency, will also vary according to the age, body weight, and response of the individual patient. A program comparable to that discussed above may be used in veterinary medicine.

[0146] Compounds disclosed herein can be evaluated for efficacy and toxicity using known methods. For example, the toxicology of a particular compound, or of a subset of the compounds, sharing certain chemical moieties, may be established by determining *in vitro* toxicity towards a cell line, such as a mammalian, including a human cell line. The results of such studies are often predictive of toxicity in animals, such as mammals, or more specifically, humans. Alternatively, the toxicity of particular compounds in an animal model, such as mice, rats, rabbits, or monkeys, may be determined using known methods. The efficacy of a particular compound may be established using several recognized methods, such as *in vitro* methods, animal models, or human clinical trials. When selecting a model to determine efficacy, the skilled artisan can be guided by the state of the art to choose an appropriate model, dose, route of administration and/or regime.

Combination Therapies

[0147] In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be used in combination with one or more additional agent(s) for treating and/or inhibiting replication HBV and/or HDV. Additional agents include, but

are not limited to, an interferon, nucleoside/nucleotide analogs, a capsid assembly modulator, a sequence specific oligonucleotide (such as anti-sense oligonucleotide and siRNA), nucleic acid polymers (NAPs) (such as STOPSTTM compounds and other nucleic acid polymers that reduce HBsAg levels) an entry inhibitor and/or a small molecule immunomodulator. Examples of additional agents include recombinant interferon alpha 2b, IFN- α , PEG-IFN- α -2a, lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide, tenofovir disoproxil, JNJ-6379, GLS4, ABI-HO731, JNJ-440, NZ-4, RG7907, AB-423, AB-506 and ABI-H2158. Examples of NAPs include, but are not limited to, REP 2139, REP 2165 and those STOPSTTM compounds described in U.S. Application No. 16/676,929, filed November 7, 2019, which is hereby incorporated by reference for the purpose of its disclosure of the STOPSTTM compounds described in the aforementioned U.S. application.

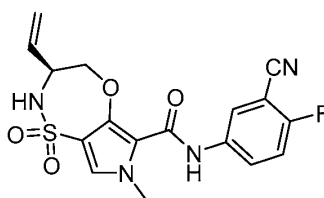
[0148] In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, as described herein can be administered with one or more additional agent(s) together in a single pharmaceutical composition. In some embodiments, a compound, or a pharmaceutically acceptable salt thereof, can be administered with one or more additional agent(s) as two or more separate pharmaceutical compositions. Further, the order of administration of a compound, or a pharmaceutically acceptable salt thereof, as described herein with one or more additional agent(s) can vary.

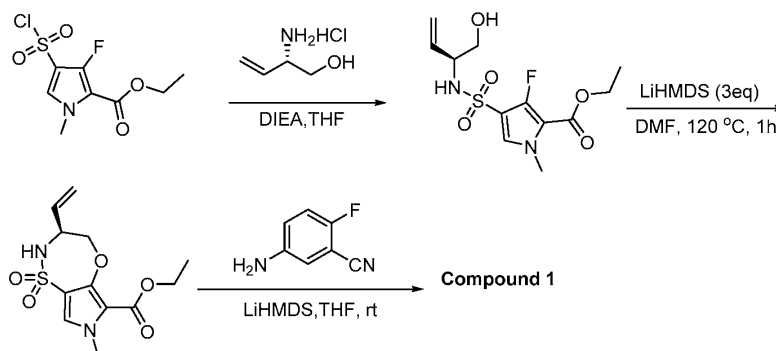
EXAMPLES

[0149] Additional embodiments are disclosed in further detail in the following examples, which are not in any way intended to limit the scope of the claims.

Example 1

(3S)-N-(3-cyano-4-fluorophenyl)-3-ethenyl-7-methyl-1,1-dioxo-2H,3H,4H-11 λ 6-pyrrolo[3,4-b][1,4,5]oxathiazepine-6-carboxamide (Compound 1)





[0150] To a stirred solution/mixture of ethyl 4-(chlorosulfonyl)-3-fluoro-1-methylpyrrole-2-carboxylate (1000 mg, 3.7 mmol, 1.00 eq.) and (2S)-1-hydroxybut-3-en-2-aminium chloride (687.4 mg, 5.6 mmol, 1.50 eq.) in THF (20 mL) were added DIEA (2.39 g, 18.5 mmol, 5.00 eq.) dropwise at room temperature (rt) under N₂ atmosphere. The resulting was stirred overnight at rt under N₂ atmosphere. The mixture was diluted with H₂O (30 mL) and extracted with EtOAc (3 x 20 mL). The combined organic layers were washed with brine (1 x 30 mL), dried over anhydrous Na₂SO₄, and filtered. The filtrate was concentrated under reduced pressure, and the product was obtained (ethyl 3-fluoro-4-[[[(2S)-1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxylate (1.2914 g, 88%)) as an off-white solid. LC-MS (Column: HALO C18, 3.0*30mm, 2.0um; Column Oven: 40C; Mobile Phase A: Water/0.1% FA, Mobile Phase B: ACN/0.1% FA; Flow rate: 1.5mL/min; Gradient:5%B to 100%B in 1.2 min, hold 0.5 min; 254nm): Rt = 0.569 min). (ES, *m/z*): 321 [M+H]⁺, exact mass = 320.

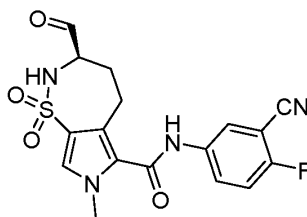
[0151] To a stirred solution of ethyl 3-fluoro-4-[[[(2S)-1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxylate (250.70 mg, 0.783 mmol, 1.00 eq.) in DMF(10mL) was added LiHMDS (1.0mo/L in THF, 2.40 mL, 2.400 mmol, 3.00 eq.) dropwise at rt. The mixture was stirred at 120 °C for 1 h. The mixture was cooled to rt and diluted with H₂O (10 mL), then extracted with EtOAc (3 x 10 mL). The combined organic layers were washed with brine (20mL), dried over anhydrous Na₂SO₄, and filtered. The filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography, eluted with PE/EtOAc(3:1) to afford ethyl (3S)-3-ethenyl-7-methyl-1,1-dioxo-2H,3H,4H-1λ6-pyrrolo[3,4-b][1,4,5]oxathiazepine-6-carboxylate (31.6 mg) as an off-white solid. LC-MS (Column: Ascentis Express C18, 3.0*50 mm, 2.7um;

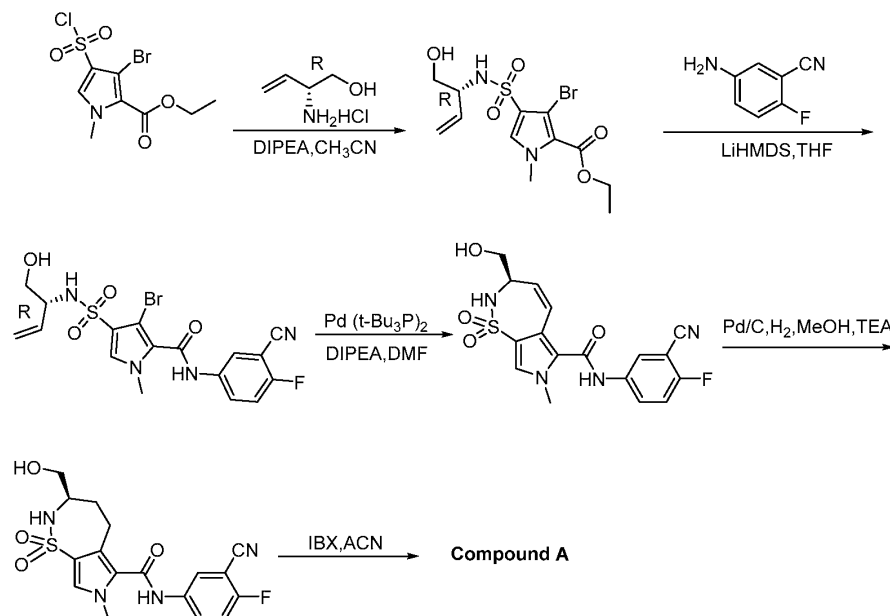
Column Oven: 40C; Mobile Phase A: Water/0.1% FA, Mobile Phase B: ACN/0.1% FA; Flow rate: 1.2 mL/min; Gradient: 10%B to 100%B in 2.0 min, hold 0.6 min; 254nm): Rt = 1.004 min). (ES, m/z): 301 $[M+H]^+$, exact mass = 300.

[0152] To a stirred solution of ethyl (3S)-3-ethenyl-7-methyl-1,1-dioxo-2H,3H,4H-1 λ 6-pyrrolo[3,4-b][1,4,5]oxathiazepine-6-carboxylate (31.60 mg, 0.105 mmol, 1.00 eq.) and 5-amino-2-fluorobenzonitrile (17.19 mg, 0.126 mmol, 1.20 eq.) in THF (3mL) were added LiHMDS (1.0 mol/L in THF, 0.65 mL, 6.00 eq.) dropwise at rt under N₂ atmosphere. The mixture was stirred at rt for 12h, diluted with H₂O (10mL) and extracted with EtOAc (3 x 10 mL). The combined organic layers were washed with brine (20 mL), dried over anhydrous Na₂SO₄. After filtration, the filtrate was concentrated under reduced pressure. The residue was purified by silica gel column chromatography, eluted with PE/EtOAc (1:1) to afford Compound **1** (20.2 mg) as an off-white solid. LC-MS (Column: ACE Excel 3 SuperC18, 3.0*50 mm, 3.0 μ m; Column Oven: 40C; Mobile Phase A: Water/5mM NH₄HCO₃, Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient: 10%B to 95%B in 2.1 min, hold 0.6 min; 254nm): Rt = 3.064 min). (ES, m/z): 391 $[M+H]^+$, exact mass = 390. ¹H-NMR: (DMSO-*d*₆, 300MHz, *ppm*): δ 9.56 (s, 1H), 8.20 (dd, J = 5.7, 2.6 Hz, 1H), 8.12-8.01 (m, 1H), 7.92 (d, J = 9.6 Hz, 1H), 7.59 – 7.49 (m, 2H), 5.93 – 5.78 (m, 1H), 5.42 (d, J = 17.4 Hz, 1H), 5.29 (d, J = 10.7 Hz, 1H), 4.69 (d, J = 12.7 Hz, 1H), 4.41-4.29 (m, 1H), 3.93 (dd, J = 12.7, 9.3 Hz, 1H), 3.85 (s, 3H).

Example 2

(3R)-N-(3-cyano-4-fluorophenyl)-3-formyl-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (Compound A)





[0153] To a solution of (2R)-2-aminobut-3-en-1-ol hydrochloride (1.46 g, 11.814 mmol, 1.30 eq.) in CH₃CN (30 mL) was added DIPEA (3.52 g, 27.235 mmol, 3.00 eq.), ethyl 3-bromo-4-(chlorosulfonyl)-1-methylpyrrole-2-carboxylate (3.00 g, 9.075 mmol, 1.00 eq.). The solution was stirred for 1 h at 80 °C. The mixture was concentrated. The residue was applied onto a silica gel column with EA/PE (40:60). Ethyl 3-bromo-4-[(2R)-1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxylate (3.1 g, 89.60%) was obtained as a white solid. LC-MS (Column: Ascentis Express C18, 3.0*50 mm, 2.7μm; Column Oven: 40C; Mobile Phase A: Water/0.1% FA, Mobile Phase B: ACN/0.1% FA; Flow rate: 1.2 mL/min; Gradient: 10%B to 100%B in 2.0 min, hold 0.6 min; 254nm; Rt = 1.295 min). (ES, *m/z*): 381 [M+H]⁺, exact mass = 380.0.

[0154] To a solution of ethyl 3-bromo-4-[(2R)-1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxylate (100 mg, 0.262 mmol, 1.00 eq.) in THF (3.00 mL) was added 5-amino-2-fluorobenzonitrile (53.56 mg, 0.393 mmol, 1.50 eq.) and LiHMDS (1 mol/L in THF, 12.59 mL, 12.590 mmol, 48.00 eq.). The solution was stirred for overnight at 25 °C. The reaction was then quenched with water (10 mL). The solution was extracted with EA (2 x 10 mL). The mixture was washed with brine (1 x 10 mL). The mixture was dried over anhydrous sodium sulfate and concentrated. The residue was applied onto a silica gel column with EA/PE (40:60). 3-bromo-N-(3-cyano-4-

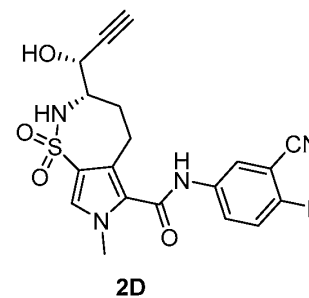
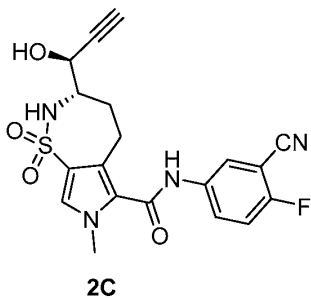
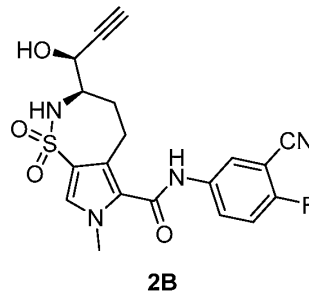
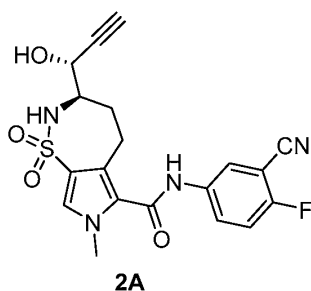
fluorophenyl)-4-[[$(2R)$ -1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxamide (66.8 mg, 52.95%) was obtained as a red solid. LC-MS (Column: HALO C18, 3.0*30mm, 2.0um; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.5mL/min; Gradient:5%B to 100%B in 1.2 min, hold 0.5 min; 254 nm; R_t = 1.408 min). (ES, m/z): 471 $[M+H]^+$, exact mass =470.0.

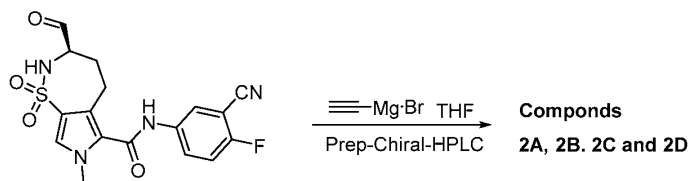
[0155] To a solution of 3-bromo-N-(3-cyano-4-fluorophenyl)-4-[[$(2R)$ -1-hydroxybut-3-en-2-yl]sulfamoyl]-1-methylpyrrole-2-carboxamide (0.20 g, 0.424 mmol, 1.00 eq.) in DMF (3.00 mL) was added DIEA (71.30 mg, 0.552 mmol, 1.30 eq.) and Pd(t-Bu₃P)₂ (43.37 mg, 0.085 mmol, 0.20 eq.). The mixture was irradiated with microwave radiation for 1 h at 130 °C. The solution was diluted with water (15 mL). The solution was extracted with EA (2 x 15 mL). The mixture was washed with brine (1 x15 mL). The mixture was dried over anhydrous sodium sulfate and concentrated. The residue was applied onto a silica gel column with CH₂Cl₂/CH₃OH (120:1). (3R)-N-(3-cyano-4-fluorophenyl)-3-(hydroxymethyl)-7-methyl-1,1-dioxo-2H,3H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (63.1 mg, 36.18%) was obtained as a red solid. LC-MS (Column: Shim-pack XR-ODS, 3.0*50 mm, 2.2um; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.2 mL/min; Gradient:5%B to 100%B in 2.0 min, hold 0.7 min; 254nm; R_t = 1.356 min). (ES, m/z):391 $[M+H]^+$, exact mass =390.1.

[0156] To a solution of (3R)-N-(3-cyano-4-fluorophenyl)-3-(hydroxymethyl)-7-methyl-1,1-dioxo-2H,3H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (200.00 mg, 0.512 mmol, 1.00 eq.) in MeOH (6 mL) was added TEA (103.68 mg, 1.025 mmol, 2.00 eq.), Pd/C (100.00 mg, 10%). The solution was stirred for 2 h at 25 °C under H₂ (1 atm) atmosphere. The filtrate was collected by filtration and concentrated. The crude product was slurry with ACN. The solid was collected by filtration. (3R)-N-(3-cyano-4-fluorophenyl)-3-(hydroxymethyl)-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (76.2 mg, 37.90%) of as an off-white solid. LC-MS (Column: Shim-pack XR-ODS, 3.0*50 mm,2.2 um; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.2 mL/min; Gradient:5%B to 100%B in 1.1 min, hold 0.7 min; 254 nm; R_t = 1.445 min). (ES, m/z):393 $[M+H]^+$, exact mass =392.1.

[0157] To a solution of (3R)-N-(3-cyano-4-fluorophenyl)-3-(hydroxymethyl)-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (400 mg, 1.019 mmol, 1.00 eq.) in ACN (40.00 mL) was followed by the addition of IBX (570.87 mg, 2.039 mmol, 2.00 eq.) at 80 °C. The solution was stirred for 40 min at 80 °C. The solids were filtered out, and the mixture was concentrated. The residue was applied onto a silica gel column with CH₂Cl₂/CH₃OH (110:1). Compound **A** (300 mg, 75.39%) was obtained as a yellow solid. LC-MS (Column: YMC-Triart C18, 3.0 μ m, 50*3.0 mm; Column Oven: 40C; Mobile Phase A: 0.04%NH₄OH, Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient:10%B to 95%B in 2.1 min, hold 0.6 min; 254nm; Rt = 0.826 min). (ES, *m/z*):391[M+H]⁺, exact mass =390.1 ¹H-NMR: ¹H NMR (400 MHz, DMSO-*d*₆) δ 10.60 (m, 1H), 9.54 (s, 1H), 8.20 (m, 1H), 7.98 - 7.94 (m, 1H), 7.57 - 7.54 (m, 2H), 5.82 (m, 1H), 4.74 - 4.22 (m, 1H), 3.69 (s, 3H), 3.11 - 3.01 (m, 1H), 2.87 - 2.67 (m, 1H), 2.09 (s, 1H), 1.56 - 1.32 (m, 1H).

Example 3
Compounds 2A, 2B, 2C and 2D





[0158] To a solution of (3R)-N-(3-cyano-4-fluorophenyl)-3-formyl-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (250 mg, 0.640 mmol, 1.00 eq.) in THF (6 mL) bromo(ethynyl)magnesium (0.5 mol/L in THF, 6.40 mL, 3.200 mmol, 5.00 eq.) added dropwise with stirring at 0 °C. The solution was stirred for 2 h at 0 °C. The reaction was then quenched with water (8mL). The solution was extracted with EA (2 x 15 mL), washed with brine (1 x 15 mL), dried over anhydrous sodium sulfate and concentrated. The residue was applied onto a silica gel column with CH₂Cl₂/CH₃OH (120:1). The crude product was purified by Prep-HPLC (conditions: 2#SHIMADZU (HPLC-01)): Column, XBridge Prep C18 OBD Column, 30*50mm, 5μm, 13nm; mobile phase, Water (10M MOL/L NH₄HCO₃) and ACN (30% Phase B up to 38% in 9 min)). This resulted in the racemic product which was purified by chiral-Prep-HPLC (conditions: Column: CHIRALPAK IG, 3*25cm, 5μm; Mobile Phase A:Hex (8 mmol/L NH₃•MeOH)-HPLC, Mobile Phase B:EtOH--HPLC; Flow rate:40 mL/min; Gradient:40 B to 40 B in 20 min; 254/220 nm; RT1:11.42; RT2:11.706. (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxyprop-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (1 mg, 0.36%, Compound **2A**) was obtained as a white solid; (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxyprop-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (7.8 mg, 2.90%, Compound **2D**) was obtained as a white solid. A mixture of two other product (9.6 mg) was also obtained.

[0159] The mixture of products (9.6 mg) was purified by Chiral-Prep-HPLC (conditions: Column: CHIRALPAK IA, 2*25cm, 5um; Mobile Phase A: MTBE(10mM NH₃•MeOH)-HPLC, Mobile Phase B: EtOH-HPLC; Flow rate:20 mL/min; Gradient:10 B to 10 B in 11 min; 220/254 nm; RT1:7.657; RT2:9.222. (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1S)-1-hydroxyprop-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (3.3 mg, 1.23%, Compound **2B**) was obtained as a white solid and (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1S)-1-hydroxyprop-2-yn-1-yl]-7-methyl-1,1-

dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (1 mg, 0.37%, Compound **2C**) was obtained as a white solid. The skilled in the art understand that Compounds **2A**, **2B**, **2C** and **2D** are diastereomers. The stereochemistry shown for each of Compounds **2A**, **2B**, **2C** and **2D** is relative and not absolute.

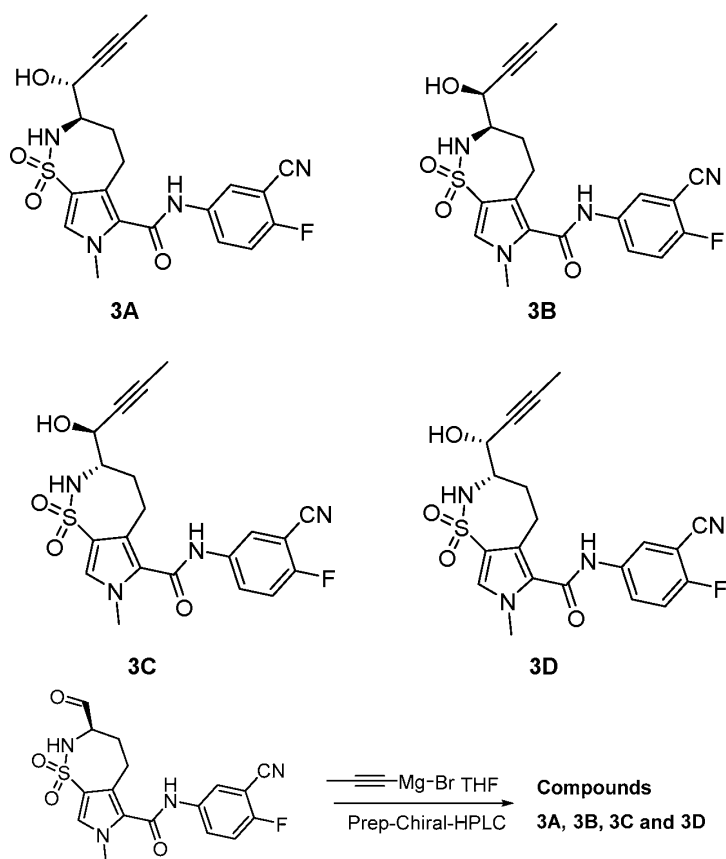
[0160] Compound **2A**: LC-MS (Column: HALO C18, 3.0*30mm, 2.0um; Column Oven:40 °C; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.5mL/min; Gradient:5%B to 100%B in 1.2 min, hold 0.5 min; 254nm; Rt = 1.110 min). (ES, m/z):417 [M+H]⁺, exact mass =416.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.61 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.92 (d, J = 10.2 Hz, 1H), 5.61 (d, J = 5.9 Hz, 1H), 4.25 (td, J = 5.6, 2.2 Hz, 1H), 3.70 (s, 3H), 3.54 (td, J = 10.0, 4.9 Hz, 1H), 3.34 (d, J = 2.2 Hz, 1H), 3.13-3.03 (m, 1H), 2.86-2.75 (m, 1H), 2.08 (q, J = 7.8, 6.5 Hz, 1H), 1.52 (q, J = 12.4 Hz, 1H).

[0161] Compound **2B**: LC-MS (Column: ACE Excel 3 SuperC18, 3.0*50 mm, 3.0 um; Column Oven: 40C; Mobile Phase A: Water/5mM N NH₄HCO₃, Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient:10%B to 95%B in 2.1 min, hold 0.6 min; 254nm; Rt = 1.458 min). (ES, m/z):417 [M+H]⁺, exact mass =416.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.61 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.46 (s, 1H), 7.06 (d, J = 10.2 Hz, 1H), 5.72 (d, J = 6.5 Hz, 1H), 4.13 (td, J = 6.7, 2.2 Hz, 1H), 3.70 (s, 3H), 3.49 (q, J = 9.9 Hz, 1H), 3.35 (d, J = 2.1 Hz, 1H), 3.09 (dd, J = 15.3, 6.6 Hz, 1H), 2.83-2.72 (m, 1H), 2.21 (dd, J = 14.3, 6.7 Hz, 1H), 1.39 (q, J = 12.3 Hz, 1H).

[0162] Compound **2C**: LC-MS (Column: ACE Excel 3 SuperC18, 3.0*50 mm, 3.0 um; Column Oven: 40C; Mobile Phase A: water/5mM NH₄HCO₃, Mobile Phase B:Acetonitrile; Flow rate: 1.2 mL/min; Gradient:10%B to 95%B in 2.1 min, hold 0.6 min; 254nm; Rt = 1.460 min). (ES, m/z):417 [M+H]⁺, exact mass =416.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.61 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.3, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.46 (s, 1H), 7.06 (d, J = 10.2 Hz, 1H), 5.72 (d, J = 6.5 Hz, 1H), 4.13 (td, J = 6.7, 2.2 Hz, 1H), 3.70 (s, 3H), 3.49 (q, J = 9.6 Hz, 1H), 3.36 (d, J = 2.1 Hz, 1H), 3.09 (dd, J = 15.1, 6.6 Hz, 1H), 2.78 (dd, J = 15.2, 12.3 Hz, 1H), 2.21 (dd, J = 14.3, 6.6 Hz, 1H), 1.39 (q, J = 12.4 Hz, 1H).

[0163] Compound **2D**: LC-MS (Column: ACE Excel 3 SuperC18, 3.0*50 mm, 3.0 μ m; Column Oven: 40°C; Mobile Phase A: water/5mM NH_4HCO_3 , Mobile Phase B: Acetonitrile; Flow rate: 1.2 mL/min; Gradient: 10%B to 95%B in 2.1 min, hold 0.6 min; 254nm; Rt = 1.453 min). (ES, m/z): 417 $[\text{M}+\text{H}]^+$, exact mass = 416.1. $^1\text{H-NMR}$: (400 MHz, $\text{DMSO-}d_6$) δ 10.61 (s, 1H), 8.20 (dd, J = 5.9, 2.7 Hz, 1H), 7.97 (ddd, J = 9.3, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.92 (d, J = 10.2 Hz, 1H), 5.62 (d, J = 6.0 Hz, 1H), 4.25 (td, J = 5.6, 2.2 Hz, 1H), 3.70 (s, 3H), 3.54 (td, J = 10.2, 5.3 Hz, 1H), 3.34 (d, J = 2.1 Hz, 1H), 3.13-3.03 (m, 1H), 2.86-2.75 (m, 1H), 2.07 (d, J = 5.0 Hz, 1H), 1.52 (q, J = 12.4 Hz, 1H).

Example 4
Compounds 3A, 3B, 3C and 3D



[0164] To a solution of (3R)-N-(3-cyano-4-fluorophenyl)-3-formyl-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (350 mg, 0.897 mmol, 1.00 eq.) in THF (7.00 mL) was added of bromo(prop-1-yn-1-yl)magnesium (8.97 mL, 4.485 mmol, 5.00 eq., 0.5 mol/L in THF) dropwise with stirring at 0 °C. The solution was stirred for 2 h at 0 °C. The reaction was quenched with water (10 mL). The solution was extracted with ethyl acetate (2 x 15 mL). The mixture was washed with brine (1 x 15 mL). The mixture was concentrated. The residue was applied onto a silica gel column with CH₂Cl₂/CH₃OH (120:1). The crude product was purified by Prep-HPLC (conditions: 2#SHIMADZU (HPLC-01)): Column, XBridge Prep OBD C18 Column, 30*150mm 5 μ m; mobile phase, Water (10M MOL/L NH₄HCO₃) and ACN (30% Phase B up to 52% in 9 min)). This resulted in racemic product was purified by Chiral-Prep-HPLC (conditions: (Prep-HPLC-009): Column, CHIRALPAK IE, 2*25cm, 5 μ m; mobile phase, Hex (8 mmol/L NH₃•MeOH) and EtOH- (hold 50% EtOH- in 19 min)). (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxybut-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (2 mg, 0.51%, Compound **3A**) was obtained as a solid, and (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxybut-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (2 mg, 0.51%, Compound **3D**) as a white solid. A mixture of two other product (27.3 mg) was also obtained.

[0165] The mixture of products (27.3 mg) was purified by Chiral-Prep-HPLC (conditions: Column: CHIRAL ART Cellulose-SB, 2*25cm, 5 μ m; Mobile Phase A:Hex (8 mmol/L NH₃•MeOH) HPLC, Mobile Phase B:EtOH--HPLC; Flow rate:20 mL/min; Gradient:35 B to 35 B in 16 min; 220/254 nm; RT1:10.984; RT2:12.642). This resulted in (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxybut-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (5.6 mg, 1.45%, Compound **3B**) as a white solid and (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1S)-1-hydroxybut-2-yn-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1 λ 6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (10.7 mg, 2.74%, Compound **3C**) as a white solid. The skilled in the art understand that Compounds **3A**, **3B**, **3C** and **3D** are diastereomers. The stereochemistry shown for each of Compounds **3A**, **3B**, **3C** and **3D** is relative and not absolute.

[0166] Compound **3A**: LC-MS (Column: Shim-pack XR-ODS,3.0*50 mm,2.2 μ m; Mobile Phase A:Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow

rate: 1.2 mL/min; Gradient: 5%B to 100%B in 2.0 min, hold 0.7 min; 254nm; Rt = 1.337 min, (ES, m/z): 431 [M+H]⁺, exact mass = 430.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.61 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.46 (s, 1H), 6.83 (d, J = 10.3 Hz, 1H), 5.44 (d, J = 5.6 Hz, 1H), 4.22 (dq, J = 5.6, 2.4 Hz, 1H), 3.70 (s, 3H), 3.50 (td, J = 10.8, 5.4 Hz, 1H), 3.08 (dd, J = 15.7, 6.3 Hz, 1H), 2.78 (dd, J = 15.3, 12.3 Hz, 1H), 2.15-2.05 (m, 1H), 1.79 (d, J = 2.1 Hz, 3H), 1.48 (m, 1H).

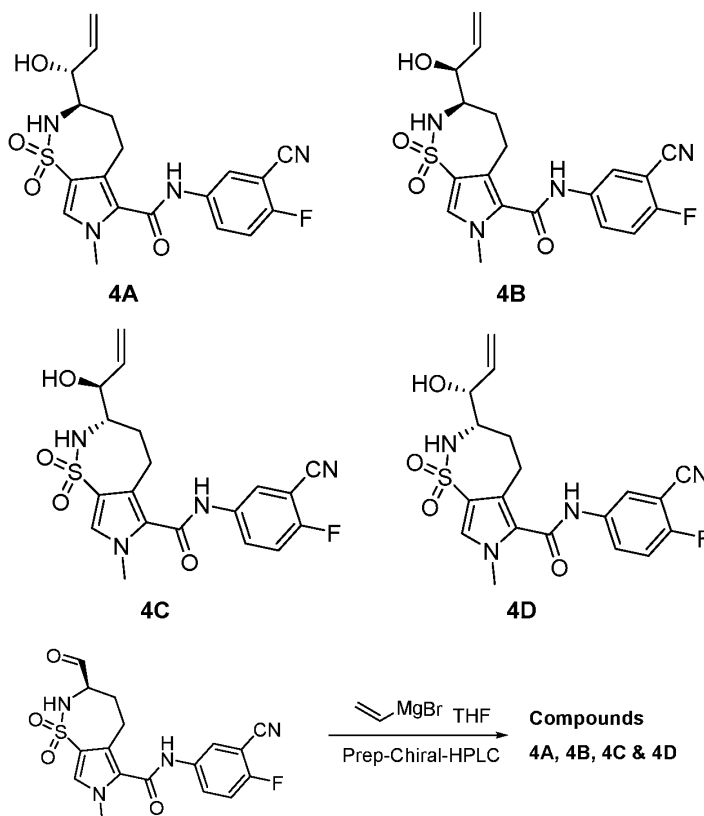
[0167] Compound **3B**: LC-MS (Column: Kinetex C18 100A 50*3.0mm, 2.6 μm; Column Oven: 40°C; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.2 mL/min; Gradient: 5%B to 100%B in 2.1min, hold 0.6 min; 254nm; Rt = 1.295 min, (ES, m/z): 431 [M+H]⁺, exact mass = 430.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.60 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.3, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.94 (d, J = 10.2 Hz, 1H), 5.48 (d, J = 6.3 Hz, 1H), 4.14 (ddt, J = 6.2, 4.1, 2.3 Hz, 1H), 3.70 (s, 3H), 3.51-3.40 (m, 1H), 3.08 (dd, J = 15.2, 6.6 Hz, 1H), 2.82-2.72 (m, 1H), 2.17 (dd, J = 14.4, 6.7 Hz, 1H), 1.82 (d, J = 2.1 Hz, 3H), 1.41 (m, 1H).

[0168] Compound **3C**: LC-MS (Column: Kinetex C18 100A 50*3.0mm, 2.6 μm; Column Oven: 40°C; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.2 mL/min; Gradient: 5%B to 100%B in 2.1min, hold 0.6 min; 254nm; Rt = 1.283 min, (ES, m/z): 431 [M+H]⁺, exact mass = 430.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.60 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.94 (d, J = 9.7 Hz, 1H), 5.48 (d, J = 6.2 Hz, 1H), 4.14 (ddq, J = 6.1, 4.1, 2.1 Hz, 1H), 3.70 (s, 3H), 3.46 (td, J = 10.3, 5.5 Hz, 1H), 3.13-3.03 (m, 1H), 2.78 (dd, J = 15.1, 12.4 Hz, 1H), 2.17 (dd, J = 14.3, 6.7 Hz, 1H), 1.82 (d, J = 2.1 Hz, 3H), 1.44-1.32 (m, 1H).

[0169] Compound **3D**: LC-MS (Column: Kinetex C18 100A 50*3.0mm, 2.6 μm; Column Oven: 40°C; Mobile Phase A: Water/0.05% TFA, Mobile Phase B: ACN/0.05% TFA; Flow rate: 1.2 mL/min; Gradient: 5%B to 100%B in 2.1min, hold 0.6 min; 254nm; Rt = 1.293 min, (ES, m/z): 431 [M+H]⁺, exact mass = 430.1. ¹H-NMR: (400 MHz, DMSO-*d*₆) δ 10.60 (s, 1H), 8.20 (dd, J = 5.8, 2.7 Hz, 1H), 7.97 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.94 (d, J = 9.7 Hz, 1H), 5.48 (d, J = 6.2 Hz, 1H), 4.14 (ddq, J = 6.1, 4.1, 2.1 Hz, 1H), 3.70 (s, 3H), 3.46 (td, J = 10.3, 5.5 Hz, 1H), 3.13-3.03 (m, 1H), 2.78 (dd, J

= 15.1, 12.4 Hz, 1H), 2.17 (dd, J = 14.3, 6.7 Hz, 1H), 1.82 (d, J = 2.1 Hz, 3H), 1.44-1.32 (m, 1H).

Example 5
Compounds 4A, 4B, 4C and 4D



[0170] To a solution of (3R)-N-(3-cyano-4-fluorophenyl)-3-formyl-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (430 mg, 1.101 mmol, 1.00 eq.) in THF (8.00 mL) was added bromo(ethenyl)magnesium (1 mol/L in THF, 5.51 mL, 5.5 mmol, 5.00 eq.) dropwise with stirring at 0 °C. The solution was stirred for 2 h at 0 °C. The reaction was then quenched with water (8 mL). The solution was extracted with ethyl acetate (2 x 15 mL). The mixture was washed with brine (1 x 15 mL). The mixture was dried over anhydrous sodium sulfate. The residue was applied onto a silica gel column with CH₂Cl₂/CH₃OH (120:1). The crude product was purified by Prep-HPLC (conditions: 2#SHIMADZU (HPLC-01)): Column, XBridge Prep OBD C18 Column, 30*150mm 5μm; mobile phase, Water (10M MOL/L NH₄HCO₃) and ACN (25% Phase B up to 50% in 9

min)). This resulted in racemic product was purified by Chiral-Prep-HPLC (conditions: (Prep-HPLC-009): Column: CHIRALPAK IC, 2*25cm,5um; Mobile Phase A: Hex (8 mmol/L NH₃•MeOH)-HPLC, Mobile Phase B: EtOH-HPLC; Flow rate: 20 mL/min; Gradient:30 B to 30 B in 18 min; 220/254 nm; RT1:4.178; RT2:11.194). (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxyprop-2-en-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (1.8 mg, 0.38%, Compound **4A**) was obtained as a white solid; (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1R)-1-hydroxyprop-2-en-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (8.6 mg, 1.83%, Compound **4D**) was obtained as a white solid. A mixture of two other product (13mg) was also obtained.

[0171] The mixture of products (13mg) was purified by Chiral-Prep-HPLC (conditions: Column: CHIRALPAK IE, 2*25cm, 5um; Mobile Phase A: Hex(8 mmol/L NH₃•MeOH)-HPLC, Mobile Phase B: EtOH--HPLC; Flow rate: 18 mL/min; Gradient:50 B to 50 B in 17 min; 220/254 nm; RT1:10.973; RT2:14.752). (3R)-N-(3-cyano-4-fluorophenyl)-3-[(1S)-1-hydroxyprop-2-en-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (1 mg, 0.21%, Compound **4B**) was obtained as a white solid; (3S)-N-(3-cyano-4-fluorophenyl)-3-[(1S)-1-hydroxyprop-2-en-1-yl]-7-methyl-1,1-dioxo-2H,3H,4H,5H-1λ6-pyrrolo[3,4-f][1,2]thiazepine-6-carboxamide (1 mg, 0.21%, Compound **4D**) was obtained as a white solid. The skilled in the art understand that Compounds 4A, 4B, 4C and 4D are diastereomers. The stereochemistry shown for each of Compounds 4A, 4B, 4C and 4D is relative and not absolute.

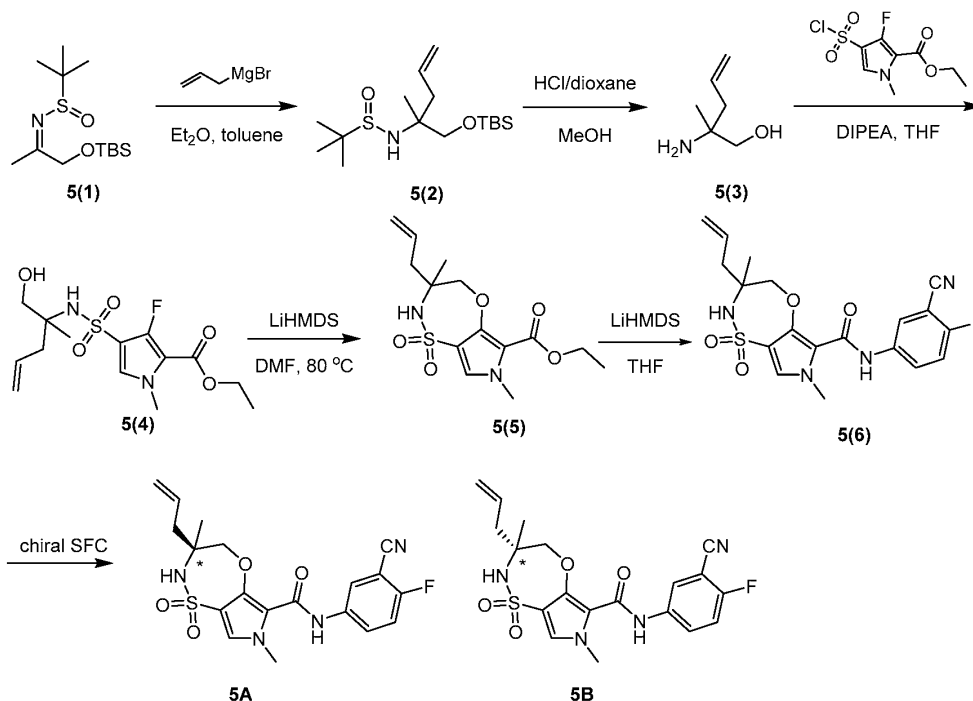
[0172] Compound **4A**: LC-MS (Column: ACE Excel 3 Super C18, 3.0*50 mm, 3.0 um; Column Oven: 40C; Mobile Phase A: Water/5mM NH₄HCO₃, Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient:10% B to 95% B in 2.1 min, hold 0.6 min; 254nm; Rt = 2.457 min). (ES, *m/z*):419 [M+H]⁺, exact mass =418.1. ¹H NMR (400 MHz, DMSO-*d*₆) δ 10.60 (s, 1H), 8.19 (dd, J = 5.8, 2.7 Hz, 1H), 7.96 (ddd, J = 9.1, 4.8, 2.7 Hz, 1H), 7.55 (t, J = 9.2 Hz, 1H), 7.45 (s, 1H), 6.95 (d, J = 10.3 Hz, 1H), 5.96 (ddd, J = 17.2, 10.5, 5.2 Hz, 1H), 5.24 (dt, J = 17.3, 1.8 Hz, 1H), 5.12 (dd, J = 10.4, 1.9 Hz, 1H), 5.04 (d, J = 5.7 Hz, 1H), 3.93-3.86 (m, 1H), 3.70 (s, 3H), 3.38 (t, J = 8.8 Hz, 1H), 3.06 (dd, J = 15.2, 6.5 Hz, 1H), 2.82-2.70 (m, 1H), 2.11 (dd, J = 14.2, 6.6 Hz, 1H), 1.38 (m, 1H).

[0173] Compound **4B**: LC-MS (Column: ACE Excel 3 Super C18, 3.0*50 mm, 3.0 μ m; Column Oven: 40C; Mobile Phase A: Water/5mM NH_4HCO_3 , Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient: 10% B to 95% B in 2.1 min, hold 0.6 min; 254nm; Rt = 1.307 min). (ES, m/z): 419 $[\text{M}+\text{H}]^+$, exact mass = 418.1. ^1H NMR (400 MHz, DMSO- d_6) δ 10.60 (s, 1H), 8.19 (dd, J = 5.8, 2.7 Hz, 1H), 7.96 (ddd, J = 9.2, 4.8, 2.6 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.95 (d, J = 10.3 Hz, 1H), 5.96 (ddd, J = 17.3, 10.5, 5.2 Hz, 1H), 5.24 (dt, J = 17.2, 1.9 Hz, 1H), 5.12 (dt, J = 10.5, 1.8 Hz, 1H), 5.04 (d, J = 5.6 Hz, 1H), 3.90 (q, J = 5.9 Hz, 1H), 3.70 (s, 3H), 3.43-3.33 (m, 1H), 3.06 (dd, J = 15.0, 6.7 Hz, 1H), 2.82-2.70 (m, 1H), 2.11 (dd, J = 15.1, 7.2 Hz, 1H), 1.35 (d, J = 12.8 Hz, 1H).

[0174] Compound **4C**: LC-MS (Column: ACE Excel 3 Super C18, 3.0*50 mm, 3.0 μ m; Column Oven: 40C; Mobile Phase A: Water/5 mM NH_4HCO_3 , Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient: 10% B to 95% B in 2.1 min, hold 0.6 min; 254nm; Rt = 2.450 min). (ES, m/z): 419 $[\text{M}+\text{H}]^+$, exact mass = 418.1. ^1H NMR (400 MHz, DMSO- d_6) δ 10.60 (s, 1H), 8.19 (dd, J = 5.7, 2.7 Hz, 1H), 7.96 (ddd, J = 9.2, 4.9, 2.7 Hz, 1H), 7.55 (t, J = 9.1 Hz, 1H), 7.45 (s, 1H), 6.78 (d, J = 10.4 Hz, 1H), 5.92 (ddd, J = 17.1, 10.5, 5.0 Hz, 1H), 5.24 (dt, J = 17.2, 1.9 Hz, 1H), 5.10 (dt, J = 10.4, 1.9 Hz, 1H), 5.01 (d, J = 5.2 Hz, 1H), 4.04 (d, J = 4.9 Hz, 1H), 3.69 (s, 3H), 3.55 (t, J = 11.8 Hz, 1H), 3.05 (dd, J = 15.1, 6.4 Hz, 1H), 2.84-2.72 (m, 1H), 1.90 (dd, J = 14.2, 6.6 Hz, 1H), 1.43 (d, J = 12.4 Hz, 1H).

[0175] Compound **4D**: LC-MS (Column: ACE Excel 3 Super C18, 3.0*50 mm, 3.0 μ m; Column Oven: 40C; Mobile Phase A: Water/5 mM NH_4HCO_3 , Mobile Phase B: ACN; Flow rate: 1.2 mL/min; Gradient: 10% B to 95% B in 2.1 min, hold 0.6 min; 254nm; Rt = 1.307 min). (ES, m/z): 419 $[\text{M}+\text{H}]^+$, exact mass = 418.1. ^1H NMR (400 MHz, DMSO- d_6) δ 10.60 (s, 1H), 8.19 (dd, J = 5.8, 2.7 Hz, 1H), 7.96 (ddd, J = 9.2, 4.8, 2.7 Hz, 1H), 7.55 (t, J = 9.2 Hz, 1H), 7.45 (s, 1H), 6.79 (d, J = 10.4 Hz, 1H), 5.92 (ddd, J = 17.2, 10.5, 5.0 Hz, 1H), 5.24 (dt, J = 17.2, 1.9 Hz, 1H), 5.10 (dt, J = 10.6, 1.8 Hz, 1H), 5.01 (d, J = 5.2 Hz, 1H), 4.04 (q, J = 4.9 Hz, 1H), 3.69 (s, 3H), 3.55 (ddd, J = 14.1, 10.2, 3.9 Hz, 1H), 3.10-3.00 (m, 1H), 2.78 (t, J = 13.7 Hz, 1H), 1.90 (dd, J = 14.2, 6.4 Hz, 1H), 1.40 (d, J = 12.8 Hz, 1H).

Example 6
Compounds 5A and 5B



[0176] To a solution of **5(1)** (4 g, 13.72 mmol, 1 eq.) in toluene (40 mL) was added dropwise allyl(bromo)magnesium (1 M in Et₂O, 34.30 mL, 2.5 eq.) at -78 °C under N₂. The mixture was stirred at -78 °C for 2 h. The reaction was quenched with sat. NH₄Cl (40 mL), then was allowed to warm to 15 °C slowly and extracted with EtOAc (30 mL x 3). The combined organic layers were concentrated under reduced pressure to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 40 g SepaFlash® Silica Flash Column, Eluent of 0~30% Ethyl acetate/Petroleum ether gradient @ 40 mL/min) to give **5(2)** (0.94 g, 2.82 mmol, 20.54%) as a colorless oil.

[0177] To a solution of **5(2)** (3.39 g, 10.16 mmol) in MeOH (30 mL) was added HCl/dioxane (4 M, 17.84 mL, 20 eq.) at 15 °C. The mixture was stirred at 15 °C for 1 h. The mixture was concentrated under reduced pressure to give crude **5(3)** (1.4 g, HCl salt) as a brown oil, which was used into the next step without further purification.

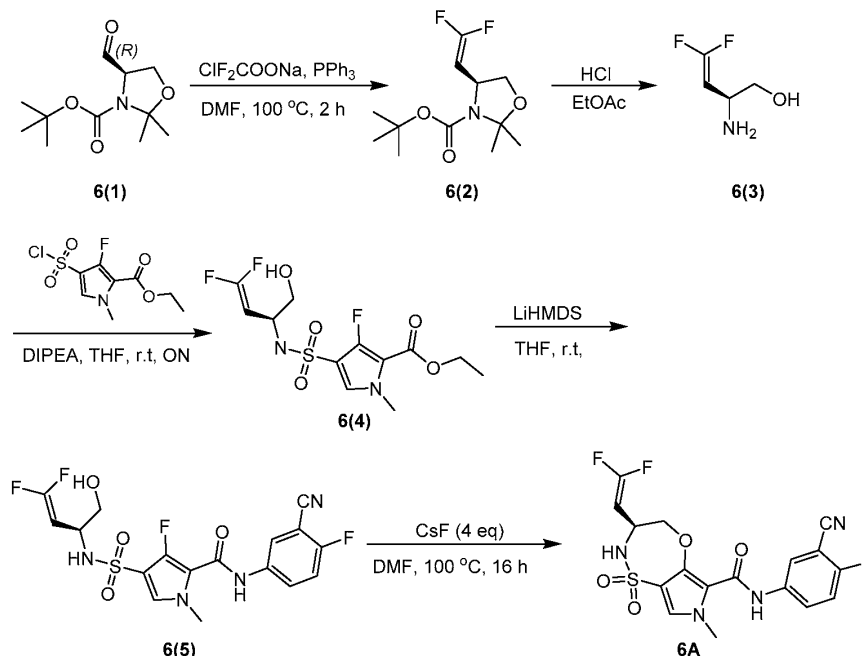
[0178] To a solution of crude **5(3)** (450 mg, 2.97 mmol, 1.5 eq.) in THF (25 mL) was added DIPEA (1.28 g, 9.89 mmol, 1.72 mL, 5 eq.) at 10 °C. The mixture was stirred at 10 °C for 1 h. Sulfonyl chloride (533.54 mg, 1.98 mmol, 1 eq.) was added to the mixture,

which was then stirred at 10 °C for 15 h. The mixture was concentrated under reduced pressure to give a residue, which was purified by flash silica gel chromatography (ISCO®; 12 g SepaFlash® Silica Flash Column, Eluent of 0~45% Ethyl acetate/Petroleum ether gradient @ 20 mL/min) to give **5(4)** (150 mg, 430.55 µmol, 21.76%) as a brown oil.

[0179] To a solution of **5(4)** (150 mg, 430.55 µmol, 1 eq.) in DMF (3 mL) was added dropwise LiHMDS (1 M, 1.5 mL, 3.48 eq.) at 10 °C. The mixture was stirred at 80 °C for 5 h. The reaction was quenched by the addition of sat. aq. NH₄Cl (15 mL) at 10 °C. The mixture was then diluted with H₂O (15 mL) and extracted with EtOAc (3 x 15 mL). The combined organic layers were washed with brine (30 mL), dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 0~25% Ethyl acetate/Petroleum ether gradient @ 20 mL/min) to give **5(5)** (130 mg, 307.36 µmol, 71.39% yield, 77.64% purity) as a yellow oil. The enantiomers were separated by chiral HPLC.

[0180] To a solution of **5(5)** (130 mg, 395.88 µmol, 1 eq.) and 5-amino-2-fluorobenzonitrile (75.45 mg, 554.23 µmol, 1.4 eq.) in THF (8 mL) was added LiHMDS (1 M, 1.58 mL, 4 eq.) at 10 °C. After the addition, the mixture was stirred at 10 °C for 2 h. TLC (Petroleum ether: Ethyl acetate (2:1)) indicated the starting material was consumed completely. The reaction was quenched with sat. aq. NH₄Cl (15 mL) and H₂O (10 mL), and then extracted with EtOAc (3 x 20 mL). The combined organic layers were washed with brine (40 mL), dried over anhydrous Na₂SO₄, filtered and concentrated to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 0~35% Ethyl acetate/Petroleum ether gradient @ 20 mL/min) to give **5(6)** (100 mg, 223.57 µmol, 56.47% yield, 93.55% purity) as a brown solid. The enantiomers, Compound **5A** and Compound **5B**, were separated by chiral HPLC. The stereochemistry shown for each of Compounds **5A** and **5B** is relative and not absolute.

Example 7
Compound 6A



[0181] To a solution of **6(1)** (0.50 g, 2.18 mmol, 1 eq.) and PPh_3 (1.72 g, 6.54 mmol, 3 eq.) in DMF (5 mL) was added sodium 2-chloro-2,2-difluoro-acetate (997.46 mg, 6.54 mmol, 3.0 eq.) at $100\text{ }^\circ\text{C}$ under N_2 . The mixture was stirred at $100\text{ }^\circ\text{C}$ for 2 h. The mixture was diluted with H_2O (20 mL) and extracted with EtOAc (3 x 25 mL). The combined organic layers were washed with brine (25 mL), dried over anhydrous Na_2SO_4 , filtered and concentrated under reduced pressure to give a residue, which was purified by flash silica gel chromatography (ISCO®; 20 g SepaFlash® Silica Flash Column, Eluent of 0~3% Ethyl acetate/Petroleum ether gradient) to give **5(2)** (0.53 g, 2.01 mmol, 92.31% yield) as a colorless oil.

[0182] A solution of **6(2)** (0.530 g, 2.01 mmol, 1 eq.) in HCl/EtOAc (4 M, 1.8 mL, 3.58 eq.) was stirred at $15\text{ }^\circ\text{C}$ for 0.5 h. The mixture was concentrated to dryness to give crude **6(3)** (0.32 g, 2.01 mmol, 99.62% yield, HCl salt) as a light yellow solid, which was used directly for next step without purification.

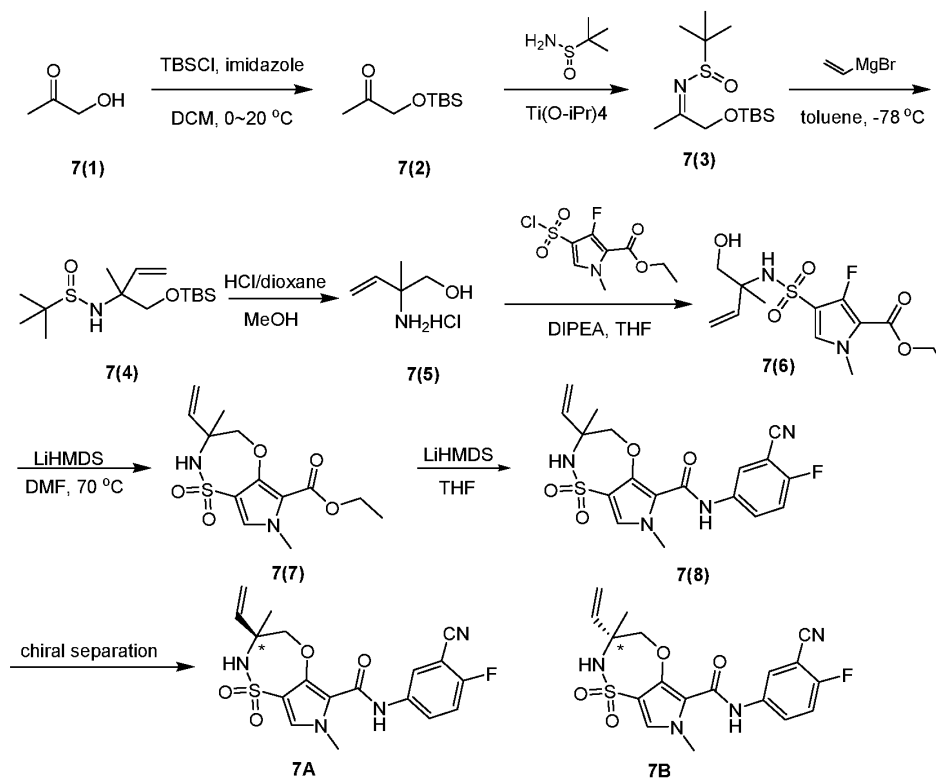
[0183] A mixture of **6(3)** (0.250 g, 2.03 mmol, 1.22 eq.) and DIPEA (1.08 g, 8.34 mmol, 1.45 mL, 5 eq.) in THF (20 mL) was stirred at $15\text{ }^\circ\text{C}$ for 1 h. Then sulfonyl chloride

(0.450 g, 1.67 mmol, 1 eq.) was added. The mixture was stirred at 15 °C for 15 h. The mixture was concentrated to give a residue, which was purified by flash silica gel chromatography (ISCO®; 20g SepaFlash® Silica Flash Column, Eluent of 0~90% Ethyl acetate/Petroleum ether gradient) to give **6(4)** (0.48 g, 80.73%) as a yellow solid.

[0184] To a solution of **6(4)** (0.25 g, 701.62 µmol, 1 eq.), 5-amino-2-fluorobenzonitrile (143.26 mg, 1.05 mmol, 1.5 eq.) in THF (5 mL) was added LiHMDS (1 M in THF, 3.5 mL, 3.5 mmol, 4.99 eq.) dropwise at 15 °C. The mixture was stirred at 15 °C for 3 h. The reaction was quenched with sat. aq. NH₄Cl (10 mL) and H₂O (5 mL). The mixture was extracted with EtOAc (3 x 20 mL). The combined organic layers were washed with brine (20 mL), dried over anhydrous Na₂SO₄, filtered and concentrated to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 30~65% Ethyl acetate/Petroleum ether gradient) to give the crude product (310 mg) as a purple solid. The crude product (190 mg) was re-purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 10~50% Ethyl acetate/Petroleum ether gradient) to give **6(5)** (180 mg) as a purple solid.

[0185] To a solution of **6(5)** (0.11 g, 246.43 µmol, 1 eq.) in DMF (2 mL) was added CsF (149.73 mg, 985.72 µmol, 4 eq.). The mixture was stirred at 100 °C for 16 h. The mixture was diluted with H₂O (15 mL) and extracted with EtOAc (3 x 15 mL). The combined organic layers were washed with brine (2 x 15 mL), dried over anhydrous Na₂SO₄, filtered and concentrated to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 12 g SepaFlash® Silica Flash Column, Eluent of 0~100% Ethyl acetate/Petroleum ether gradient @ 25 mL/min) to give the crude product (35 mg) as a red gum, which was re-purified by prep-HPLC (Instrument: BK; Column: Xtimate C18 150*25mm*5µm; Condition: water(0.05% HCl)-ACN; Begin B: 35; End B: 65; Gradient Time(min): 11.5; 100%B Hold Time(min): 2; FlowRate (ml/min): 25) to give compound **6A** (1.4 mg) as a white solid.

Example 8
Compounds 7A and 7B



[0186] To a solution of 1-hydroxypropan-2-one (20 g, 269.98 mmol, 18.52 mL, 1 eq.) and imidazole (25.73 g, 377.98 mmol, 1.4 eq.) in DCM (200 mL) was added TBSCl (44.76 g, 296.98 mmol, 36.39 mL, 1.1 eq.) at 0 °C. The mixture was stirred at 20 °C for 12 h. The mixture was concentrated under reduced pressure to give a residue. The residue was diluted with H₂O (200 mL), extracted with EtOAc (3 x 150 mL). The combined organic layers were washed with brine (200 mL), dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure to give the crude **7(2)** (47 g, 249.55 mmol, 92.43% yield) as a yellow oil, which was used into the next step without further purification.

[0187] To a mixture of crude **(2)** (37 g, 196.45 mmol, 1 eq.) in THF (370 mL) was added tetraisopropoxytitanium (139.59 g, 491.14 mmol, 144.95 mL, 2 eq.) and 2-methylpropane-2-sulfonamide (23.81 g, 196.45 mmol, 1 eq.) at 15 °C. The mixture was stirred at 70 °C for 14 h. After cooling to 15 °C, the mixture was poured into brine (400 mL) with stirring. The resulting suspension was filtered by diatomite and washed with EtOAc (6

x 50 mL). The filtrate was concentrated to give a residue. To the residue was added H₂O (200 mL). The mixture was extracted with EtOAc (3 x 150 mL). The combined organic layers were washed with brine (200 mL), dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure to give the crude product, which was purified by flash silica gel chromatography (ISCO®; 220 g SepaFlash® Silica Flash Column, Eluent of 0~15% Ethyl acetate/Petroleum ether gradient @ 100 mL/min) to give **7(3)** (9 g, 30.87 mmol, 15.71% yield) as a yellow oil.

[0188] To a solution of **(3)** (5 g, 17.15 mmol, 1 eq.) in toluene (50 mL) was added bromo(vinyl)magnesium (1.6 M in 2-MeTHF, 32.16 mL, 3 eq.) at -78 °C under N₂. The mixture was stirred at -78 °C for 2 h. The reaction was quenched with sat. NH₄Cl (100 mL) at -78 °C. The mixture was allowed to warm to 15 °C slowly. The mixture was then filtered and extracted with EtOAc (3 x 50 mL). The combined organic layers were washed with brine (100 mL), dried over anhydrous Na₂SO₄, filtered and concentrated under reduced pressure to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 40 g SepaFlash® Silica Flash Column, Eluent of 0~20% Ethyl acetate/Petroleum ether gradient @ 40 mL/min) to give **7(4)** (1.5 g, 4.69 mmol, 27.37% yield) as a yellow oil.

[0189] To a solution of **7(4)** (1.5 g, 4.69 mmol, 1 eq.) in MeOH (15 mL) was added HCl/dioxane (4 M, 11.73 mL, 10 eq.) at 15 °C. The mixture was stirred at 15 °C for 1 h. The mixture was concentrated under reduced pressure to give a residue. The residue was diluted with MTBE (30 mL) and extracted with H₂O (3 x 60 mL). The combined aqueous layers were concentrated under reduced pressure to give **7(5)** (620 mg, 4.51 mmol, 95.99% yield, HCl salt) as a brown oil, which was used into the next step without further purification

[0190] To a solution of **7(5)** (400 mg, 2.91 mmol, 1.5 eq.) in THF (25 mL) was added DIPEA (1.25 g, 9.69 mmol, 1.69 mL, 5 eq.) at 10 °C. The mixture was stirred at 10 °C for 1 h. Then to the mixture was added sulfonyl chloride (522.60 mg, 1.94 mmol, 1 eq.). The mixture was stirred at 10 °C for 19 h. The mixture was concentrated under reduced pressure to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 12 g SepaFlash® Silica Flash Column, Eluent of 0~50% Ethyl acetate/Petroleum ether gradient @ 30 mL/min) to give **7(6)** (170 mg, 508.43 µmol, 26.24% yield) as a brown oil.

[0191] To a solution of **7(6)** (170 mg, 508.43 μ mol, 1 eq.) in DMF (4 mL) was added LiHMDS (1 M, 1.77 mL, 3.48 eq.) at 10 °C. The mixture was stirred at 80 °C for 6 h. TLC (Petroleum ether: Ethyl acetate (1:1)) indicated the starting material was consumed completely. The reaction was quenched with sat. aq. NH_4Cl (15 mL) at 10 °C. The mixture was then diluted with H_2O (15 mL) and extracted with EtOAc (3 x 15 mL). The combined organic layers were washed with brine (30 mL), dried over anhydrous Na_2SO_4 , filtered and concentrated under reduced pressure to give a residue. The residue was purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 0~25% Ethyl acetate/Petroleum ether; gradient @ 20 mL/min) to give **7(7)** (130 mg, 370.37 μ mol, 72.85% yield, 89.56% purity) as a yellow oil.

[0192] To a solution of **7(7)** (130 mg, 413.54 μ mol, 1 eq.) and 5-amino-2-fluorobenzonitrile (78.81 mg, 578.96 μ mol, 1.4 eq.) in THF (8 mL) was added LiHMDS (1 M, 1.65 mL, 4 eq.) at 10 °C. The mixture was stirred at 10 °C for 2 h. TLC (Petroleum ether: Ethyl acetate (2:1)) indicated the starting material was consumed completely. The reaction was quenched with sat. aq. NH_4Cl (15 mL) and H_2O (10 mL). The mixture was then extracted with EtOAc (3 x 20 mL). The combined organic layers were washed with brine (40 mL), dried over anhydrous Na_2SO_4 , filtered and concentrated to give a residue, which was purified by flash silica gel chromatography (ISCO®; 4 g SepaFlash® Silica Flash Column, Eluent of 0~35% Ethyl acetate/Petroleum ether; gradient @ 20 mL/min) to give **7(8)** (100 mg, 58.62% yield, 98.03% purity) as a brown solid. The enantiomers, compound **7A** and compound **7B**, were further separated by chiral HPLC. The stereochemistry shown for each of Compounds **7A** and **7B** is relative and not absolute.

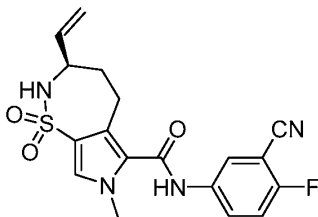
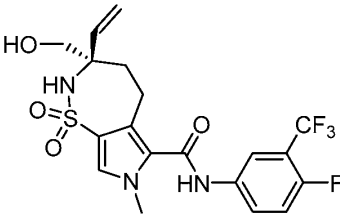
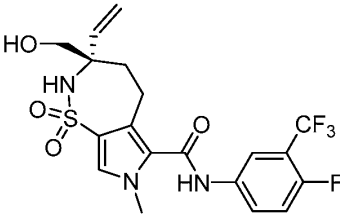
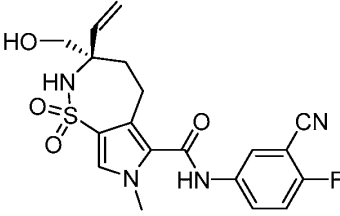
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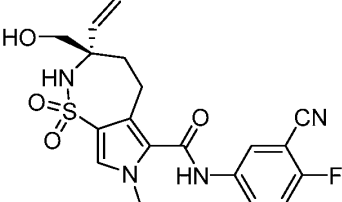
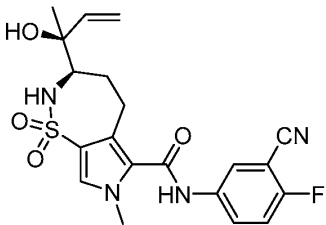
Additional Compounds

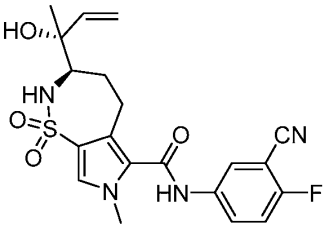
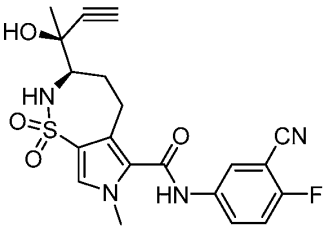
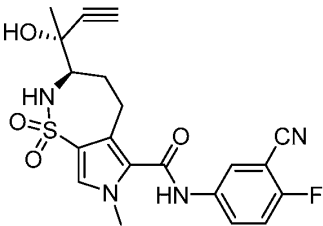
[0193] The foregoing syntheses are exemplary and can be used as a starting point to prepare a large number of additional compounds, including those provided in Table 1. Examples of compounds of Formulae (I) and (II) that can be prepared in various ways, including those synthetic schemes shown and described herein, are provided below. Those skilled in the art will be able to recognize modifications of the disclosed syntheses and to

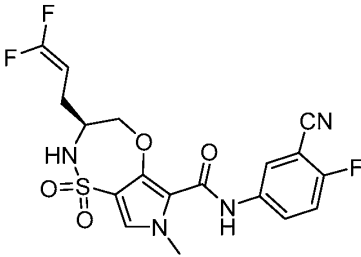
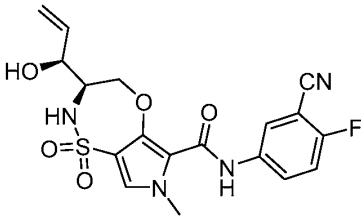
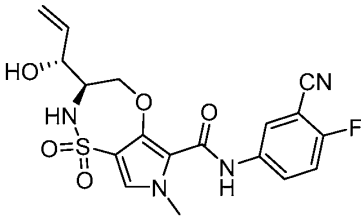
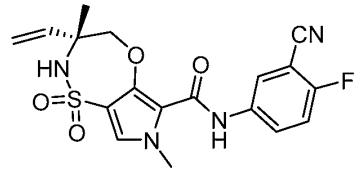
devise routes based on the disclosures herein; all such modifications and alternate routes are within the scope of the claims.

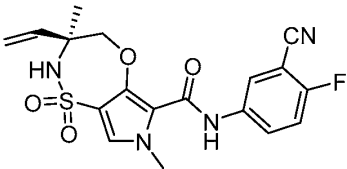
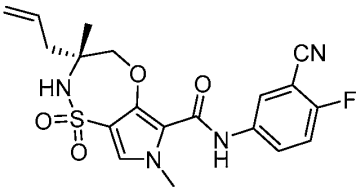
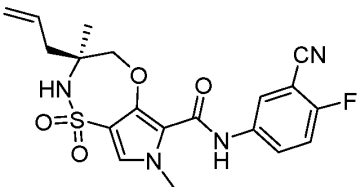
Table 1

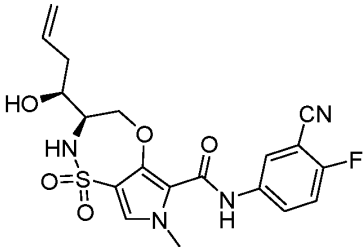
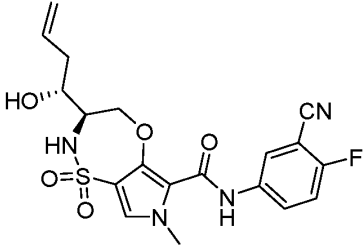
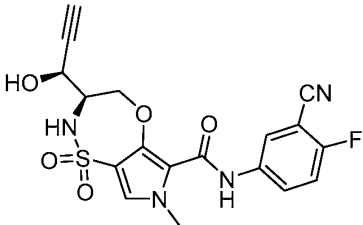
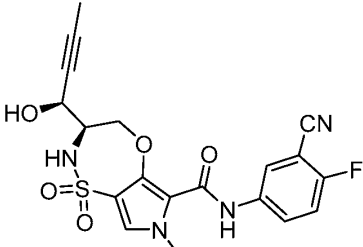
Compound No.	Structure	MS [M+1] ⁺	¹ H NMR
8		389.4	(400MHz, CDCl ₃) δ: 8.13 - 7.98 (m, 2H), 7.80 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.24 (t, J=8.7 Hz, 1H), 7.14 (s, 1H), 5.86 (ddd, J=4.9, 10.6, 17.3 Hz, 1H), 5.38 - 5.15 (m, 2H), 4.37 (br s, 1H), 4.12 (br d, J=8.4 Hz, 1H), 3.78 (s, 3H), 3.27 - 3.05 (m, 2H), 2.17 (br dd, J=6.5, 14.6 Hz, 1H), 1.55 (br d, J=3.4 Hz, 1H)
9A		462.3	(400MHz, CDOD ₃) δ: 8.07 (dd, J=2.7, 6.3 Hz, 1H), 7.93 - 7.82 (m, 1H), 7.38 - 7.27 (m, 2H), 6.02 (dd, J=11.1, 17.8 Hz, 1H), 5.33 - 5.22 (m, 2H), 3.77 (s, 3H), 3.63 - 3.55 (m, 1H), 3.54 - 3.43 (m, 1H), 3.20 - 3.00 (m, 2H), 2.30 - 2.12 (m, 2H)
9B		462.3	(400MHz, CDOD ₃) δ: 8.09 (dd, J=2.6, 6.3 Hz, 1H), 7.89 (td, J=3.8, 8.7 Hz, 1H), 7.41 - 7.28 (m, 2H), 6.04 (dd, J=11.2, 17.7 Hz, 1H), 5.35 - 5.21 (m, 2H), 3.79 (s, 3H), 3.64 - 3.58 (m, 1H), 3.56 - 3.48 (m, 1H), 3.20 - 3.00 (m, 2H), 2.33 - 2.14 (m, 2H)
10A		419.1	(400MHz, CDOD ₃) δ: 8.11 (dd, J=2.7, 5.7 Hz, 1H), 7.91 (ddd, J=2.8, 4.7, 9.2 Hz, 1H), 7.36 (t, J=9.0 Hz, 1H), 7.30 (s, 1H), 6.02 (dd, J=11.2, 17.7 Hz, 1H), 5.34 - 5.20 (m, 2H), 3.77 (s, 3H), 3.62 - 3.46 (m, 2H), 3.22 - 2.99 (m, 2H), 2.32

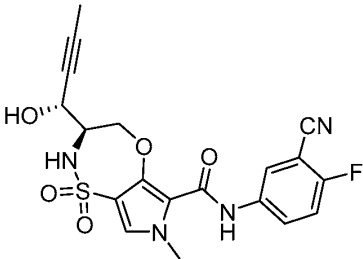
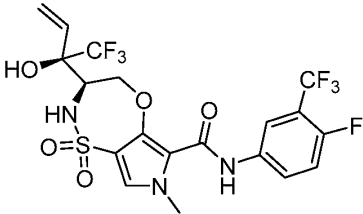
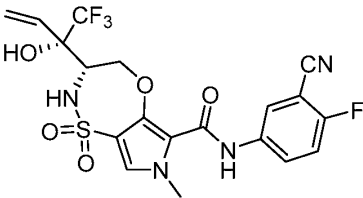
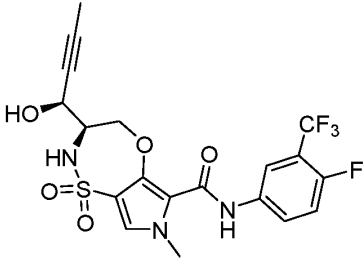
Compound No.	Structure	MS [M+1] +	¹ H NMR
			- 2.13 (m, 2H)
10B		419.1	(400MHz, CDCl ₃) δ: 8.07 - 7.96 (m, 1H), 7.85 (br s, 1H), 7.27 - 7.14 (m, 2H), 5.94 (br dd, J=11.7, 17.1 Hz, 1H), 5.40 - 5.20 (m, 2H), 5.08 (br s, 1H), 3.81 (s, 3H), 3.70 - 3.50 (m, 2H), 3.11 (br d, J=5.6 Hz, 2H), 2.32 (br s, 2H), 2.17 - 2.05 (m, 1H)
11A		433.4	(400MHz, DMSO-d ₆) δ: 10.60 (s, 1H), 8.18 (dd, J=2.4, 5.6 Hz, 1H), 8.02 - 7.87 (m, 1H), 7.54 (t, J=9.1 Hz, 1H), 7.46 (s, 1H), 6.92 (d, J=10.8 Hz, 1H), 5.92 (dd, J=10.7, 17.1 Hz, 1H), 5.20 (dd, J=1.9, 17.3 Hz, 1H), 5.06 - 4.95 (m, 1H), 4.76 (s, 1H), 3.68 (s, 3H), 3.03 (br dd, J=6.8, 15.1 Hz, 1H), 2.73 (br d, J=13.5 Hz, 1H), 2.13 - 2.04 (m, 1H), 1.26 - 1.09 (m, 5H)

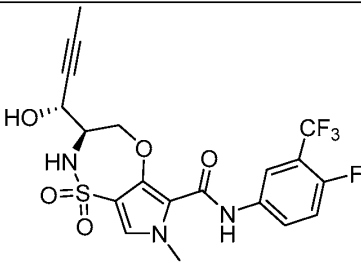
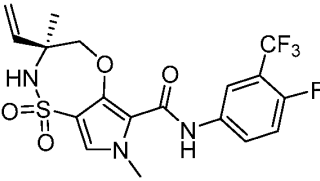
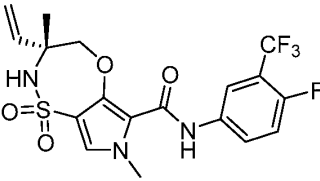
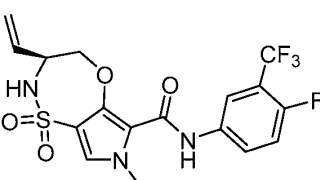
Compound No.	Structure	MS [M+1] +	¹ H NMR
11B		433.4	(400MHz, DMSO-d ₆) δ: 10.59 (s, 1H), 8.18 (dd, J=2.6, 5.8 Hz, 1H), 7.95 (ddd, J=2.7, 4.9, 9.2 Hz, 1H), 7.53 (t, J=9.1 Hz, 1H), 7.45 (s, 1H), 6.81 (d, J=10.6 Hz, 1H), 6.00 (dd, J=10.7, 17.3 Hz, 1H), 5.22 (dd, J=1.9, 17.3 Hz, 1H), 5.02 (dd, J=2.0, 10.6 Hz, 1H), 4.62 (s, 1H), 3.69 (s, 3H), 3.03 (br dd, J=6.6, 14.6 Hz, 1H), 2.79 - 2.65 (m, 1H), 2.07 (s, 1H), 1.47 - 1.32 (m, 1H), 1.15 (s, 3H)
12A		431.1	(400MHz, DMSO-d ₆) δ: 10.61 (s, 1H), 8.19 (dd, J=2.7, 5.8 Hz, 1H), 7.96 (ddd, J=2.7, 4.8, 9.1 Hz, 1H), 7.54 (t, J=9.1 Hz, 1H), 7.46 (s, 1H), 6.90 (d, J=10.8 Hz, 1H), 5.53 (s, 1H), 3.69 (s, 3H), 3.40 - 3.36 (m, 1H), 3.28 (s, 1H), 3.09 (br dd, J=6.9, 13.8 Hz, 1H), 2.80 - 2.70 (m, 1H), 2.25 (br dd, J=6.9, 13.8 Hz, 1H), 1.49 (q, J=12.0 Hz, 1H), 1.41 (s, 3H)
12B		431.1	(400MHz, DMSO-d ₆) δ: 10.61 (br s, 1H), 8.19 (dd, J=2.6, 5.8 Hz, 1H), 8.00 - 7.92 (m, 1H), 7.54 (t, J=9.1 Hz, 1H), 7.45 (s, 1H), 6.93 (br s, 1H), 5.52 (br s, 1H), 3.69 (s, 3H), 3.51 - 3.41 (m, 1H), 3.34 (br s, 1H), 3.10 (br dd, J=6.6, 14.8 Hz, 1H), 2.82 - 2.65 (m, 2H), 2.25 (br dd, J=6.6, 13.9 Hz, 1H), 1.48 (q, J=11.6 Hz, 1H), 1.35 (s, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
13		441.0	(400MHz, CDCl ₃) δ: 8.78 (s, 1H), 7.96 (dd, J=2.7, 5.3 Hz, 1H), 7.72 (ddd, J=2.9, 4.4, 8.9 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.09 (s, 1H), 4.77 - 4.64 (m, 2H), 4.43 - 4.18 (m, 2H), 3.96 (s, 4H), 2.40 (br t, J=7.5 Hz, 2H)
14A		421.2	(400MHz, CD ₃ OD) δ: 8.17 (dd, J=2.7, 5.6 Hz, 1H), 7.91 (ddd, J=2.8, 4.7, 8.8 Hz, 1H), 7.36 (t, J=8.9 Hz, 1H), 7.30 (s, 1H), 6.14 - 6.00 (m, 1H), 5.42 (d, J=17.1 Hz, 1H), 5.30 (d, J=10.5 Hz, 1H), 4.95 (d, J=2.0 Hz, 1H), 4.26 (t, J=6.0 Hz, 1H), 4.18 (dd, J=8.7, 12.8 Hz, 1H), 3.94 (s, 3H), 3.83 - 3.75 (m, 1H)
14B		421.1	(400MHz, DMSO-d ₆) δ: 9.54 (s, 1H), 8.20 (dd, J=2.6, 5.6 Hz, 1H), 8.05 (ddd, J=2.8, 4.8, 9.0 Hz, 1H), 7.68 - 7.36 (m, 3H), 6.03 - 5.85 (m, 1H), 5.57 - 5.07 (m, 3H), 4.72 (br d, J=12.3 Hz, 1H), 4.25 (br s, 1H), 3.97 - 3.70 (m, 5H)
15A		405.0	(400MHz, CDCl ₃) δ: 8.73 (s, 1H), 7.90 (dd, J=2.7, 5.4 Hz, 1H), 7.76 (ddd, J=2.8, 4.5, 9.0 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.06 (s, 1H), 6.10 (dd, J=10.8, 17.4 Hz, 1H), 5.48 (d, J=17.4 Hz, 1H), 5.40 (d, J=10.8 Hz, 1H), 4.87 (s, 1H), 4.78 - 4.72 (m, 1H), 4.68 - 4.61 (m, 1H), 3.96 (s, 3H), 2.02 (s, 1H), 1.59 (s, 3H)

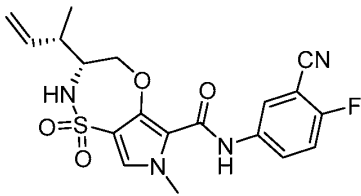
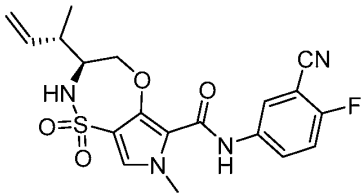
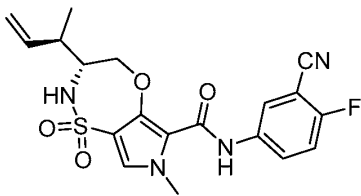
Compound No.	Structure	MS [M+1] +	¹ H NMR
15B		405.0	(400MHz, CDCl ₃) δ: 8.73 (s, 1H), 7.90 (dd, J=2.8, 5.4 Hz, 1H), 7.80 - 7.73 (m, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.06 (s, 1H), 6.10 (dd, J=10.8, 17.4 Hz, 1H), 5.48 (d, J=17.3 Hz, 1H), 5.40 (d, J=10.8 Hz, 1H), 4.88 (s, 1H), 4.78 - 4.72 (m, 1H), 4.67-4.61 (m, 1H), 3.96 (s, 3H), 2.02 (s, 1H), 1.59 (s, 3H)
16A		419.0	(400MHz, CDCl ₃) δ: 8.76 (s, 1H), 7.91 (dd, J=2.8, 5.4 Hz, 1H), 7.75 (ddd, J=2.8, 4.6, 9.0 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.05 (s, 1H), 5.99-5.85 (m, 1H), 5.35-5.23 (m, 2H), 4.75 (s, 1H), 4.70-4.58 (m, 2H), 3.96 (s, 3H), 2.67 (dd, J=7.4, 13.8 Hz, 1H), 2.41 (dd, J=7.5, 13.8 Hz, 1H), 1.45 (s, 3H)
16B		419.0	(400MHz, CDCl ₃) δ: 8.77 (s, 1H), 7.91 (dd, J=2.8, 5.4 Hz, 1H), 7.75 (ddd, J=2.9, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.05 (s, 1H), 5.98-5.84 (m, 1H), 5.35-5.24 (m, 2H), 4.75 (s, 1H), 4.70-4.58 (m, 2H), 3.96 (s, 3H), 2.67 (dd, J=7.3, 13.4 Hz, 1H), 2.41 (dd, J=7.8, 13.6 Hz, 1H), 2.02 (s, 1H), 1.45 (s, 3H)

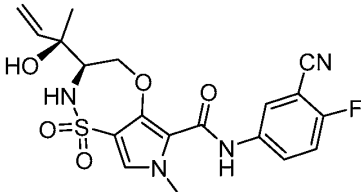
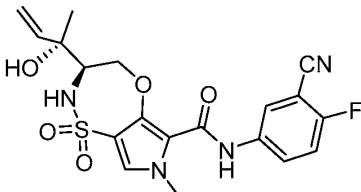
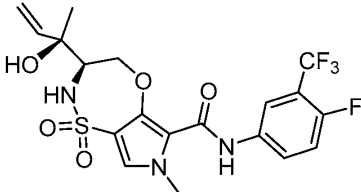
Compound No.	Structure	MS [M+1] +	¹ H NMR
17A		435.5	(400MHz, CDCl ₃) δ: 8.87 (s, 1H), 7.96 (dd, J=2.6, 5.4 Hz, 1H), 7.74 (ddd, J=2.9, 4.4, 9.0 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.09 (s, 1H), 5.85 (br d, J=8.0 Hz, 1H), 5.35-5.19 (m, 2H), 5.05 (d, J=9.6 Hz, 1H), 4.89 (dd, J=2.2, 12.9 Hz, 1H), 4.35 (dd, J=9.1, 12.9 Hz, 1H), 4.02-3.83 (m, 5H), 2.59 (br s, 1H), 2.45-2.27 (m, 1H), 2.09 (d, J=3.6 Hz, 1H)
17B		435.5	(400MHz, CDCl ₃) δ: 8.91 (s, 1H), 7.95 (dd, J=2.7, 5.4 Hz, 1H), 7.75 (ddd, J=2.8, 4.6, 9.1 Hz, 1H), 7.21 (t, J=8.7 Hz, 1H), 7.12 (s, 1H), 5.95-5.71 (m, 1H), 5.39-5.21 (m, 2H), 5.01 (d, J=10.4 Hz, 1H), 4.72 (dd, J=1.6, 12.8 Hz, 1H), 4.29 (dd, J=9.2, 12.8 Hz, 1H), 4.08-3.88 (m, 5H), 2.63-2.44 (m, 2H), 2.09-1.94 (m, 1H)
18		419.2	(400MHz, CD ₃ OD) δ: 8.17 (dd, J=2.8, 5.6 Hz, 1H), 7.92 (ddd, J=2.7, 4.7, 9.1 Hz, 1H), 7.36 (t, J=8.9 Hz, 1H), 7.30 (s, 1H), 5.04 (dd, J=2.0, 12.8 Hz, 1H), 4.57 (dd, J=2.1, 5.7 Hz, 1H), 4.26 (dd, J=8.8, 12.8 Hz, 1H), 3.98-3.90 (m, 4H), 3.05 (d, J=2.1 Hz, 1H)
19A		433.0	(400MHz, DMSO-d ₆) δ: 9.54 (s, 1H), 8.21 (dd, J=2.8, 5.8 Hz, 1H), 8.05 (ddd, J=2.8, 5.0, 9.3 Hz, 1H), 7.79 (br d, J=4.3 Hz, 1H), 7.56-7.48 (m, 2H), 5.88 (d, J=6.0 Hz, 1H), 4.89 (dd, J=1.9, 12.7 Hz, 1H), 4.34-4.24 (m, 1H), 4.02 (dd, J=9.3, 12.8 Hz, 1H), 3.83 (s, 3H), 3.70-3.61 (m, 1H), 1.86 (d, J=2.0 Hz, 3H)

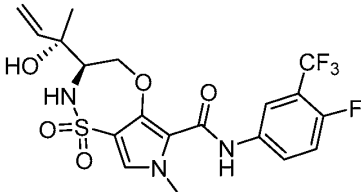
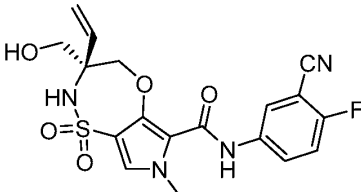
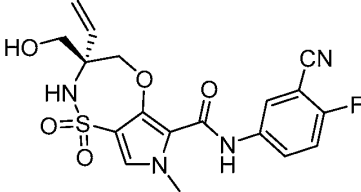
Compound No.	Structure	MS [M+1] +	¹ H NMR
19B		433.3	(400MHz, CDCl ₃) δ: 8.87 (s, 1H), 7.98 (dd, J=2.8, 5.4 Hz, 1H), 7.72 (ddd, J=2.7, 4.5, 9.0 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.11 (s, 1H), 4.95 (dd, J=2.1, 12.9 Hz, 2H), 4.78-4.65 (m, 2H), 4.30 (dd, J=8.6, 12.8 Hz, 1H), 4.11 (ddd, J=2.3, 5.6, 8.2 Hz, 1H), 3.97 (s, 3H), 2.35 (br s, 1H), 1.89 (d, J=2.1 Hz, 3H)
20		532.1	(400MHz, CDCl ₃) δ: 8.69 (s, 1H), 7.86 (dd, J=2.6, 6.1 Hz, 1H), 7.75-7.68 (m, 1H), 7.21 (t, J=9.2 Hz, 1H), 7.10 (s, 1H), 6.09 (dd, J=10.9, 17.1 Hz, 1H), 5.81 (d, J=17.1 Hz, 1H), 5.70 (d, J=10.9 Hz, 1H), 4.98-4.89 (m, 2H), 4.39 (dd, J=9.1, 12.9 Hz, 1H), 4.29 (br d, J=7.5 Hz, 1H), 3.98 (s, 3H), 3.40 (s, 1H)
21		489.2	¹ H NMR (400MHz, DMSO-d ₆) δ: 9.52 (s, 1H), 8.18 (dd, J=2.8, 5.8 Hz, 1H), 8.02 (ddd, J=2.8, 4.9, 9.1 Hz, 2H), 7.57-7.44 (m, 2H), 6.96 (s, 1H), 6.10 (dd, J=10.9, 17.0 Hz, 1H), 5.67 (dd, J=1.3, 17.0 Hz, 1H), 5.49 (d, J=11.9 Hz, 1H), 4.72 (d, J=11.8 Hz, 1H), 4.11 (br s, 1H), 3.82 (s, 3H), 3.74 (dd, J=8.9, 12.7 Hz, 1H)
22A		476.1	(400MHz, CDCl ₃) δ: 8.84 (s, 1H), 7.84 (dd, J=2.6, 6.1 Hz, 1H), 7.73 (td, J=3.5, 8.8 Hz, 1H), 5.04 (br s, 1H), 4.95 (dd, J=1.4, 12.5 Hz, 1H), 4.71 (br s, 1H), 4.27-4.16 (m, 1H), 4.16-4.08 (m, 1H), 3.98 (s, 3H), 2.33 (br s, 1H), 1.91 (d, J=2.1 Hz, 3H)

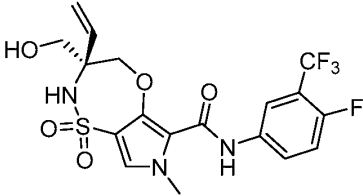
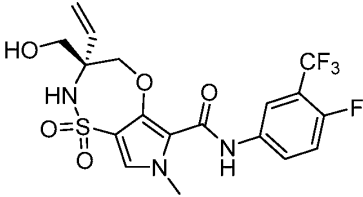
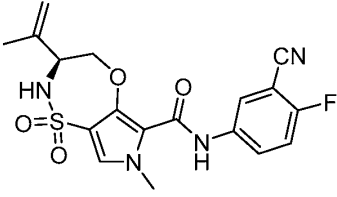
Compound No.	Structure	MS [M+1] +	¹ H NMR
22B		476.3	(400MHz, CDCl ₃) δ: 8.85 (s, 1H), 7.91-7.79 (m, 1H), 7.78-7.69 (m, 1H), 7.19 (t, J=9.3 Hz, 1H), 7.10 (s, 1H), 5.05-4.83 (m, 2H), 4.78-4.63 (m, 2H), 4.30 (dd, J=8.7, 12.8 Hz, 1H), 4.11 (br s, 1H), 3.98 (s, 3H), 1.89 (d, J=2.0 Hz, 3H)
23A		448.2	(400MHz, CDCl ₃) δ: 8.69 (s, 1H), 7.81-7.69 (m, 2H), 7.18 (t, J=9.3 Hz, 1H), 7.04 (s, 1H), 6.09 (dd, J=10.8, 17.3 Hz, 1H), 5.52-5.34 (m, 2H), 4.85 (s, 1H), 4.78-4.70 (m, 1H), 4.68-4.58 (m, 1H), 3.95 (s, 3H), 1.58 (s, 3H)
23B		448.2	(400MHz, CDCl ₃) δ: 8.69 (s, 1H), 7.82-7.68 (m, 2H), 7.18 (t, J=9.4 Hz, 1H), 7.04 (s, 1H), 6.09 (dd, J=10.8, 17.4 Hz, 1H), 5.54-5.33 (m, 2H), 4.86 (s, 1H), 4.79-4.58 (m, 2H), 3.95 (s, 3H), 1.58 (s, 3H)
24		434.2	(400MHz, CDCl ₃) δ: 8.81 (s, 1H), 7.83 (dd, J=2.6, 6.1 Hz, 1H), 7.79-7.71 (m, 1H), 7.28 (s, 2H), 7.21 (t, J=9.4 Hz, 1H), 7.10 (s, 1H), 6.01-5.86 (m, 1H), 5.58-5.36 (m, 2H), 4.83-4.69 (m, 2H), 4.61 (br d, J=8.3 Hz, 1H), 4.21 (dd, J=8.5, 12.9 Hz, 1H), 3.99 (s, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
25A		435.4	(400MHz, CDCl ₃) δ: 8.88 (br s, 1H), 7.98 (br d, J=2.0 Hz, 1H), 7.76 (br d, J=3.8 Hz, 1H), 7.25-7.18 (m, 1H), 7.11 (br s, 1H), 5.99-5.73 (m, 1H), 5.47-5.21 (m, 2H), 5.16-4.81 (m, 2H), 4.37 (br dd, J=9.4, 12.6 Hz, 1H), 4.15-3.79 (m, 5H), 2.63 (td, J=2.9, 5.6 Hz, 1H), 2.47-2.27 (m, 1H), 2.21-1.99 (m, 1H)
25B		435.4	(400MHz, CDCl ₃) δ: 8.90 (br s, 1H), 7.95 (br d, J=2.6 Hz, 1H), 7.81-7.67 (m, 1H), 7.21 (br t, J=8.7 Hz, 1H), 7.12 (s, 1H), 5.92-5.75 (m, 1H), 5.38-5.21 (m, 2H), 4.99 (br d, J=10.3 Hz, 1H), 4.72 (br d, J=13.0 Hz, 1H), 4.29 (br dd, J=9.5, 12.3 Hz, 1H), 4.08-3.85 (m, 5H), 2.63-2.43 (m, 2H), 2.00 (br s, 1H)
26		532.2	(400MHz, CDCl ₃) δ: 8.70 (s, 1H), 7.86 (dd, J=2.6, 6.1 Hz, 1H), 7.71 (td, J=3.5, 8.7 Hz, 1H), 7.20 (t, J=9.3 Hz, 1H), 7.10 (s, 1H), 6.08 (dd, J=10.9, 17.0 Hz, 1H), 5.81 (d, J=17.1 Hz, 1H), 5.70 (d, J=10.8 Hz, 1H), 4.94 (dd, J=2.4, 13.1 Hz, 2H), 4.46-4.35 (m, 1H), 4.27 (br d, J=7.8 Hz, 1H), 3.97 (s, 3H), 3.45 (br d, J=1.8 Hz, 1H)
27A		419.1	(400MHz, DMSO-d ₆) δ: 9.52 (s, 1H), 8.19 (dd, J=2.7, 5.7 Hz, 1H), 8.03 (ddd, J=2.8, 4.8, 9.2 Hz, 1H), 7.67 (br s, 1H), 7.56-7.48 (m, 2H), 5.90-5.76 (m, 1H), 5.18-5.04 (m, 2H), 4.70-4.59 (m, 1H), 3.92 (dd, J=9.4, 12.8 Hz, 1H), 3.83 (s, 3H), 3.59-3.50 (m, 1H), 1.08 (d, J=6.8 Hz, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
27B		419.1	(400MHz, DMSO-d ₆) δ: 9.54 (s, 1H), 8.19 (dd, J=2.4, 5.7 Hz, 1H), 8.05 (ddd, J=2.8, 5.0, 9.2 Hz, 1H), 7.63-7.57 (m, 1H), 7.56-7.50 (m, 2H), 5.92-5.79 (m, 1H), 5.15-5.00 (m, 2H), 4.65 (br d, J=12.1 Hz, 1H), 3.90-3.85 (m, 1H), 3.83 (s, 3H), 3.71-3.59 (m, 1H), 1.10-1.04 (m, 3H)
27C		419.1	(400MHz, DMSO-d ₆) δ: 9.54 (s, 1H), 8.19 (dd, J=2.6, 5.6 Hz, 1H), 8.05 (ddd, J=2.9, 4.9, 9.0 Hz, 1H), 7.66-7.42 (m, 3H), 5.93-5.77 (m, 1H), 5.18-5.00 (m, 2H), 4.65 (d, J=12.3 Hz, 1H), 3.90-3.85 (m, 1H), 3.83 (s, 3H), 3.70-3.62 (m, 1H), 1.07 (d, J=7.0 Hz, 3H)
27D		419.1	(400MHz, DMSO-d ₆) δ: 9.52 (s, 1H), 8.19 (dd, J=2.7, 5.7 Hz, 1H), 8.03 (ddd, J=2.6, 4.9, 9.2 Hz, 1H), 7.67 (d, J=9.5 Hz, 1H), 7.57-7.48 (m, 2H), 5.90-5.74 (m, 1H), 5.16-5.03 (m, 2H), 4.65 (dd, J=1.4, 12.8 Hz, 1H), 3.92 (dd, J=9.4, 12.6 Hz, 1H), 3.83 (s, 3H), 3.59-3.49 (m, 1H), 1.08 (d, J=6.8 Hz, 3H)

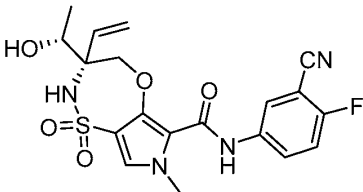
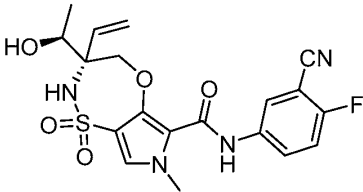
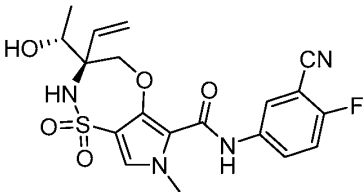
Compound No.	Structure	MS [M+1] +	¹ H NMR
28A		435.2	(400MHz, DMSO-d ₆) δ: 9.49 (s, 1H), 8.19 (dd, J=2.7, 5.7 Hz, 1H), 8.04 (ddd, J=2.7, 4.8, 9.1 Hz, 1H), 7.59 (br s, 1H), 7.50 (s, 2H), 5.94 (dd, J=10.8, 17.0 Hz, 1H), 5.30 (d, J=2.0 Hz, 1H), 5.33-5.22 (m, 1H), 5.04 (dd, J=2.0, 10.6 Hz, 1H), 4.84 (d, J=12.0 Hz, 1H), 3.82 (s, 3H), 3.72 (dd, J=9.1, 12.4 Hz, 1H), 3.64 (br s, 1H), 1.31 (s, 3H)
28B		435.3	(400MHz, DMSO-d ₆) δ: 9.51 (s, 1H), 8.19 (dd, J=2.6, 5.8 Hz, 1H), 8.02 (br s, 1H), 7.58-7.41 (m, 3H), 6.07 (dd, J=10.6, 17.3 Hz, 1H), 5.30 (dd, J=1.5, 17.3 Hz, 1H), 5.13 (dd, J=1.6, 10.7 Hz, 1H), 5.05 (s, 1H), 4.84 (d, J=12.0 Hz, 1H), 3.91 (br d, J=12.6 Hz, 1H), 3.82 (s, 3H), 3.65-3.55 (m, 1H), 3.31 (s, 3H), 1.20 (s, 3H)
29A		478.2	(400MHz, DMSO-d ₆) δ: 9.55 (s, 1H), 8.19 (dd, J=2.6, 6.3 Hz, 1H), 7.99-7.89 (m, 1H), 7.55-7.44 (m, 3H), 6.08 (dd, J=10.7, 17.2 Hz, 1H), 5.30 (dd, J=1.5, 17.3 Hz, 1H), 5.17-5.08 (m, 1H), 5.04 (s, 1H), 4.82 (d, J=12.4 Hz, 1H), 3.92 (dd, J=8.9, 12.4 Hz, 1H), 3.82 (s, 3H), 3.60 (br t, J=9.5 Hz, 1H), 1.20 (s, 3H)

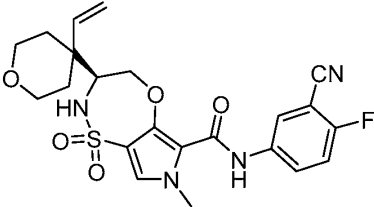
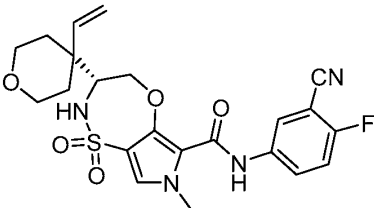
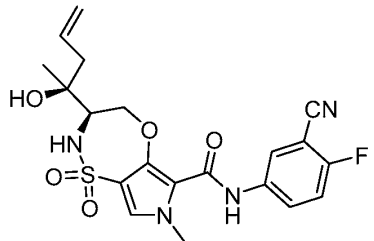
Compound No.	Structure	MS [M+1] +	¹ H NMR
29B		478.2	(400MHz, DMSO-d ₆) δ: 9.54 (s, 1H), 8.19 (dd, J=2.5, 6.5 Hz, 1H), 7.98-7.89 (m, 1H), 7.59 (br d, J=9.3 Hz, 1H), 7.53-7.42 (m, 2H), 5.94 (dd, J=10.6, 17.0 Hz, 1H), 5.35-5.21 (m, 2H), 5.04 (dd, J=2.1, 10.7 Hz, 1H), 4.81 (d, J=12.3 Hz, 1H), 3.82 (s, 3H), 3.72 (dd, J=9.1, 12.3 Hz, 1H), 3.67-3.59 (m, 1H), 2.07 (s, 3H), 1.31 (s, 3H)
30A		421.2	(400MHz, DMSO-d ₆) δ: 9.41 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.02 (ddd, J=2.7, 4.8, 9.2 Hz, 1H), 7.83 (br s, 1H), 7.54 (t, J=9.1 Hz, 1H), 7.47 (s, 1H), 5.89 (dd, J=11.0, 17.5 Hz, 1H), 5.47 (d, J=1.1 Hz, 1H), 5.42 (d, J=1.3 Hz, 1H), 5.27 (dd, J=1.1, 11.0 Hz, 1H), 5.16 (t, J=5.9 Hz, 1H), 4.76-4.65 (m, 2H), 3.81 (s, 3H), 3.76 (dd, J=5.9, 10.8 Hz, 1H), 3.40-3.37 (m, 1H), 3.33 (s, 1H), 3.31 (s, 16H)
30B		421.2	(400MHz, DMSO-d ₆) δ: 9.42 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.02 (ddd, J=2.7, 4.9, 9.2 Hz, 1H), 7.84 (br s, 1H), 7.54 (t, J=9.1 Hz, 1H), 7.47 (s, 1H), 5.89 (dd, J=11.0, 17.5 Hz, 1H), 5.45 (dd, J=1.1, 17.5 Hz, 1H), 5.27 (dd, J=1.2, 10.9 Hz, 1H), 5.17 (t, J=6.0 Hz, 1H), 4.78-4.62 (m, 2H), 3.81 (s, 3H), 3.76 (dd, J=6.0, 10.8 Hz, 1H), 3.39 (br d, J=6.3 Hz, 1H)

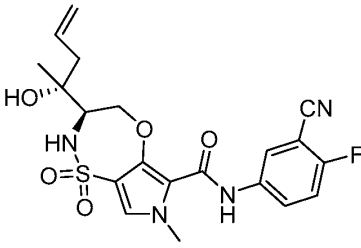
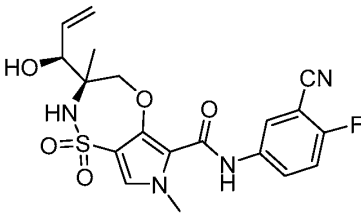
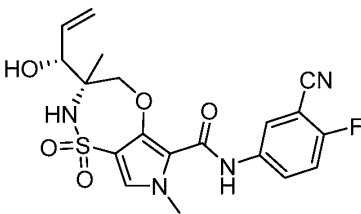
Compound No.	Structure	MS [M+1] +	¹ H NMR
31A		464.3	(400MHz, DMSO-d ₆) δ: 9.46 (s, 1H), 8.18 (dd, J=2.5, 6.4 Hz, 1H), 7.95-7.90 (m, 1H), 7.85 (br s, 1H), 7.52 (t, J=9.8 Hz, 1H), 7.47 (s, 1H), 5.90 (dd, J=10.9, 17.5 Hz, 1H), 5.45 (dd, J=0.9, 17.5 Hz, 1H), 5.28 (dd, J=0.9, 10.9 Hz, 1H), 5.17 (br t, J=5.9 Hz, 1H), 4.77-4.64 (m, 2H), 3.82 (s, 3H), 3.77 (br dd, J=5.9, 10.8 Hz, 1H), 3.39 (dd, J=5.5, 10.6 Hz, 1H)
31B		464.3	(400MHz, DMSO-d ₆) δ: 9.45 (s, 1H), 8.17 (dd, J=2.4, 6.4 Hz, 1H), 7.95-7.89 (m, 1H), 7.85 (s, 1H), 7.52 (t, J=9.8 Hz, 1H), 7.46 (s, 1H), 5.89 (dd, J=11.0, 17.5 Hz, 1H), 5.44 (d, J=17.5 Hz, 1H), 5.27 (d, J=10.9 Hz, 1H), 5.16 (t, J=5.9 Hz, 1H), 4.75-4.64 (m, 2H), 3.81 (s, 3H), 3.76 (dd, J=6.1, 10.7 Hz, 1H), 3.38 (dd, J=5.9, 10.7 Hz, 1H)
32		405.1	(400MHz, CDCl ₃) δ: 8.84 (s, 1H), 7.95 (dd, J=2.8, 5.4 Hz, 1H), 7.73 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.10 (s, 1H), 5.12 (d, J=1.4 Hz, 1H), 5.04-4.93 (m, 1H), 4.80 (dd, J=2.3, 12.8 Hz, 1H), 4.66 (d, J=9.0 Hz, 1H), 4.44 (br t, J=8.1 Hz, 1H), 4.21 (dd, J=8.4, 12.8 Hz, 1H), 3.97 (s, 3H), 1.99-1.86 (m, 1H), 1.95 (s, 3H)

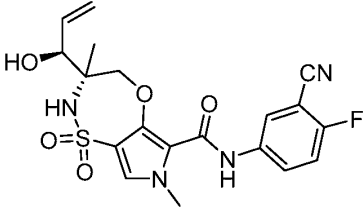
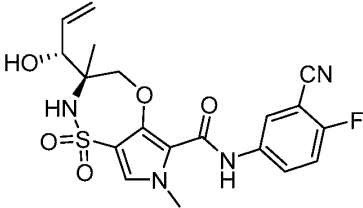
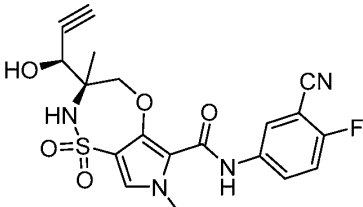
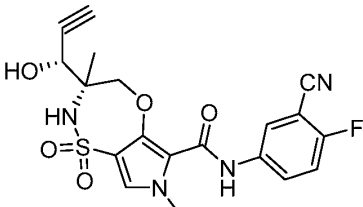
Compound No.	Structure	MS [M+1] +	¹ H NMR
33		448.2	(400MHz, CDCl ₃) δ: 8.82 (s, 1H), 7.84 (dd, J=2.6, 6.0 Hz, 1H), 7.74 (td, J=3.6, 8.7 Hz, 1H), 7.21 (t, J=9.3 Hz, 1H), 7.10 (s, 1H), 5.13 (d, J=1.4 Hz, 1H), 5.03 (s, 1H), 4.81 (dd, J=2.3, 12.8 Hz, 1H), 4.66 (br d, J=3.9 Hz, 1H), 4.46 (br s, 1H), 4.21 (dd, J=8.4, 12.8 Hz, 1H), 3.99 (s, 3H), 1.96 (s, 3H)
34		489.1	(400MHz, DMSO-d ₆) δ: 9.52 (s, 1H), 8.18 (dd, J=2.8, 5.8 Hz, 1H), 8.08-7.97 (m, 2H), 7.60-7.45 (m, 2H), 6.97 (s, 1H), 6.10 (dd, J=10.9, 17.0 Hz, 1H), 5.67 (dd, J=1.2, 17.1 Hz, 1H), 5.49 (d, J=11.8 Hz, 1H), 4.72 (d, J=11.6 Hz, 1H), 4.11 (br t, J=9.6 Hz, 1H), 3.83 (s, 3H), 3.75 (dd, J=9.0, 12.6 Hz, 1H)
35A		433.2	(400MHz, DMSO-d ₆) δ: 9.55 (s, 1H), 8.20 (dd, J=2.6, 5.6 Hz, 1H), 8.10-7.95 (m, 1H), 7.64 (br s, 1H), 7.57-7.41 (m, 2H), 5.94 (s, 1H), 5.02 (d, J=12.1 Hz, 1H), 3.98 (dd, J=8.8, 12.7 Hz, 1H), 3.82 (s, 3H), 3.70 (br d, J=8.5 Hz, 1H), 3.53 (s, 1H), 1.41 (s, 3H)
35B		433.2	(400MHz, DMSO-d ₆) δ: 9.53 (s, 1H), 8.20 (dd, J=2.7, 5.8 Hz, 1H), 8.06 (ddd, J=2.8, 4.9, 9.3 Hz, 1H), 7.62 (br s, 1H), 7.57-7.46 (m, 2H), 5.98 (s, 1H), 5.02 (d, J=11.5 Hz, 1H), 4.08-3.94 (m, 1H), 3.83 (s, 3H), 3.63 (br s, 1H), 1.48 (s, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
36A		476.3	(400MHz, DMSO-d ₆) δ: 9.59 (s, 1H), 8.20 (dd, J=2.6, 6.6 Hz, 1H), 7.99-7.92 (m, 1H), 7.72-7.56 (m, 1H), 7.53-7.45 (m, 2H), 5.93 (s, 1H), 5.00 (d, J=12.0 Hz, 1H), 3.99 (dd, J=8.8, 12.6 Hz, 1H), 3.82 (s, 3H), 3.70 (br d, J=8.9 Hz, 1H), 3.52 (s, 1H), 1.41 (s, 3H)
36B		476.3	(400MHz, DMSO-d ₆) δ: 9.57 (s, 1H), 8.20 (dd, J=2.6, 6.4 Hz, 1H), 7.97 (br dd, J=3.6, 8.8 Hz, 1H), 7.60 (br s, 1H), 7.53-7.42 (m, 2H), 5.98 (s, 1H), 4.99 (d, J=11.6 Hz, 1H), 4.00 (dd, J=8.9, 12.5 Hz, 1H), 3.83 (s, 3H), 3.70-3.57 (m, 1H), 1.48 (s, 3H)
37		487.1	(400MHz, CDCl ₃) δ: 8.70 (s, 1H), 8.00 (dd, J=2.7, 5.6 Hz, 1H), 7.66-7.59 (m, 1H), 7.19 (t, J=8.6 Hz, 1H), 7.13 (s, 1H), 6.84 (d, J=4.4 Hz, 1H), 5.29 (d, J=4.3 Hz, 1H), 5.22 (dd, J=2.5, 13.9 Hz, 1H), 5.01 (br d, J=13.6 Hz, 1H), 4.59 (s, 1H), 3.98 (s, 3H), 2.85 (s, 1H)
38A		435.1	(400MHz, DMSO-d ₆) δ: 9.36 (s, 1H), 8.17 (dd, J=2.6, 5.6 Hz, 1H), 8.07-7.98 (m, 1H), 7.70 (s, 1H), 7.53 (t, J=9.1 Hz, 1H), 7.48 (s, 1H), 5.79 (dd, J=11.1, 17.6 Hz, 1H), 5.44 (d, J=17.8 Hz, 1H), 5.34 (d, J=11.6 Hz, 1H), 5.20 (d, J=5.9 Hz, 1H), 4.89-4.82 (m, 1H), 4.06 (quin, J=6.1 Hz, 1H), 3.82 (s, 3H), 1.05 (d, J=6.3 Hz, 3H)

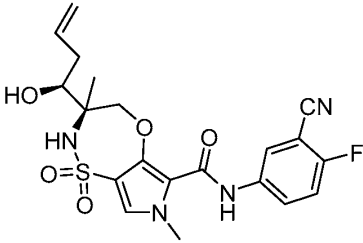
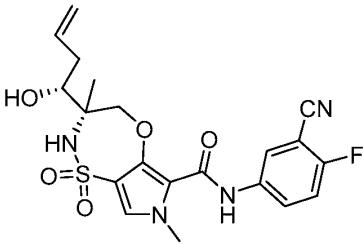
Compound No.	Structure	MS [M+1] +	¹ H NMR
38B		435.3	(400MHz, DMSO-d ₆) δ: 9.36 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.02 (ddd, J=2.8, 4.9, 9.1 Hz, 1H), 7.70 (br s, 1H), 7.53 (t, J=9.1 Hz, 1H), 7.47 (s, 1H), 5.79 (dd, J=11.1, 17.6 Hz, 1H), 5.44 (dd, J=1.1, 17.6 Hz, 1H), 5.34 (d, J=11.3 Hz, 1H), 5.21 (br d, J=5.6 Hz, 1H), 4.88-4.72 (m, 2H), 4.10-4.01 (m, 1H), 3.82 (s, 3H), 1.05 (d, J=6.1 Hz, 3H)
38C		435.3	(400MHz, DMSO-d ₆) δ: 9.37 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.02 (ddd, J=2.8, 4.8, 9.2 Hz, 1H), 7.57-7.43 (m, 3H), 5.87 (dd, J=11.1, 17.8 Hz, 1H), 5.42-5.25 (m, 2H), 4.93-4.82 (m, 2H), 4.73 (d, J=13.6 Hz, 1H), 3.89 (quin, J=6.2 Hz, 1H), 3.81 (s, 3H), 1.13 (d, J=6.4 Hz, 3H)
38D		435.1	(400MHz, DMSO-d ₆) δ: 9.37 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.02 (ddd, J=2.6, 4.9, 9.2 Hz, 1H), 7.56-7.45 (m, 3H), 5.37 (d, J=17.3 Hz, 1H), 5.28 (d, J=11.4 Hz, 1H), 4.89-4.83 (m, 2H), 4.73 (d, J=13.6 Hz, 1H), 3.89 (quin, J=6.3 Hz, 1H), 3.81 (s, 3H), 1.13 (d, J=6.4 Hz, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
39A		475.2	(400MHz, DMSO-d ₆) δ: 9.56 (s, 1H), 8.19 (dd, J=2.8, 5.8 Hz, 1H), 8.08-8.01 (m, 1H), 7.62-7.49 (m, 3H), 5.71 (dd, J=11.1, 17.8 Hz, 1H), 5.35 (d, J=11.4 Hz, 1H), 5.17 (d, J=17.5 Hz, 1H), 4.71 (br d, J=12.3 Hz, 1H), 3.82 (s, 3H), 3.76 (br dd, J=9.1, 12.8 Hz, 1H), 3.73-3.65 (m, 2H), 3.59 (br s, 1H), 3.49-3.38 (m, 2H), 1.89-1.74 (m, 2H), 1.72-1.57 (m, 2H)
39B		475.2	(400MHz, DMSO-d ₆) δ: 9.55 (s, 1H), 8.18 (dd, J=2.6, 5.6 Hz, 1H), 8.07-8.01 (m, 1H), 7.60-7.49 (m, 3H), 5.70 (dd, J=11.1, 17.8 Hz, 1H), 5.34 (d, J=11.0 Hz, 1H), 5.17 (br d, J=17.8 Hz, 1H), 4.71 (br d, J=12.6 Hz, 1H), 3.82 (s, 3H), 3.76 (br dd, J=9.0, 12.6 Hz, 1H), 3.72-3.64 (m, 2H), 3.62-3.54 (m, 1H), 3.48-3.38 (m, 2H), 1.89-1.55 (m, 4H)
40A		449.1	(400MHz, DMSO-d ₆) δ: 9.51 (s, 1H), 8.20 (dd, J=2.8, 5.8 Hz, 1H), 8.06 (ddd, J=2.8, 4.9, 9.2 Hz, 1H), 7.57-7.43 (m, 3H), 5.93-5.78 (m, 1H), 5.13-5.03 (m, 2H), 4.98 (d, J=12.1 Hz, 1H), 4.81 (s, 1H), 3.96 (dd, J=8.8, 12.7 Hz, 1H), 3.83 (s, 3H), 3.58 (br t, J=9.1 Hz, 1H), 2.37-2.28 (m, 1H), 2.10 (br dd, J=7.1, 13.9 Hz, 1H), 1.19 (s, 3H)

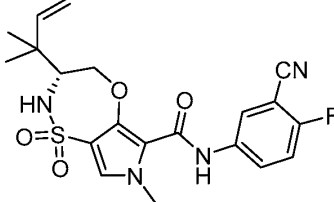
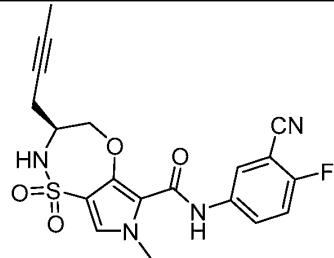
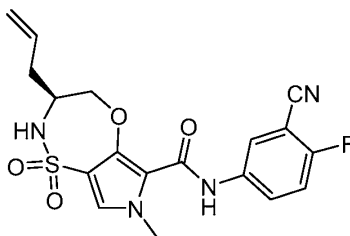
Compound No.	Structure	MS [M+1] +	¹ H NMR
40B		449.3	(400MHz, DMSO-d ₆) δ: 9.52 (s, 1H), 8.20 (dd, J=2.8, 5.8 Hz, 1H), 8.04 (ddd, J=2.7, 4.8, 9.2 Hz, 1H), 7.57-7.44 (m, 3H), 5.89 (dt, J=2.2, 9.2 Hz, 1H), 5.16-5.05 (m, 2H), 4.99 (d, J=11.9 Hz, 1H), 4.87 (s, 1H), 3.94 (dd, J=8.9, 12.5 Hz, 1H), 3.83 (s, 3H), 3.62 (br t, J=9.6 Hz, 1H), 2.39-2.22 (m, 2H), 1.01 (s, 3H)
41A		435.1	(400MHz, CDCl ₃) δ: 8.79 (s, 1H), 7.93 (dd, J=2.8, 5.4 Hz, 1H), 7.74 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.06 (s, 1H), 6.08 (ddd, J=6.4, 10.6, 17.2 Hz, 1H), 5.56-5.41 (m, 2H), 5.04 (s, 1H), 4.92 (d, J=13.4 Hz, 1H), 4.75 (d, J=13.3 Hz, 1H), 4.50 (d, J=6.4 Hz, 1H), 3.96 (s, 3H), 2.14 (br s, 1H), 1.43 (s, 4H)
41B		435.0	(400MHz, CDCl ₃) δ: 8.79 (s, 1H), 7.93 (dd, J=2.8, 5.5 Hz, 1H), 7.74 (ddd, J=2.9, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.6 Hz, 1H), 7.06 (s, 1H), 6.08 (ddd, J=6.3, 10.6, 17.1 Hz, 1H), 5.57-5.35 (m, 2H), 5.06 (s, 1H), 4.91 (d, J=13.3 Hz, 1H), 4.75 (d, J=13.4 Hz, 1H), 4.50 (br d, J=6.3 Hz, 1H), 3.95 (s, 3H), 2.18 (br d, J=2.6 Hz, 1H), 1.43 (s, 3H)

Compound No.	Structure	MS [M+1] +	¹ H NMR
41C		435.1	(400MHz, CDCl ₃) δ: 8.76 (s, 1H), 7.91 (dd, J=2.8, 5.4 Hz, 1H), 7.75 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.22 (t, J=8.7 Hz, 1H), 7.09 (s, 1H), 5.99 (ddd, J=7.4, 10.2, 17.3 Hz, 1H), 5.53-5.42 (m, 2H), 5.28 (s, 1H), 4.83 (d, J=13.5 Hz, 1H), 4.64 (d, J=13.5 Hz, 1H), 4.38 (dd, J=4.9, 7.3 Hz, 1H), 3.98 (s, 3H), 2.56 (d, J=4.9 Hz, 1H), 1.45 (s, 3H)
41D		435.1	(400MHz, CDCl ₃) δ: 8.75 (s, 1H), 7.90 (dd, J=2.8, 5.4 Hz, 1H), 7.74 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.07 (s, 1H), 6.09-5.84 (m, 1H), 5.97 (ddd, J=7.4, 10.2, 17.3 Hz, 1H), 5.52-5.40 (m, 2H), 5.30 (br s, 1H), 4.81 (d, J=13.5 Hz, 1H), 4.62 (d, J=13.5 Hz, 1H), 4.36 (br d, J=7.4 Hz, 1H), 3.96 (s, 3H), 2.61 (br s, 1H), 2.02 (s, 1H), 1.49-1.36 (m, 1H), 1.43 (s, 2H)
42A		433.1	(400MHz, DMSO-d ₆) δ: 9.40 (br s, 1H), 8.18 (dd, J=2.6, 5.6 Hz, 1H), 8.10-7.87 (m, 1H), 7.58-7.36 (m, 2H), 6.04 (br s, 1H), 4.77-4.45 (m, 3H), 3.81 (s, 3H), 3.33 (br s, 1H), 1.43-1.29 (m, 3H)
42B		433.1	(400MHz, DMSO-d ₆) δ: 9.41 (br s, 1H), 8.18 (dd, J=2.6, 5.8 Hz, 1H), 8.10-7.84 (m, 2H), 7.58-7.40 (m, 2H), 6.02 (br s, 1H), 4.78-4.48 (m, 3H), 3.81 (s, 3H), 3.22 (br s, 1H), 1.36 (s, 3H)

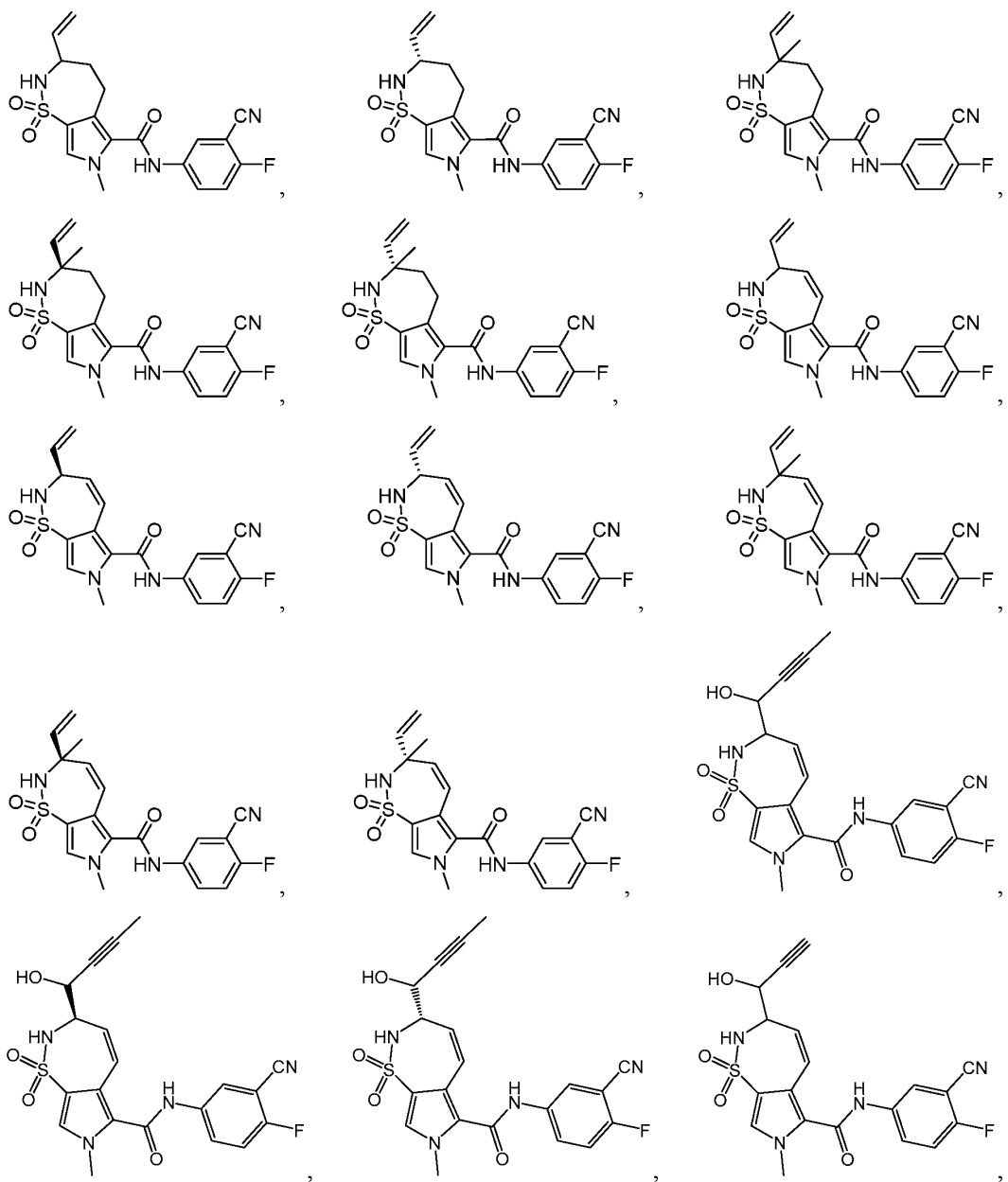
Compound No.	Structure	MS [M+1] +	¹ H NMR
42C		433.1	(400MHz, DMSO-d ₆) δ: 9.45 (s, 1H), 8.17 (dd, J=2.6, 5.8 Hz, 1H), 8.00 (ddd, J=2.7, 4.8, 9.1 Hz, 1H), 7.82 (br d, J=3.0 Hz, 1H), 7.52 (t, J=9.1 Hz, 1H), 7.44 (s, 1H), 5.90-5.75 (m, 1H), 4.66-4.54 (m, 3H), 3.80 (s, 3H), 3.51 (d, J=2.1 Hz, 1H), 1.33 (s, 3H)
42D		433.1	(400MHz, DMSO-d ₆) δ: 9.45 (s, 1H), 8.17 (dd, J=2.7, 5.7 Hz, 1H), 8.00 (ddd, J=2.7, 4.9, 9.2 Hz, 1H), 7.83 (br s, 1H), 7.52 (t, J=9.1 Hz, 1H), 7.44 (s, 1H), 5.83 (d, J=6.3 Hz, 1H), 4.68-4.54 (m, 3H), 3.80 (s, 3H), 3.51 (d, J=2.1 Hz, 1H), 1.33 (s, 3H)
43A		523.1	(400MHz, CD ₃ CN) δ: 9.06-8.95 (m, 1H), 8.03 (dd, J=2.8, 5.6 Hz, 1H), 7.95 (ddd, J=2.8, 4.8, 9.1 Hz, 1H), 7.31 (t, J=9.1 Hz, 1H), 7.19 (s, 1H), 5.71 (dd, J=11.0, 17.8 Hz, 2H), 5.51 (d, J=11.0 Hz, 1H), 5.34 (d, J=17.8 Hz, 1H), 4.82 (dd, J=1.6, 12.8 Hz, 1H), 3.99-3.87 (m, 4H), 3.76 (d, J=7.9 Hz, 1H), 3.04-2.90 (m, 4H), 2.46-2.33 (m, 1H), 2.30-2.18 (m, 3H)
43B		523.1	(400MHz, CD ₃ CN) δ: 9.01 (s, 1H), 8.02 (dd, J=2.7, 5.6 Hz, 1H), 7.98-7.90 (m, 1H), 7.31 (t, J=9.1 Hz, 1H), 7.19 (s, 1H), 5.71 (br dd, J=11.0, 17.6 Hz, 2H), 5.51 (d, J=11.0 Hz, 1H), 5.34 (d, J=17.8 Hz, 1H), 4.82 (dd, J=1.4, 12.6 Hz, 1H), 3.99-3.88 (m, 4H), 3.81-3.70 (m, 1H), 3.05-2.91 (m, 4H), 2.47-2.33 (m, 1H), 2.31-2.18 (m, 4H)

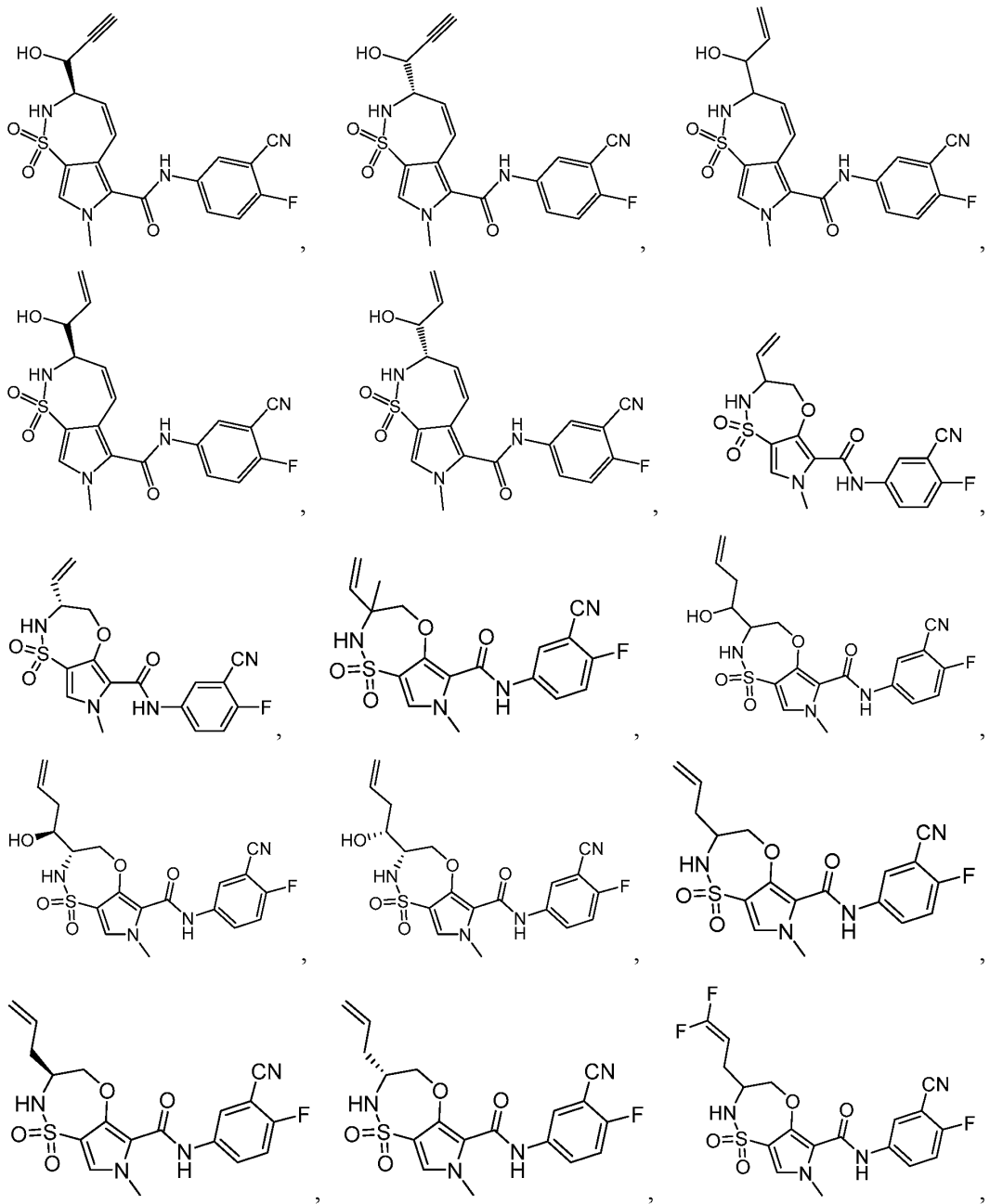
Compound No.	Structure	MS [M+1] +	¹ H NMR
44A		449.1	(400MHz, CDCl ₃) δ: 8.82 (s, 1H), 7.96 (dd, J=2.7, 5.4 Hz, 1H), 7.74 (ddd, J=2.7, 4.6, 9.1 Hz, 1H), 7.22 (t, J=8.7 Hz, 1H), 7.07 (s, 1H), 6.00-5.80 (m, 1H), 5.35-5.21 (m, 2H), 5.07-4.84 (m, 2H), 4.73 (d, J=13.3 Hz, 1H), 4.06 (td, J=2.4, 10.4 Hz, 1H), 3.97 (s, 3H), 2.79 (br dd, J=3.6, 13.8 Hz, 1H), 2.18-2.00 (m, 2H), 1.47 (s, 3H)
44B		449.1	(400MHz, CDCl ₃) δ: 8.82 (s, 1H), 7.96 (dd, J=2.8, 5.5 Hz, 1H), 7.74 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.22 (t, J=8.7 Hz, 1H), 7.07 (s, 1H), 6.00-5.74 (m, 1H), 5.34-5.20 (m, 2H), 5.01 (d, J=13.4 Hz, 1H), 4.91 (s, 1H), 4.73 (d, J=13.4 Hz, 1H), 4.06 (br d, J=10.6 Hz, 1H), 3.97 (s, 3H), 2.79 (br dd, J=3.6, 13.8 Hz, 1H), 2.20-2.00 (m, 2H), 1.47 (s, 3H)

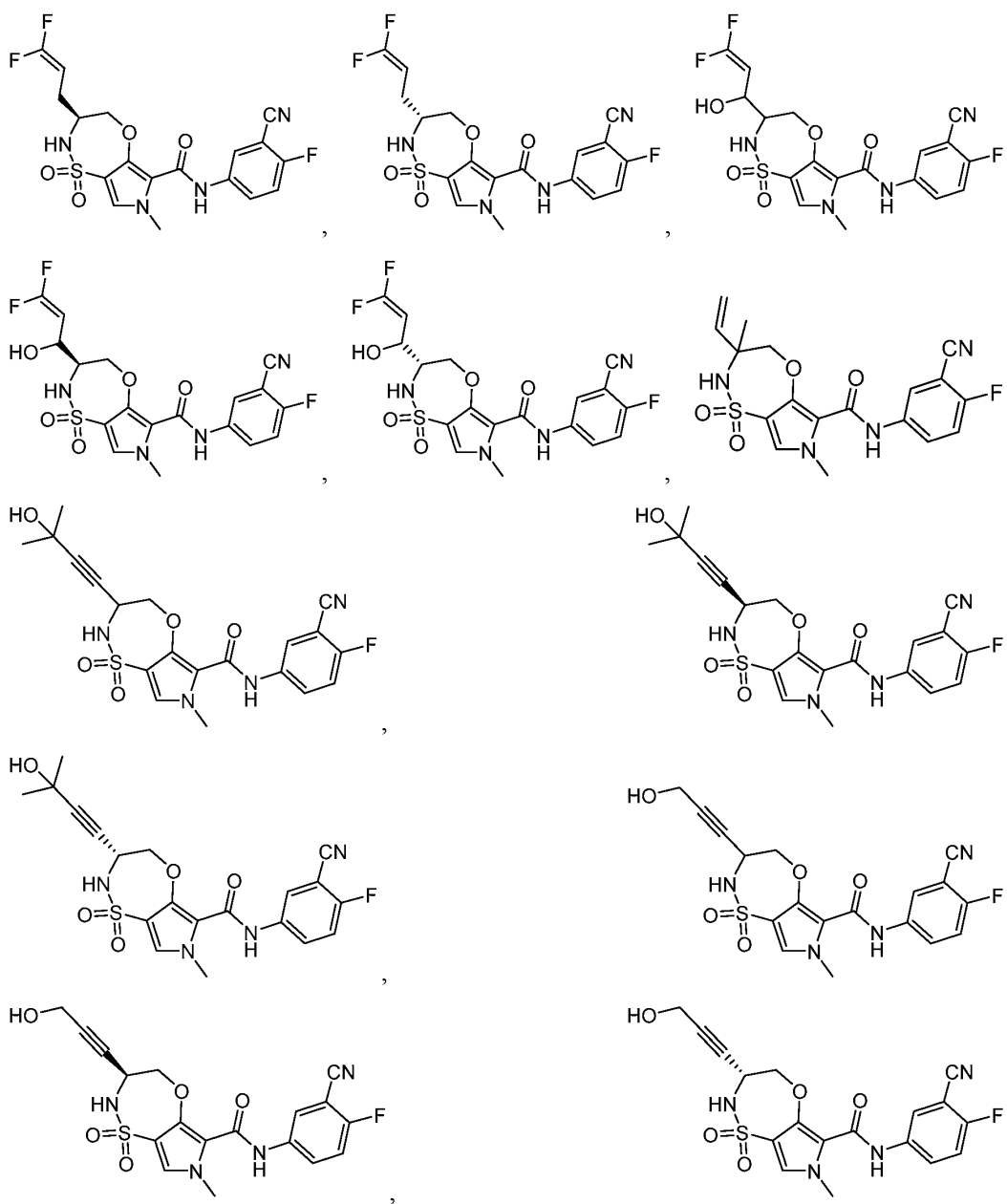
Compound No.	Structure	MS [M+1] +	¹ H NMR
44C		449.1	(400MHz, CDCl ₃) δ: 8.77 (s, 1H), 7.91 (dd, J=2.7, 5.4 Hz, 1H), 7.76 (ddd, J=2.8, 4.6, 9.1 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.08 (s, 1H), 5.87 (dddd, J=5.9, 8.3, 10.6, 16.4 Hz, 1H), 5.38-5.11 (m, 3H), 4.78 (d, J=13.3 Hz, 1H), 4.57 (d, J=13.3 Hz, 1H), 3.96 (s, 3H), 3.74 (br d, J=9.4 Hz, 1H), 2.58-2.42 (m, 2H), 2.35 (td, J=9.3, 14.0 Hz, 1H), 1.50 (s, 3H)
44D		449.1	(400MHz, CDCl ₃) δ: 8.77 (s, 1H), 7.91 (dd, J=2.7, 5.4 Hz, 1H), 7.76 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.08 (s, 1H), 5.97-5.78 (m, 1H), 5.44-5.08 (m, 3H), 4.78 (d, J=13.3 Hz, 1H), 4.57 (d, J=13.3 Hz, 1H), 3.96 (s, 3H), 3.80-3.69 (m, 1H), 2.58-2.42 (m, 2H), 2.35 (td, J=9.3, 14.1 Hz, 1H), 1.50 (s, 3H)
45A		433.1	(400MHz, CDCl ₃) δ: 8.83 (s, 1H), 7.94 (dd, J=2.7, 5.4 Hz, 1H), 7.74 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.21 (t, J=8.7 Hz, 1H), 7.09 (s, 1H), 5.81 (dd, J=10.8, 17.4 Hz, 1H), 5.27-5.16 (m, 2H), 4.80 (dd, J=1.9, 12.6 Hz, 1H), 4.52-4.46 (m, 1H), 4.00-3.92 (m, 4H), 3.79 (br t, J=7.4 Hz, 1H), 1.27 (s, 3H), 1.20 (s, 3H)

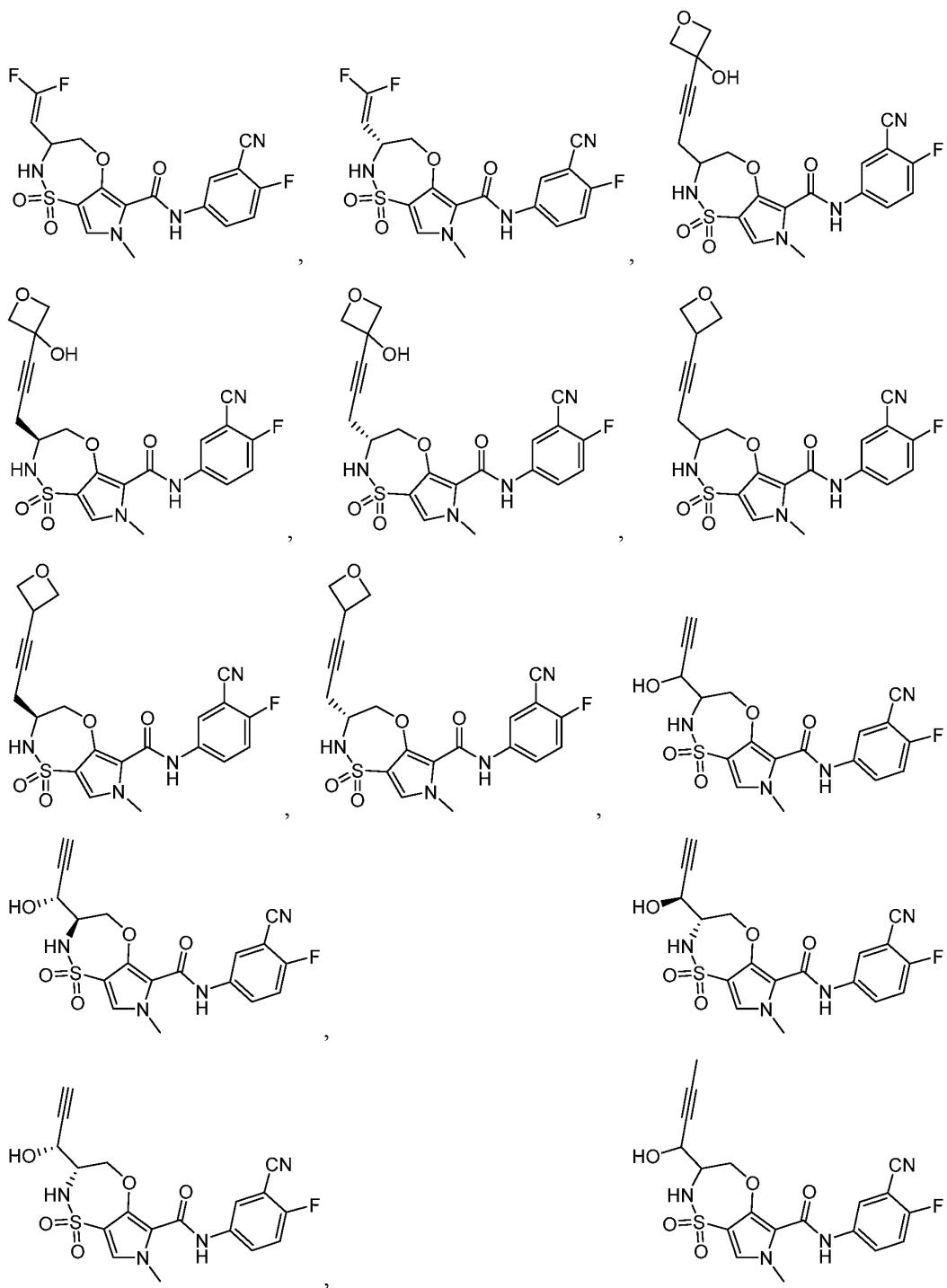
Compound No.	Structure	MS [M+1] +	¹ H NMR
45B		433.2	(400MHz, CDCl ₃) δ: 8.83 (s, 1H), 7.94 (dd, J=2.8, 5.4 Hz, 1H), 7.74 (ddd, J=2.8, 4.5, 9.1 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.09 (s, 1H), 5.81 (dd, J=10.7, 17.4 Hz, 1H), 5.29-5.13 (m, 2H), 4.80 (dd, J=1.8, 12.6 Hz, 1H), 4.51 (d, J=10.5 Hz, 1H), 4.01-3.92 (m, 4H), 3.84-3.76 (m, 1H), 1.27 (s, 3H), 1.20 (s, 3H)
46		416.8	(400MHz, DMSO-d ₆) δ: 9.55 (s, 1H), 8.18 (m, 1H), 8.02 (m, 1H), 7.80 (m, 1H), 7.50-7.54 (m, 2H), 4.70 (m, 1H), 3.92-3.97 (m, 1H), 3.81 (s, 3H), 3.73 (m, 1H), 2.33-2.42 (m, 2H), 1.78 (s, 3H)
47		405.0	(400MHz, DMSO-d ₆) δ: 8.83 (s, 1H), 7.95 (dd, J=2.8, 5.3 Hz, 1H), 7.73 (ddd, J=2.8, 4.6, 9.2 Hz, 1H), 7.20 (t, J=8.7 Hz, 1H), 7.08 (s, 1H), 5.89 - 5.75 (m, 1H), 5.29 (s, 1H), 5.27 - 5.22 (m, 1H), 4.74 (dd, J=2.3, 12.8 Hz, 1H), 4.69 - 4.59 (m, 1H), 4.15 (dd, J=7.8, 12.8 Hz, 1H), 4.09 - 3.99 (m, 1H), 3.96 (s, 3H), 2.60 - 2.35 (m, 2H)

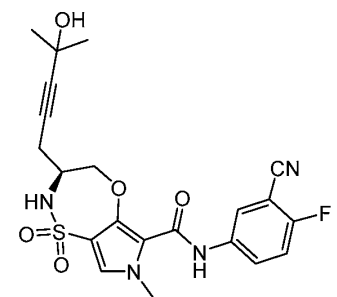
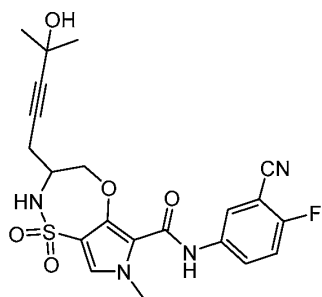
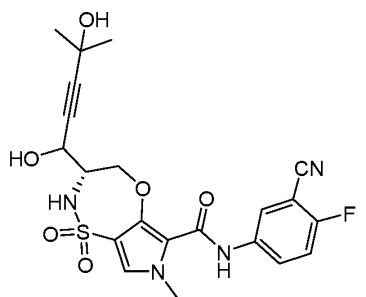
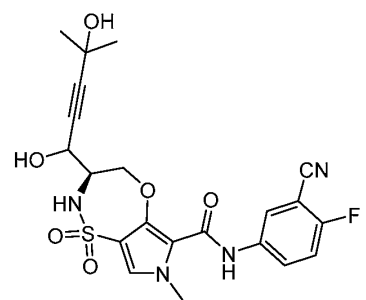
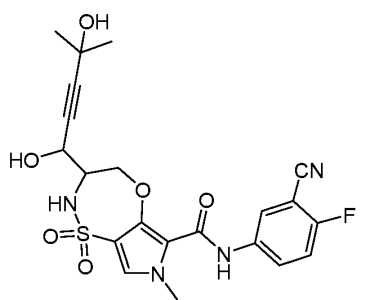
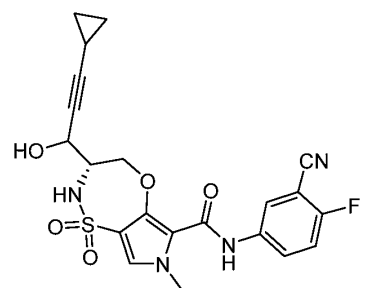
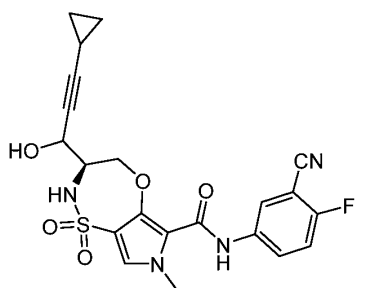
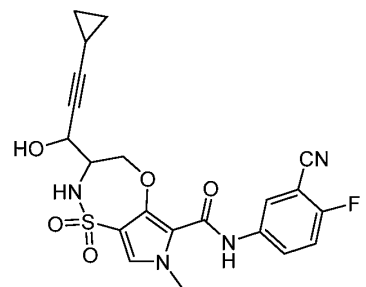
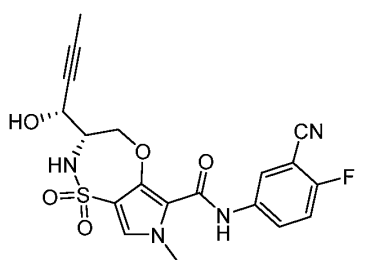
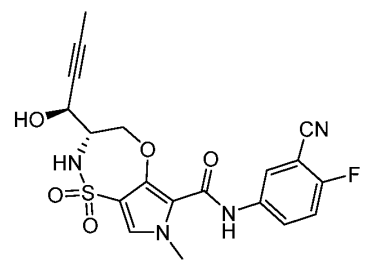
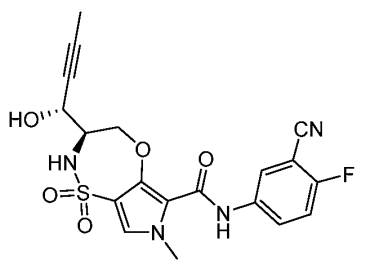
The stereochemistry shown for each of compounds in Table 1 is relative and not absolute.

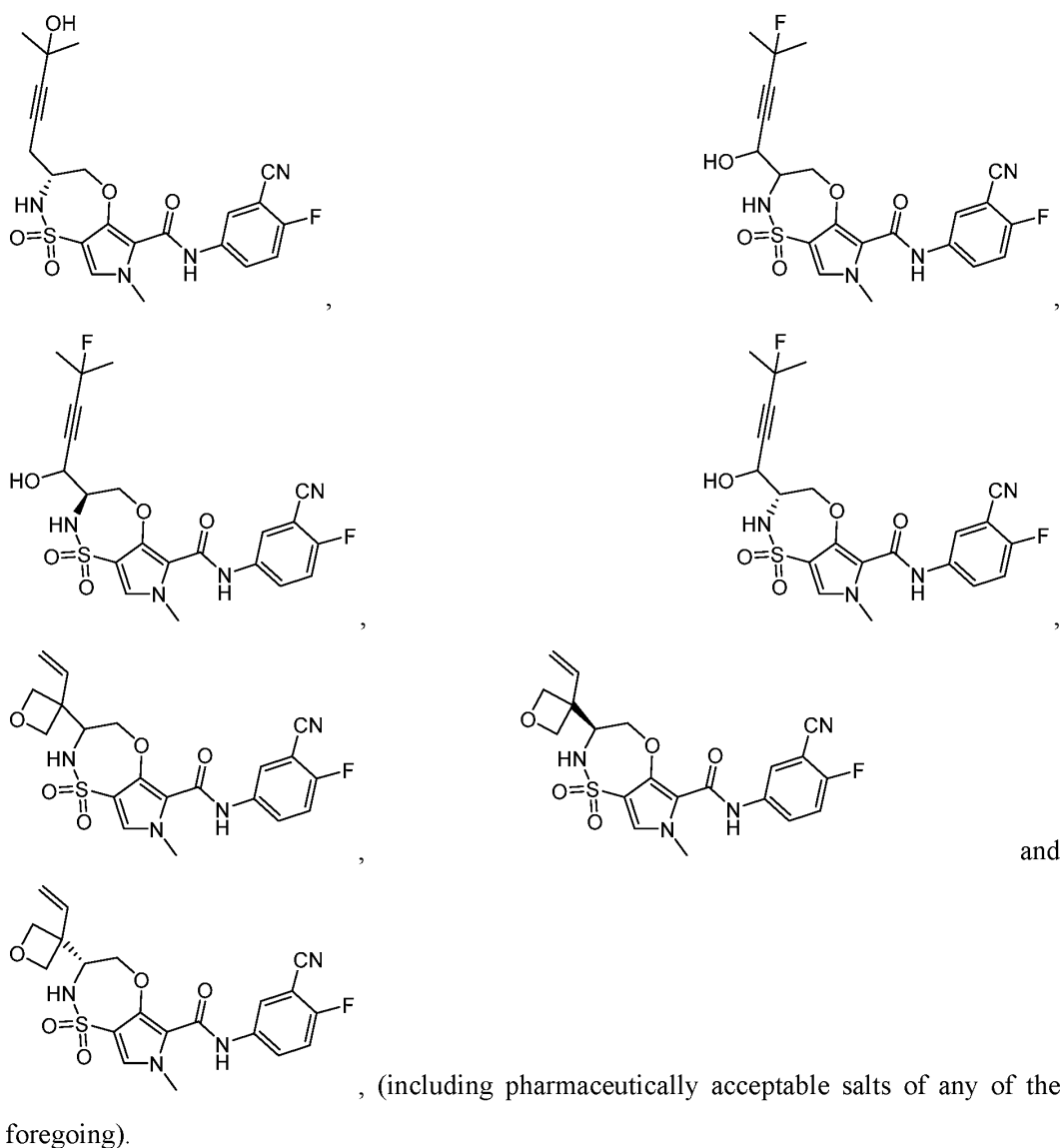












Example A
HBV-DNA Antiviral Assay using HepG2.2.15 cells

[0194] The following assay procedure describes the HBV antiviral assay. This assay uses HepG2.2.15 cells, which have been transfected with HBV genome, and extracellular HBV DNA quantification as endpoint. Cell viability is assessed in parallel by measuring the intracellular ATP content using the CellTiter-Glo® reagent from Promega.

[0195] On day 0, HepG2.2.15 cells were seeded in 96-well plates at a density of 6.0×10^4 cells/well (0.1 ml/well). The cells were incubated at 37°C and 5% CO₂.

[0196] On day 1, the test articles were diluted and added to cell culture wells (8 concentrations, 4-fold dilution, in duplicate). GLS4, Tenofovir and Sorafenib were used as reference compounds. 100 µl of culture medium containing the compounds was added to the plate, and the final total volume per well was 200 µl. The final concentration of DMSO in the culture medium was 0.5%. The plate map of compound treatment is shown below. The cells were cultured at 37 °C and 5% CO₂ for 3 days.

The plate map of compound treatment

	1	2	3	4	5	6	7	8	9	10	11	12
A	High dose			4-fold dilution, 8 dilution points, duplicate						Low dose		
B	compound 1									ETV(1µM)	0.5%DMSO control	Blank
C												
D												
E	compound 2									ETV(1µM)	0.5%DMSO control	Blank
F												
G												
H	compound 3									ETV(1µM)	0.5%DMSO control	Blank
	High dose			4-fold dilution, 8 dilution points, duplicate						Low dose		

[0197] On day 4, the plates were refreshed with culture media with compounds.

[0198] On day 7, cell viability was assessed using the CellTiter-Glo[®], and the cell culture supernatants were collected for determination of HBV DNA by qPCR.

HBV DNA quantification by qPCR

[0199] Extracellular DNA was isolated with QIAamp 96 DNA Blood Kit per the manufacturer's manual. HBV DNA was then quantified by qPCR with HBV specific primers and probes as specified in Table 2 using the FastStart Universal MasterMix from Roche on an ABI-7900HT. The PCR cycle program consisted of 95°C for 10 min, followed by 40 cycles at 95°C for 15 sec and 60°C for 1 min.

Table 2: HBV DNA Primers and Probe

Items	Name	Sequence (5'→3')
HBV Primer	HBV-forward	GTGTCTGCGGCGTTTTATCA
	HBV-reverse	GACAAACGGGCAACATACCTT
HBV Probe	HBV probe	FAM-CCTCTKCATCCTGCTGCTATGCCTCATC- TAMRA

[0200] A DNA standard was prepared by dilution of the pAAV2 HBV1.3 plasmid with concentrations ranging from 10 to 1×10^7 copies/ μ L and used to generate a standard curve by plotting Ct value vs. the concentration of the HBV plasmid DNA standard. The quantity of HBV DNA in each sample was determined by interpolating from the standard curve.

Cell viability

[0201] After harvest of the supernatants, the cell viability was detected by CellTiter-Glo[®] according to the manufacturer's manual. In brief, 50 μ L of fresh cell culture medium was added to the culture plates, followed by addition of 50 μ L CellTiter-Glo into each well. The plates were incubated at room temperature for 10 mins. The luminescence signal was collected on a BioTek Synergy 2 plate reader.

Data analysis

[0202] Cell viability was calculated as follows: % Cell viability = (luminescence value of test sample - average luminescence value of blank) / (average luminescence value of 0.5% DMSO control - average luminescence of blank) $\times$ 100%. HBV DNA inhibition was calculated as follows: 100-(HBV DNA copy number of test sample - HBV DNA copy number of ETV)/ HBV DNA copy number of 0.5%DMSO control - HBV DNA copy number of ETV) $\times$ 100%. The CC₅₀, EC₅₀ and EC₉₀ values were determined by dose-response curves fitted by GraphPad Prism using "log (agonist) vs. response -- Variable slope".

[0203] Compounds of Formula (I) and Formula (II) are active against HBV as shown in Table 3, where 'A' indicates an EC₅₀ < 1 μ M, 'B' indicates an EC₅₀ of ≥ 1 μ M and < 10 μ M and 'C' indicates an EC₅₀ ≥ 10 μ M and < 50 μ M.

Table 3 – Activity of compounds

No.	EC ₅₀
1	A
2A, 2B, 2C 2D	A*

No.	EC ₅₀
3A, 3B, 3C 3D	A*
4A, 4B, 4C 4D	A*
6A	A

* Indicate that at least 2 of the diastereomers have the indicated activity

EXAMPLE B

HBV-DNA Antiviral Assay using HepG2.117 cells

[0204] The following assay procedure describes the HBV antiviral assay, using HepG2.117 cells, which carry a stably integrated genotype D HBV genome under the control of a Tet-off promoter, and intracellular HBV DNA quantification as endpoint. Cell viability is assessed in parallel by measuring the intracellular ATP content using ATPlite (Perkin Elmer).

[0205] On day 0, HepG2.117 cells (which are maintained in routine cell culture with doxycycline present in the medium at a final concentration of 1 µg/mL) were seeded in 96-well plates (white with clear bottom) at a density of 2.0×10^4 cells/well (0.1 mL/well) in medium without doxycycline to induce pgRNA transcription and subsequent formation of HBV particles. The cells were incubated at 37 °C and 5% CO₂.

[0206] On day 1, medium was removed from each well, the test articles were diluted in culture medium without doxycycline and 100 µL was added to cell culture wells (9 concentrations, 4-fold dilution). For each plate, 6 untreated (merely DMSO) wells were included. The final concentration of DMSO in the culture medium was 2%. Each plate was prepared in duplicate (one for HBV DNA extraction, one for ATPlite measurement). The cells were incubated at 37 °C and 5% CO₂ for 3 days.

[0207] On day 4, cell viability was assessed using ATPlite and cell lysates were prepared for HBV DNA extraction and subsequent quantification by qPCR.

HBV DNA quantification by qPCR

[0208] Medium was removed from each well and 100 μ L of 0.33% NP-40 in H₂O was added to each well. Plates were sealed, incubated at 4 °C for 5 mins, vortexed extensively and centrifuged briefly. Next, 35 μ L of lysate was added to 65 μ L QuickExtract DNA Extraction Solution (Epicentre) in a PCR plate for each well. PCR plate was incubated at 65 °C for 6 mins, 98 °C for 2 mins and finally cooled to 4 °C. HBV DNA was then quantified by qPCR with HBV-specific primers and probes as specified in Table 4 using the Bio-Rad SSOAdvanced Universal Probes Supermix on a CFX96 machine (Bio-Rad). The PCR cycle program consisted of 95 °C for 3 mins, followed by 40 cycles at 95 °C for 10 sec and 60 °C for 30 sec.

Table 4: HBV DNA Primers and Probe for HepG2.117 assay

Items	Name	Sequence (5'→3')
HBV Primer	HBV-forward	GTGTCTGCGGCGTTTTATCA
	HBV-reverse	GACAAACGGGCAACATACCTT
HBV Probe	HBV probe	FAM/CCTCTKCAT/ZEN/CCTGCTGCTATGCCTCATC/3IABk FQ/

[0209] A DNA standard was prepared by dilution of an IDT gBlock corresponding to the amplicon with concentrations ranging from 10² to 10⁸ copies/input (i.e. per 4 μ L) and used to generate a standard curve by plotting C_q values vs. HBV DNA standard concentration. The quantity of HBV DNA in each sample was determined by interpolating from the standard curve.

Cell viability

[0210] Using the other plates, the cell viability was quantified by ATPlite according to the manufacturer's manual. In brief, 50 μ L of cell lysis solution was added to the culture plates and shaken for 5', followed by addition of 50 μ L substrate into each well and further shaking. The plates were incubated at room temperature for 10 mins and luminescence signal was subsequently measured on a VarioSkan Lux (ThermoFisher) plate reader.

Data analysis

[0211] Cell viability was calculated as follows: % Cell viability = (luminescence value of test sample) / (average luminescence value of 2% DMSO control) x 100%. HBV DNA inhibition was calculated as follows: 100 - (HBV DNA copy number of test sample) / (average HBV DNA copy number of 2% DMSO control) x 100%. No normalization to entecavir was required due to the excellent dynamic window of this assay. The CC₅₀, EC₅₀ and EC₉₀ values were determined by dose-response curves fitted by GraphPad Prism using "log (agonist) vs. response -- Variable slope".

[0212] Compounds of Formulae (I) and (II) are active against HBV as shown in Table 5, where 'A' indicates an EC₅₀ < 1 nM, 'B' indicates an EC₅₀ of ≥1 nM and < 10 nM, 'C' indicates an EC₅₀ ≥ 10 nM and < 100 nM, and 'D' indicates an EC₅₀ ≥ 100 nM and < 1000 nM.

Table 5

Compound	EC ₅₀ HepG2.117 (nM)	Compound	EC ₅₀ HepG2.117 (nM)	Compound	EC ₅₀ HepG2.117 (nM)
1	B	19B	B	31A	B
8	B	20	B	31B	B
9A	C	21	C	32	B
9B	C	22A	C	33	B
10A	C	22B	B	34	A
10B	C	23A	B	35A	B
11A	B	23B	B	35B	B
11B	B	24	B	36A	C
12A	C	25A	C	36B	B
12B	C	25B	C	37	C
13	B	26	C	38A	C
14A	B	27A	C	38B	C
14B	B	27B	B	38C	B
15A	B	27C	C	38D	D
15B	B	27D	A	39A	C
16A	B	28A	B	39B	C
16B	B	28B	B	40A	B
17A	B	29A	C	40B	B
17B	B	29B	B	41A	B
18	B	30A	B	41B	B
19A	B	30B	C	41C	C

Compound	EC ₅₀ HepG2.117 (nM)
41D	C
42A	B
42B	B
42C	B
42D	C
43A	C

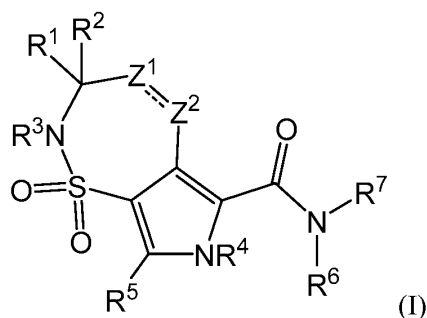
Compound	EC ₅₀ HepG2.117 (nM)
43B	C
44A	C
44B	B
44C	B
44C	C
45A	B

Compound	EC ₅₀ HepG2.117 (nM)
45B	B
46	B
47	B

[0213] Although the foregoing has been described in some detail by way of illustrations and examples for purposes of clarity and understanding, it will be understood by those of skill in the art that numerous and various modifications can be made without departing from the spirit of the present disclosure. Therefore, it should be clearly understood that the forms disclosed herein are illustrative only and are not intended to limit the scope of the present disclosure, but rather to also cover all modification and alternatives coming with the true scope and spirit of the invention.

WHAT IS CLAIMED IS:

1. A compound of Formula (I), or a pharmaceutically acceptable salt thereof, having the structure:



wherein:

----- is a single or a double bond,

wherein when ----- is a single bond, then Z¹ is CR^{8A}R^{9A} and Z² is CR^{8B}R^{9B},

and

wherein when ----- is a double bond, then Z¹ and Z² are each independently CR¹⁰,

R¹ is a substituted or an unsubstituted C₂₋₈ alkenyl or a substituted or an unsubstituted C₂₋₈ alkynyl, wherein the substituted C₂₋₈ alkenyl and the substituted C₂₋₈ alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A};

R² is hydrogen, deuterium or a substituted or an unsubstituted C₁₋₄ alkyl, wherein the substituted C₁₋₄ alkyl is substituted with one or more substituents selected from halogen, hydroxy and R^{11B};

R³ is hydrogen, deuterium or an unsubstituted C₁₋₄ alkyl;

R⁴ is hydrogen, deuterium or an unsubstituted C₁₋₄ alkyl;

R⁵ is hydrogen, deuterium, halogen, an unsubstituted C₁₋₄ alkyl, cyano, an unsubstituted C₁₋₄ haloalkyl or an unsubstituted C₃₋₈ monocyclic cycloalkyl;

R⁶ is a substituted phenyl or a substituted pyridyl, wherein the substituted phenyl and the substituted pyridyl is substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C₁₋₄ haloalkyl and an unsubstituted C₁₋₄ alkyl; and

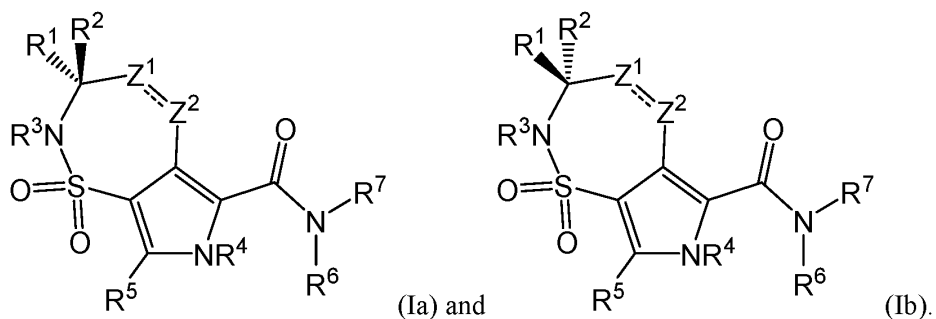
R^7 is hydrogen, deuterium or an unsubstituted C_{1-4} alkyl;

R^{8A} , R^{8B} , R^{9A} and R^{9B} are independently hydrogen, deuterium, halogen, an unsubstituted C_{1-4} alkyl or hydroxy;

each R^{10} are independently hydrogen, deuterium, halogen or an unsubstituted C_{1-4} alkyl; and

R^{11A} and R^{11B} are independently an optionally substituted $-O$ -acyl, an unsubstituted O -linked α -amino acid, $-O-P(=O)(OH)_2$ or $-CH_2-P(=O)(OH)_2$.

2. The compound of Claim 1, or a pharmaceutically acceptable salt thereof, wherein the compound of Formula (I) has a structure selected from the group consisting of:



3. The compound of Claim 1 or 2, or a pharmaceutically acceptable salt thereof, wherein ----- is a single or a double bond; Z^1 is $CR^{8A}R^{9A}$, and Z^2 is $CR^{8B}R^{9B}$.

4. The compound of Claim 3, or a pharmaceutically acceptable salt thereof, wherein R^{8A} is hydrogen.

5. The compound of Claim 3, or a pharmaceutically acceptable salt thereof, wherein R^{8A} is deuterium.

6. The compound of Claim 3, or a pharmaceutically acceptable salt thereof, wherein R^{8A} is halogen.

7. The compound of Claim 3, or a pharmaceutically acceptable salt thereof, wherein R^{8A} is an unsubstituted C_{1-4} alkyl.

8. The compound of Claim 3, or a pharmaceutically acceptable salt thereof, wherein R^{8A} is hydroxy.

9. The compound of any one of Claims 3-8, or a pharmaceutically acceptable salt thereof, wherein R^{9A} is hydrogen.

10. The compound of any one of Claims 3-8, or a pharmaceutically acceptable salt thereof, wherein R^{9A} is deuterium.
11. The compound of any one of Claims 3-8, or a pharmaceutically acceptable salt thereof, wherein R^{9A} is halogen.
12. The compound of any one of Claims 3-8, or a pharmaceutically acceptable salt thereof, wherein R^{9A} is an unsubstituted C₁₋₄ alkyl.
13. The compound of any one of Claims 3-8, or a pharmaceutically acceptable salt thereof, wherein R^{9A} is hydroxy.
14. The compound of any one of Claims 3-13, or a pharmaceutically acceptable salt thereof, wherein R^{8B} is hydrogen.
15. The compound of any one of Claims 3-13, or a pharmaceutically acceptable salt thereof, wherein R^{8B} is deuterium.
16. The compound of any one of Claims 3-13, or a pharmaceutically acceptable salt thereof, wherein R^{8B} is halogen.
17. The compound of any one of Claims 3-13, or a pharmaceutically acceptable salt thereof, wherein R^{8B} is an unsubstituted C₁₋₄ alkyl.
18. The compound of any one of Claims 3-13, or a pharmaceutically acceptable salt thereof, wherein R^{8B} is hydroxy.
19. The compound of any one of Claims 3-18, or a pharmaceutically acceptable salt thereof, wherein R^{9B} is hydrogen.
20. The compound of any one of Claims 3-18, or a pharmaceutically acceptable salt thereof, wherein R^{9B} is deuterium.
21. The compound of any one of Claims 3-18, or a pharmaceutically acceptable salt thereof, wherein R^{9B} is halogen.
22. The compound of any one of Claims 3-18, or a pharmaceutically acceptable salt thereof, wherein R^{9B} is an unsubstituted C₁₋₄ alkyl.
23. The compound of any one of Claims 3-18, or a pharmaceutically acceptable salt thereof, wherein R^{9B} is hydroxy.
24. The compound of Claim 1 or 2, or a pharmaceutically acceptable salt thereof, wherein is a double bond, Z¹ and Z² are each independently CR¹⁰.

25. The compound of Claim 24, or a pharmaceutically acceptable salt thereof, wherein R¹⁰ is hydrogen.

26. The compound of Claim 24, or a pharmaceutically acceptable salt thereof, wherein R¹⁰ is deuterium.

27. The compound of Claim 24, or a pharmaceutically acceptable salt thereof, wherein R¹⁰ is halogen.

28. The compound of Claim 24, or a pharmaceutically acceptable salt thereof, wherein R¹⁰ is an unsubstituted C₁₋₄ alkyl.

29. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R¹ is an unsubstituted C₂₋₈ alkenyl.

30. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R¹ is an unsubstituted C₂₋₈ alkynyl.

31. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R¹ is a substituted C₂₋₈ alkenyl, wherein the substituted C₂₋₈ alkenyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A}.

32. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R¹ is a substituted C₃₋₄ alkenyl, wherein the substituted C₃₋₄ alkenyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A}.

33. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with one or more halogens.

34. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with one or more hydroxys.

35. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted monocyclic C₃₋₆ cycloalkyl.

36. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted monocyclic C₃₋₆ cycloalkyl.

37. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted bicyclic C₃₋₈ cycloalkyl selected from an unsubstituted fused bicyclic C₃₋₈ cycloalkyl, an unsubstituted bridged bicyclic C₃₋₈ cycloalkyl and an unsubstituted spiro bicyclic C₃₋₈ cycloalkyl.

38. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted bicyclic C₃₋₈ cycloalkyl selected from a substituted fused bicyclic C₃₋₈ cycloalkyl, a substituted bridged bicyclic C₃₋₈ cycloalkyl and a substituted spiro bicyclic C₃₋₈ cycloalkyl.

39. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted monocyclic heterocyclyl.

40. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted monocyclic heterocyclyl.

41. The compound of any one of Clams 31-32, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with R^{11A}.

42. The compound of Claim 41, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is an optionally substituted –O–acyl.

43. The compound of Claim 41, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is an unsubstituted O-linked α -amino acid.

44. The compound of Claim 41, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is –O–P(=O)(OH)₂.

45. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R¹ is a substituted C₂₋₈ alkynyl, wherein the substituted C₂₋₈ alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A}.

46. The compound of any one of Clams 1-28, or a pharmaceutically acceptable salt thereof, wherein R^1 is a substituted C_{3-4} alkynyl, wherein the substituted C_{3-4} alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{11A} .

47. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with one or more halogens.

48. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with one or more hydroxys.

49. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted monocyclic C_{3-6} cycloalkyl.

50. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted monocyclic C_{3-6} cycloalkyl.

51. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted bicyclic C_{3-8} cycloalkyl selected from an unsubstituted fused bicyclic C_{3-8} cycloalkyl, an unsubstituted bridged bicyclic C_{3-8} cycloalkyl and an unsubstituted spiro bicyclic C_{3-8} cycloalkyl.

52. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted bicyclic C_{3-8} cycloalkyl selected from a substituted fused bicyclic C_{3-8} cycloalkyl, a substituted bridged bicyclic C_{3-8} cycloalkyl and a substituted spiro bicyclic C_{3-8} cycloalkyl.

53. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted monocyclic heterocyclyl.

54. The compound of any one of Clams 45-46, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted monocyclic heterocyclyl.

55. The compound of any one of Clams 46-47, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with R^{11A} .

56. The compound of Claim 55, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is an optionally substituted $-O-$ acyl.

57. The compound of Claim 55, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is an unsubstituted O-linked α -amino acid.

58. The compound of Claim 55, or a pharmaceutically acceptable salt thereof, wherein R^{11A} is $-O-P(=O)(OH)_2$.

59. The compound of any one of Claims 1-58, or a pharmaceutically acceptable salt thereof, wherein R^2 is hydrogen.

60. The compound of any one of Claims 1-58, or a pharmaceutically acceptable salt thereof, wherein R^2 is deuterium.

61. The compound of any one of Claims 1-58, or a pharmaceutically acceptable salt thereof, wherein R^2 is an unsubstituted C_{1-4} alkyl.

62. The compound of any one of Claims 1-61, or a pharmaceutically acceptable salt thereof, wherein R^3 is hydrogen.

63. The compound of any one of Claims 1-61, or a pharmaceutically acceptable salt thereof, wherein R^3 is deuterium.

64. The compound of any one of Claims 1-61, or a pharmaceutically acceptable salt thereof, wherein R^3 is an unsubstituted C_{1-4} alkyl.

65. The compound of any one of Claims 1-61, or a pharmaceutically acceptable salt thereof, wherein R^3 is a substituted C_{1-4} alkyl.

66. The compound of any one of Claims 1-65, or a pharmaceutically acceptable salt thereof, wherein R^4 is hydrogen.

67. The compound of any one of Claims 1-65, or a pharmaceutically acceptable salt thereof, wherein R^4 is deuterium.

68. The compound of any one of Claims 1-65, or a pharmaceutically acceptable salt thereof, wherein R^4 is an unsubstituted C_{1-4} alkyl.

69. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R^5 is hydrogen.

70. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R^5 is deuterium.

71. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R⁵ is halogen.

72. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₁₋₄ alkyl.

73. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R⁵ is cyano.

74. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₁₋₄ haloalkyl.

75. The compound of any one of Claims 1-68, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₃₋₈ monocyclic cycloalkyl.

76. The compound of any one of Claims 1-75, or a pharmaceutically acceptable salt thereof, wherein R⁶ is a substituted phenyl.

77. The compound of Claim 76, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a mono-substituted phenyl.

78. The compound of Claim 76, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a di-substituted phenyl.

79. The compound of any one of Claims 1-75, or a pharmaceutically acceptable salt thereof, wherein R⁶ is a substituted pyridyl.

80. The compound of Claim 79, or a pharmaceutically acceptable salt thereof, wherein the substituted pyridyl is a mono-substituted pyridyl.

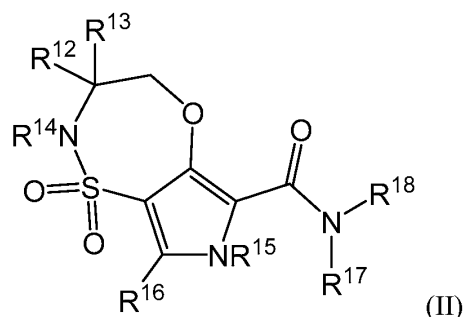
81. The compound of Claim 79, or a pharmaceutically acceptable salt thereof, wherein the substituted pyridyl is a di-substituted pyridyl.

82. The compound of any one of Claims 1-81, or a pharmaceutically acceptable salt thereof, wherein R⁷ is hydrogen.

83. The compound of any one of Claims 1-81, or a pharmaceutically acceptable salt thereof, wherein R⁷ is deuterium.

84. The compound of any one of Claims 1-81, or a pharmaceutically acceptable salt thereof, wherein R⁷ is an unsubstituted C₁₋₄ alkyl.

85. A compound of Formula (II), or a pharmaceutically acceptable salt thereof, having the structure:



wherein:

R^{12} is a substituted or an unsubstituted C_{2-8} alkenyl or a substituted or an unsubstituted C_{2-8} alkynyl, wherein the substituted C_{2-8} alkenyl and the substituted C_{2-8} alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A} ;

R^{13} is hydrogen, deuterium or a substituted or an unsubstituted C_{1-4} alkyl, wherein the substituted C_{1-4} alkyl is substituted with one or more substituents selected from halogen, hydroxy and R^{19B} ;

R^{14} is hydrogen, deuterium or an unsubstituted C_{1-4} alkyl;

R^{15} is hydrogen, deuterium or an unsubstituted C_{1-4} alkyl;

R^{16} is hydrogen, deuterium, halogen, an unsubstituted C_{1-4} alkyl, cyano, an unsubstituted C_{1-4} haloalkyl or an unsubstituted C_{3-8} monocyclic cycloalkyl;

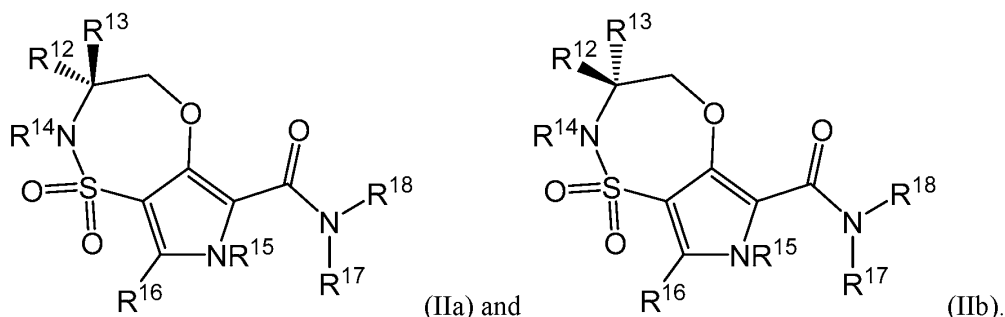
R^{17} is a substituted phenyl or a substituted pyridyl, wherein the substituted phenyl and the substituted pyridyl is substituted with one or more substituents independently selected from halogen, cyano, an unsubstituted C_{1-4} haloalkyl and an unsubstituted C_{1-4} alkyl;

R^{18} is hydrogen, deuterium or an unsubstituted C_{1-4} alkyl; and

R^{19A} and R^{19B} are independently an optionally substituted $-O-$ acyl, an unsubstituted O -linked α -amino acid, $-O-P(=O)(OH)_2$ or $-CH_2-P(=O)(OH)_2$; and

provided that when R^{12} is an unsubstituted 2-butyryl, R^{13} is hydrogen, R^{14} and R^{18} are each hydrogen, R^{15} is methyl and R^{16} is hydrogen, then R^{17} is not 3,4-difluorophenyl.

86. The compound of Claim 85, or a pharmaceutically acceptable salt thereof, wherein the compound of Formula (II) has a structure selected from the group consisting of:



87. The compound of any one of Claims 85-86, or a pharmaceutically acceptable salt thereof, wherein R^{12} is an unsubstituted C_{2-8} alkenyl.

88. The compound of any one of Claims 85-86, or a pharmaceutically acceptable salt thereof, wherein R^{12} is an unsubstituted C_{2-8} alkynyl.

89. The compound of any one of Claims 85-86, or a pharmaceutically acceptable salt thereof, wherein R^{12} is a substituted C_{2-8} alkenyl, wherein the substituted C_{2-8} alkenyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A} .

90. The compound of any one of Claims 85-86, or a pharmaceutically acceptable salt thereof, wherein R^{12} is a substituted C_{3-4} alkenyl, wherein the substituted C_{3-4} alkenyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A} .

91. The compound of any one of Claims 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with one or more halogens.

92. The compound of any one of Claims 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with one or more hydroxys.

93. The compound of any one of Claims 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted monocyclic C_{3-6} cycloalkyl.

94. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted monocyclic C₃₋₆ cycloalkyl.

95. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted bicyclic C₃₋₈ cycloalkyl selected from an unsubstituted fused bicyclic C₃₋₈ cycloalkyl, an unsubstituted bridged bicyclic C₃₋₈ cycloalkyl and an unsubstituted spiro bicyclic C₃₋₈ cycloalkyl.

96. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted bicyclic C₃₋₈ cycloalkyl selected from a substituted fused bicyclic C₃₋₈ cycloalkyl, a substituted bridged bicyclic C₃₋₈ cycloalkyl and a substituted spiro bicyclic C₃₋₈ cycloalkyl.

97. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with an unsubstituted monocyclic heterocyclyl.

98. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with a substituted monocyclic heterocyclyl.

99. The compound of any one of Clams 89-90, or a pharmaceutically acceptable salt thereof, wherein the substituted alkenyl is substituted with R^{19A}.

100. The compound of Claim 99, or a pharmaceutically acceptable salt thereof, wherein R^{19A} is an optionally substituted –O–acyl.

101. The compound of Claim 99, or a pharmaceutically acceptable salt thereof, wherein R^{19A} is an unsubstituted O-linked α -amino acid.

102. The compound of Claim 99, or a pharmaceutically acceptable salt thereof, wherein R^{19A} is –O–P(=O)(OH)₂.

103. The compound of any one of Clams 85-86, or a pharmaceutically acceptable salt thereof, wherein R¹² is a substituted C₂₋₈ alkynyl, wherein the substituted C₂₋₈ alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C₃₋₆ cycloalkyl, an optionally substituted bicyclic C₃₋₈ cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A}.

104. The compound of any one of Clams 85-86, or a pharmaceutically acceptable salt thereof, wherein R^{12} is a substituted C_{3-4} alkynyl, wherein the substituted C_{3-4} alkynyl is substituted with one or more substituents independently selected from halogen, hydroxy, an optionally substituted monocyclic C_{3-6} cycloalkyl, an optionally substituted bicyclic C_{3-8} cycloalkyl, an optionally substituted monocyclic heterocyclyl and R^{19A} .

105. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with one or more halogens.

106. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with one or more hydroxys.

107. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted monocyclic C_{3-6} cycloalkyl.

108. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted monocyclic C_{3-6} cycloalkyl.

109. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted bicyclic C_{3-8} cycloalkyl selected from an unsubstituted fused bicyclic C_{3-8} cycloalkyl, an unsubstituted bridged bicyclic C_{3-8} cycloalkyl and an unsubstituted spiro bicyclic C_{3-8} cycloalkyl.

110. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted bicyclic C_{3-8} cycloalkyl selected from a substituted fused bicyclic C_{3-8} cycloalkyl, a substituted bridged bicyclic C_{3-8} cycloalkyl and a substituted spiro bicyclic C_{3-8} cycloalkyl.

111. The compound of any one of Clams 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with an unsubstituted monocyclic heterocyclyl.

112. The compound of any one of Claims 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with a substituted monocyclic heterocyclyl.

113. The compound of any one of Claims 103-104, or a pharmaceutically acceptable salt thereof, wherein the substituted alkynyl is substituted with R^{19A}.

114. The compound of Claim 113, or a pharmaceutically acceptable salt thereof, wherein R¹⁹ is an optionally substituted –O–acyl.

115. The compound of Claim 113, or a pharmaceutically acceptable salt thereof, wherein R^{19A} is an unsubstituted O-linked α -amino acid.

116. The compound of Claim 113, or a pharmaceutically acceptable salt thereof, wherein R^{19A} is –O–P(=O)(OH)₂.

117. The compound of any one of Claims 85-116, or a pharmaceutically acceptable salt thereof, wherein R¹³ is hydrogen.

118. The compound of any one of Claims 85-116, or a pharmaceutically acceptable salt thereof, wherein R¹³ is deuterium.

119. The compound of any one of Claims 85-116, or a pharmaceutically acceptable salt thereof, wherein R¹³ is an unsubstituted C₁₋₄ alkyl.

120. The compound of any one of Claims 85-116, or a pharmaceutically acceptable salt thereof, wherein R¹³ is a substituted C₁₋₄ alkyl.

121. The compound of any one of Claims 85-120, or a pharmaceutically acceptable salt thereof, wherein R¹⁴ is hydrogen.

122. The compound of any one of Claims 85-120, or a pharmaceutically acceptable salt thereof, wherein R¹⁴ is deuterium.

123. The compound of any one of Claims 85-120, or a pharmaceutically acceptable salt thereof, wherein R¹⁴ is an unsubstituted C₁₋₄ alkyl.

124. The compound of any one of Claims 85-123, or a pharmaceutically acceptable salt thereof, wherein R¹⁵ is hydrogen.

125. The compound of any one of Claims 85-123, or a pharmaceutically acceptable salt thereof, wherein R¹⁵ is deuterium.

126. The compound of any one of Claims 85-123, or a pharmaceutically acceptable salt thereof, wherein R¹⁵ is an unsubstituted C₁₋₄ alkyl.

127. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is hydrogen.

128. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is deuterium.

129. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is halogen.

130. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₁₋₄ alkyl.

131. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is cyano.

132. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₁₋₄ haloalkyl.

133. The compound of any one of Claims 85-126, or a pharmaceutically acceptable salt thereof, wherein R⁵ is an unsubstituted C₃₋₈ monocyclic cycloalkyl.

134. The compound of any one of Claims 85-133, or a pharmaceutically acceptable salt thereof, wherein R¹⁷ is a substituted phenyl.

135. The compound of Claim 134, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a mono-substituted phenyl.

136. The compound of Claim 134, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a di-substituted phenyl.

137. The compound of any one of Claims 85-133, or a pharmaceutically acceptable salt thereof, wherein R¹⁶ is a substituted pyridyl.

138. The compound of Claim 137, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a mono-substituted pyridyl.

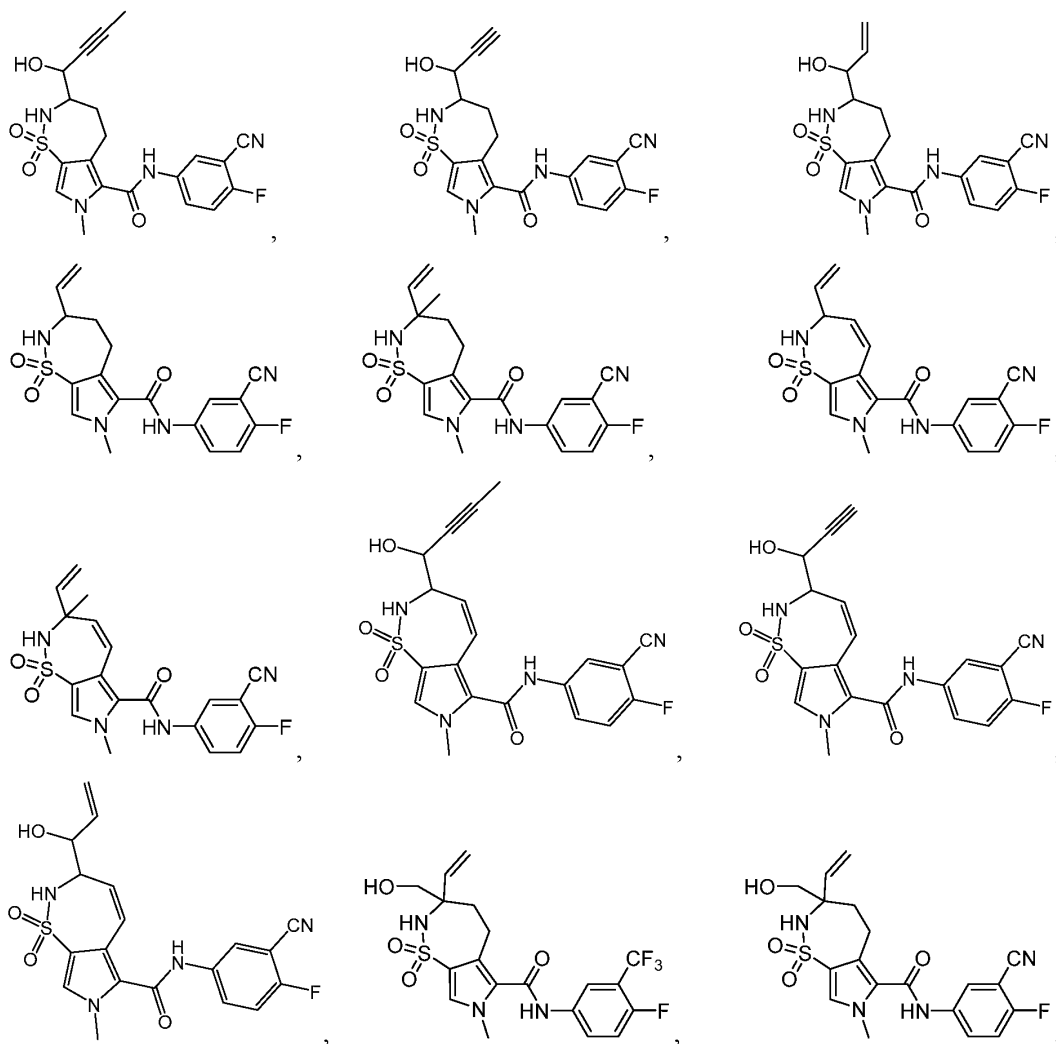
139. The compound of Claim 137, or a pharmaceutically acceptable salt thereof, wherein the substituted phenyl is a di-substituted pyridyl.

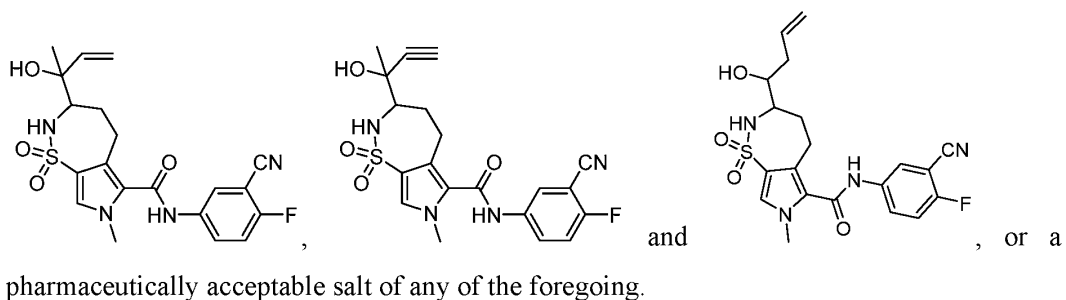
140. The compound of any one of Claims 85-139, or a pharmaceutically acceptable salt thereof, wherein R¹⁸ is hydrogen.

141. The compound of any one of Claims 85-139, or a pharmaceutically acceptable salt thereof, wherein R¹⁸ is deuterium.

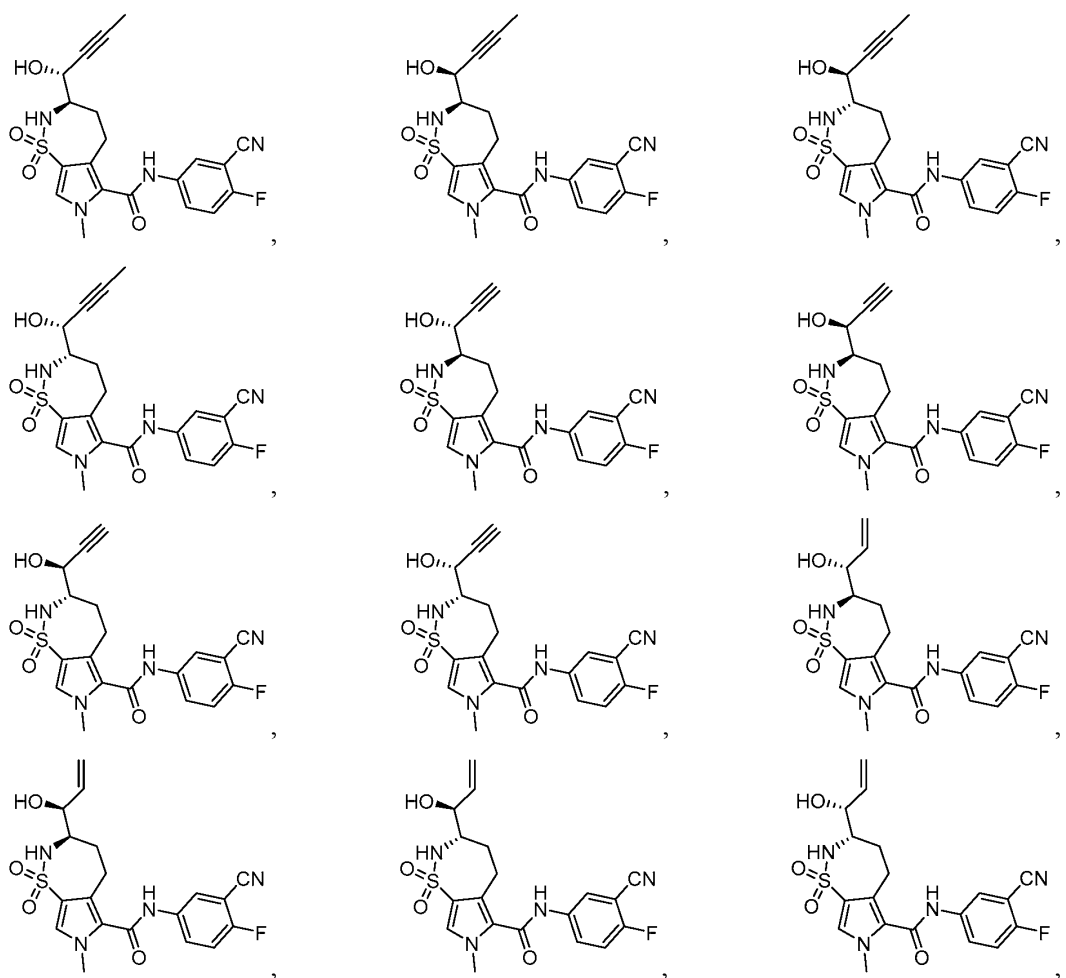
142. The compound of any one of Claims 85-139, or a pharmaceutically acceptable salt thereof, wherein R¹⁸ is an unsubstituted C₁₋₄ alkyl.

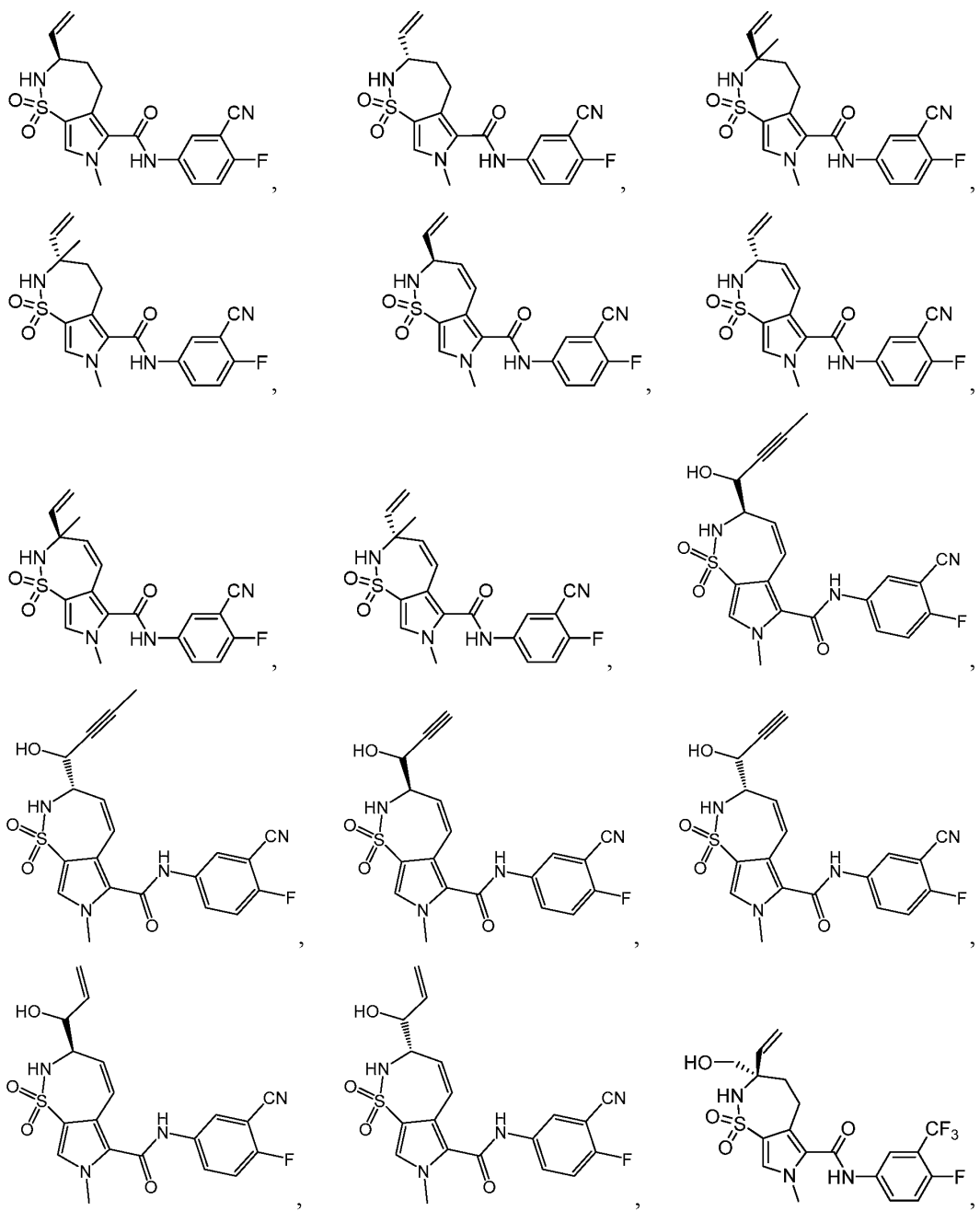
143. The compound of Claim 1, wherein the compound is selected from the group consisting of:

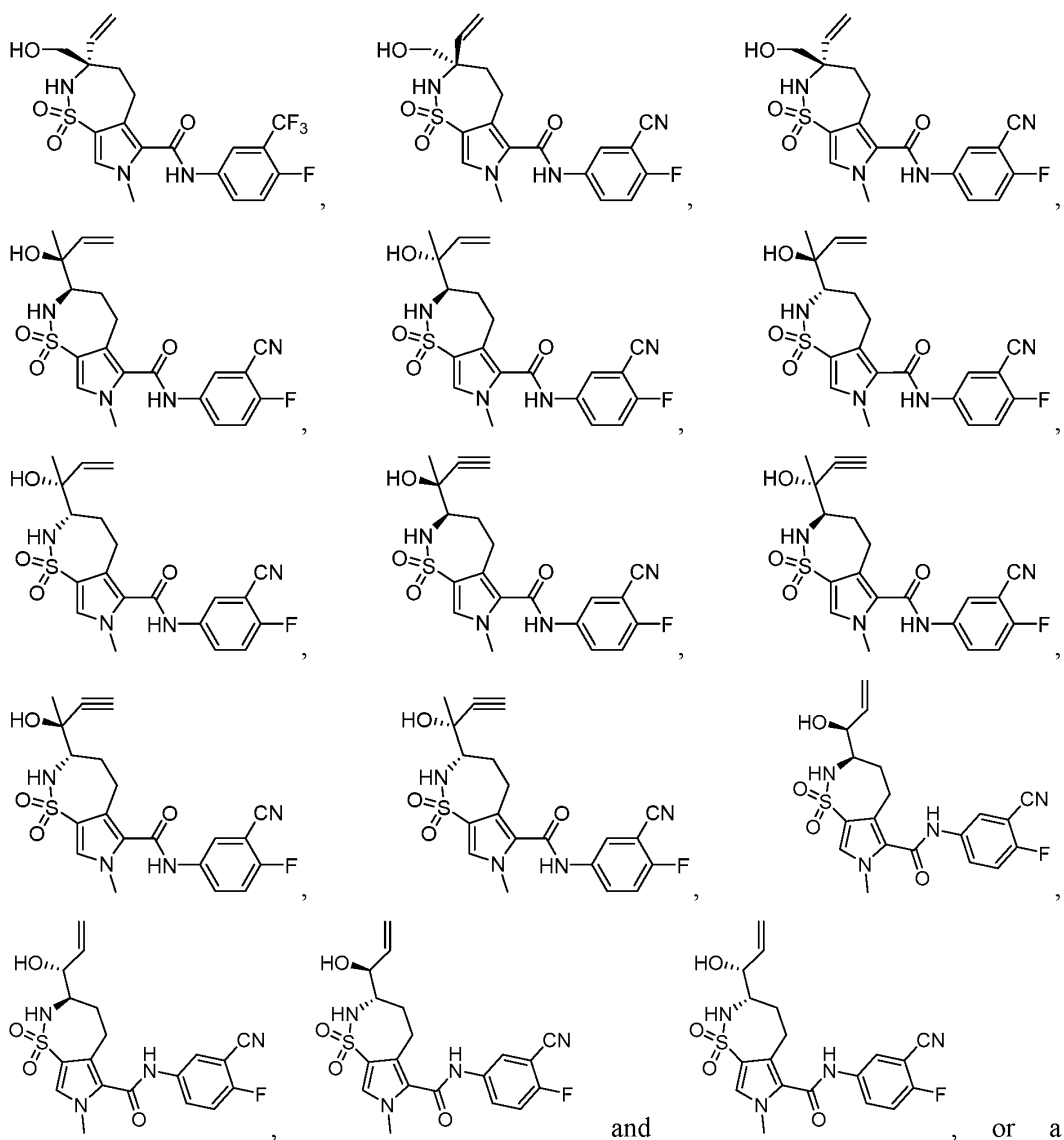




144. The compound of Claim 1, wherein the compound is selected from the group consisting of:

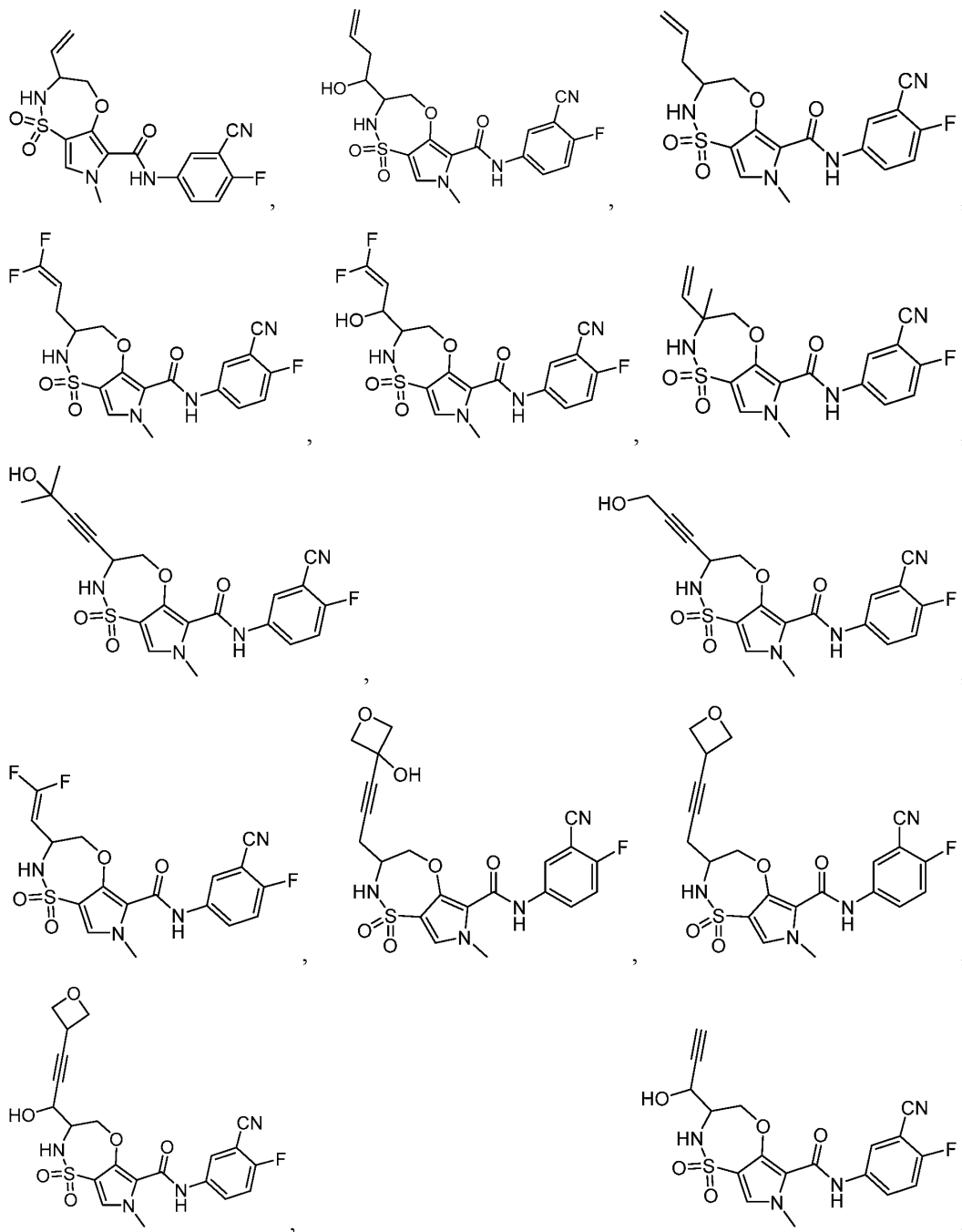


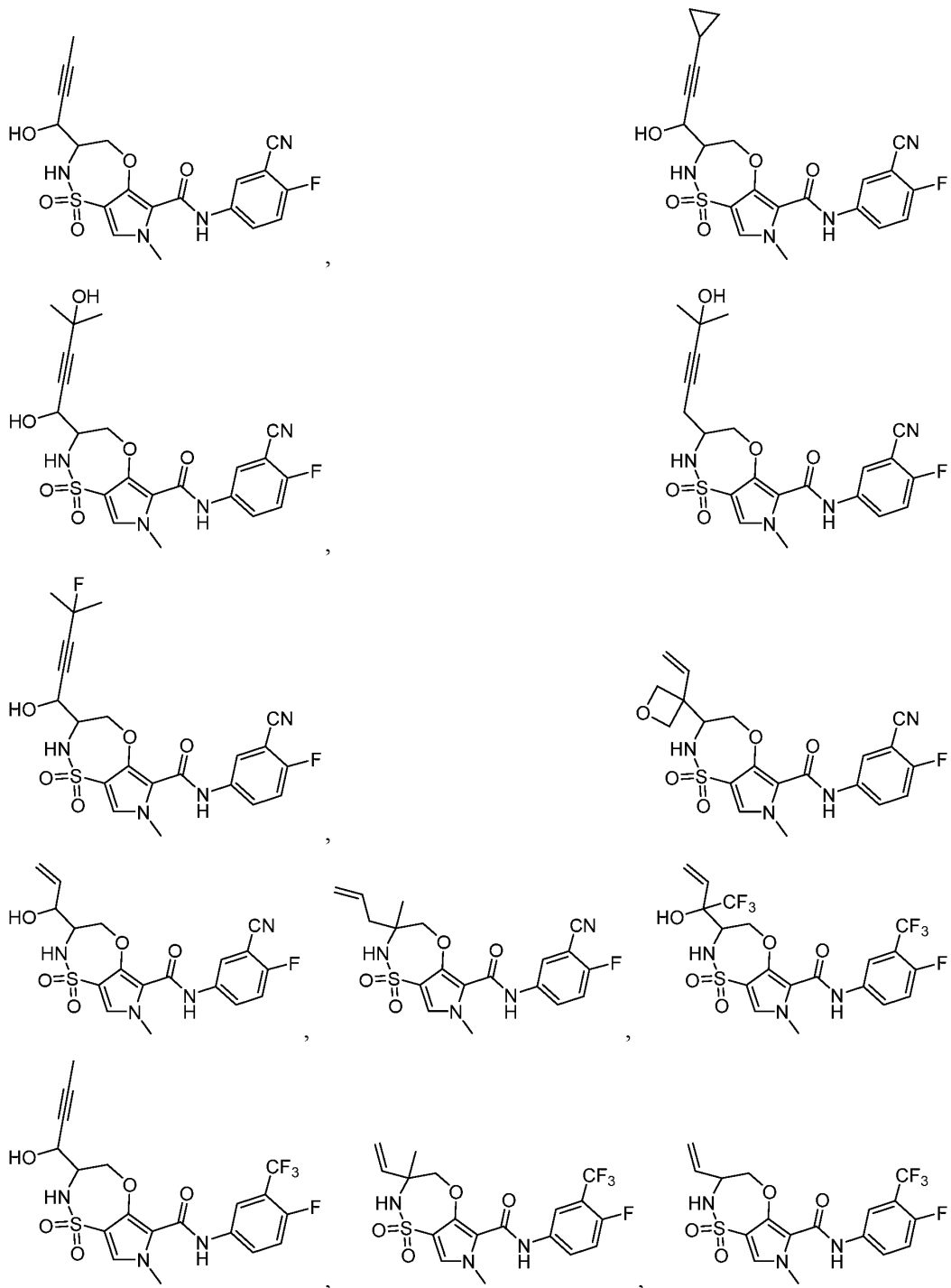


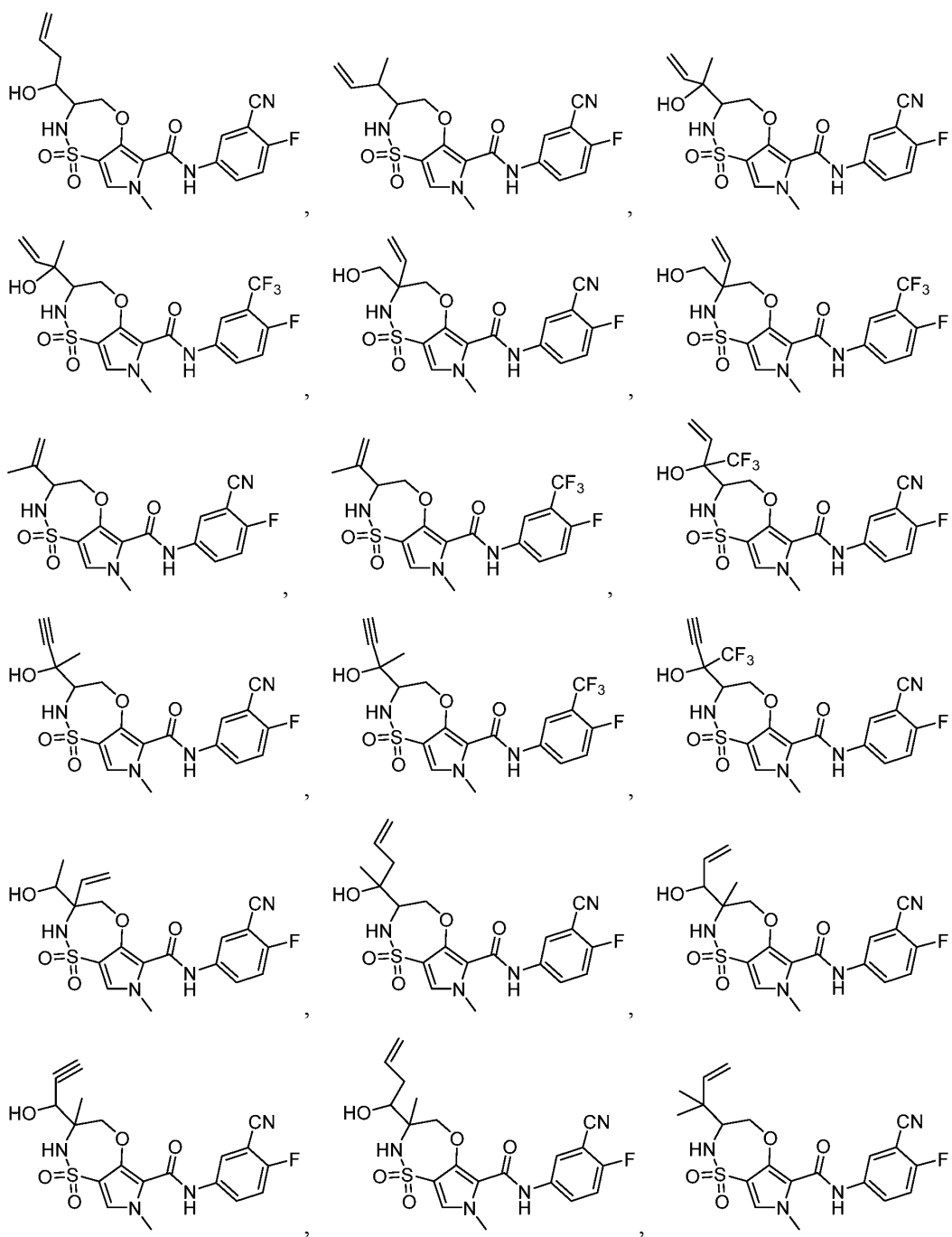


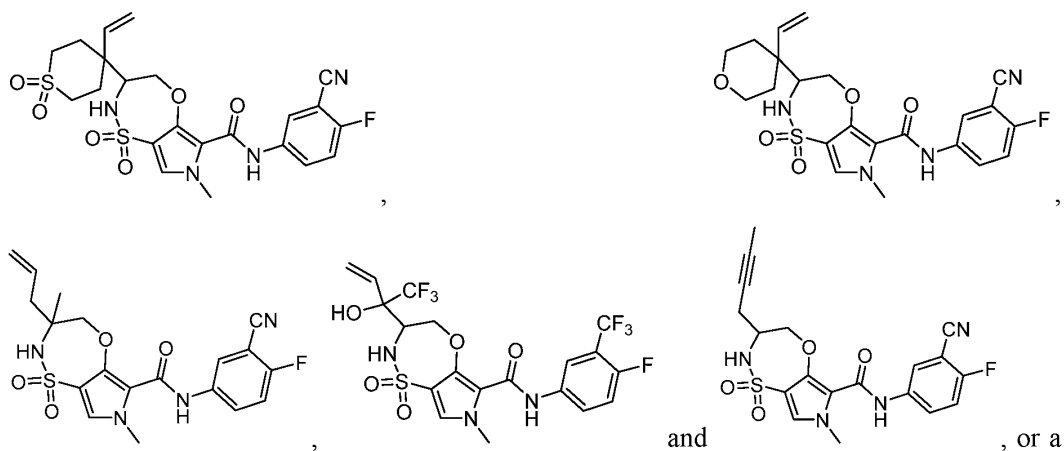
pharmaceutically acceptable salt of any of the foregoing.

145. The compound of Claim 85, wherein the compound is selected from the group consisting of:



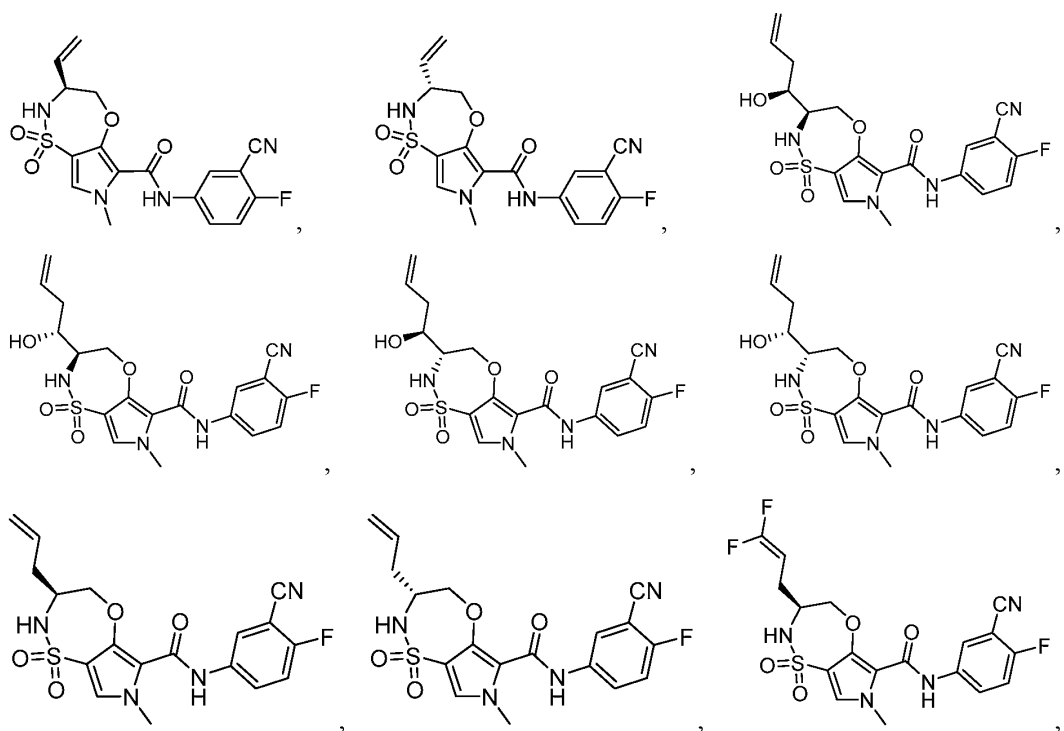


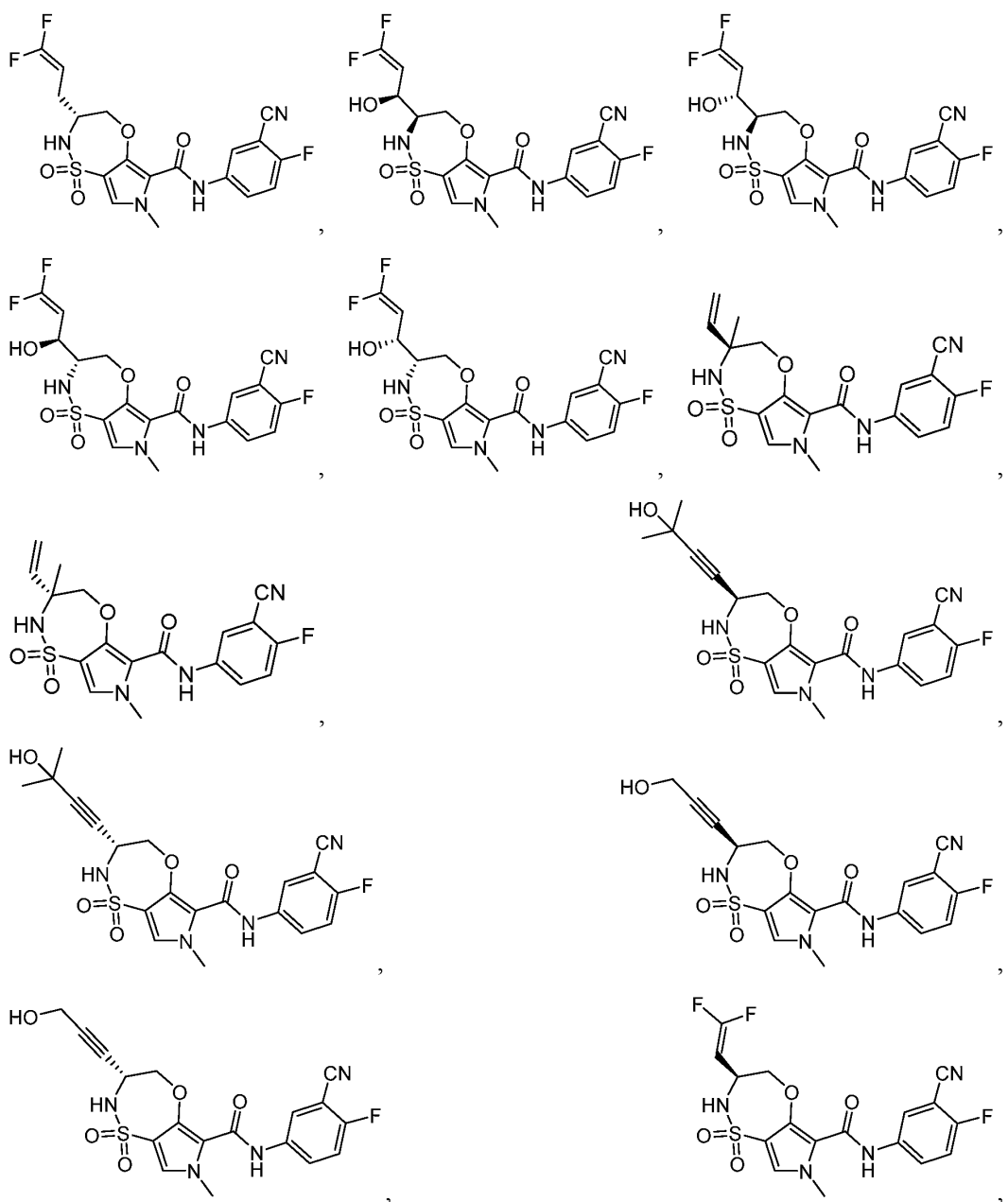


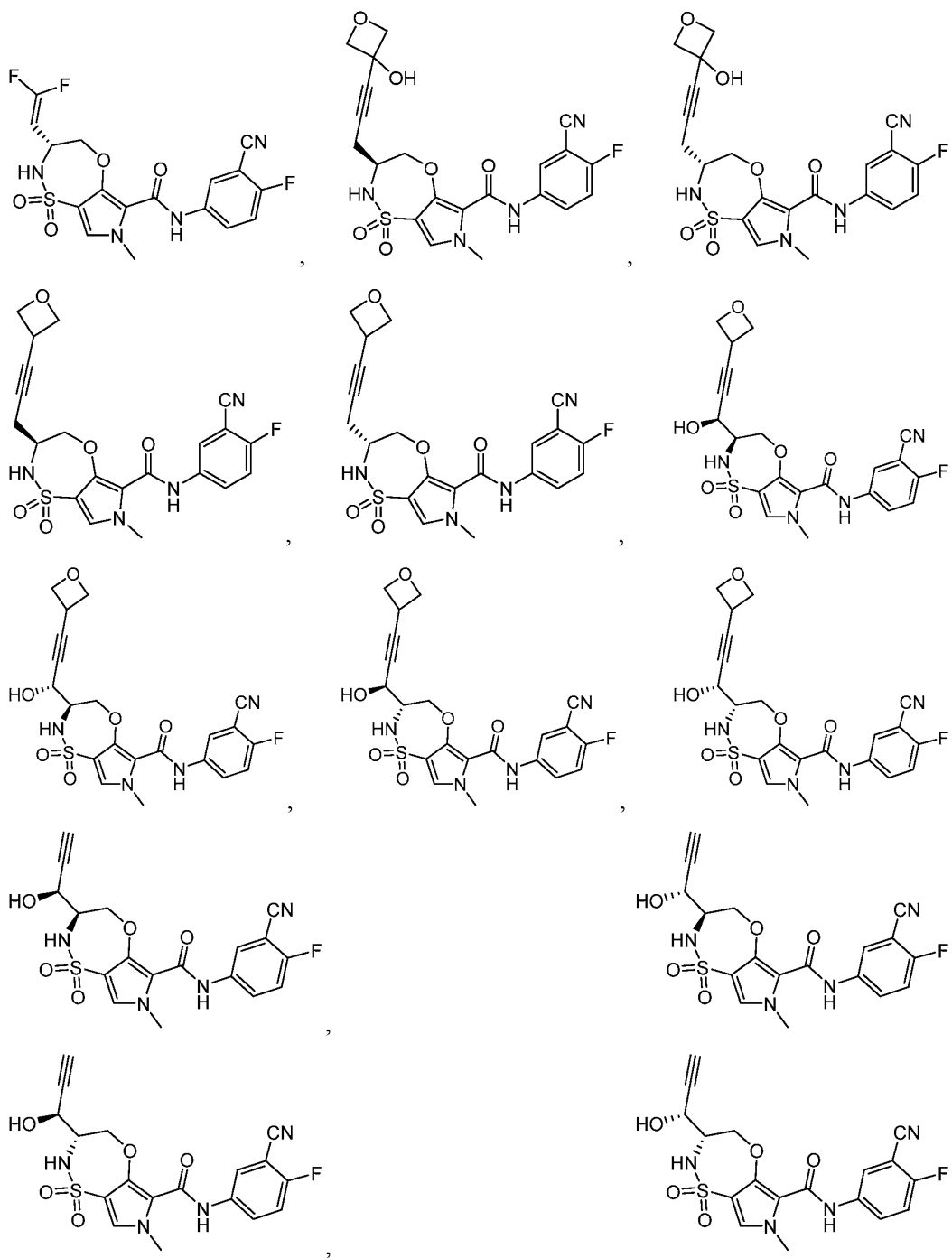


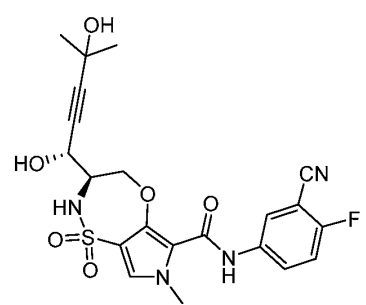
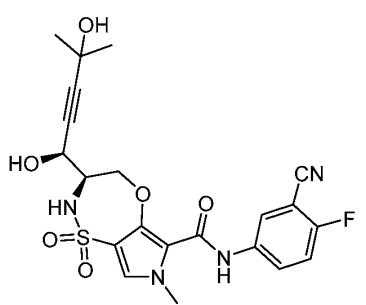
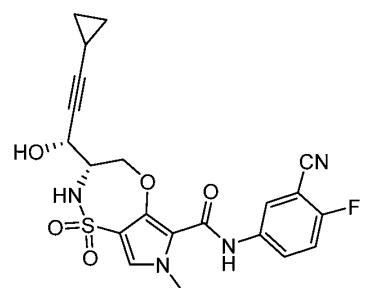
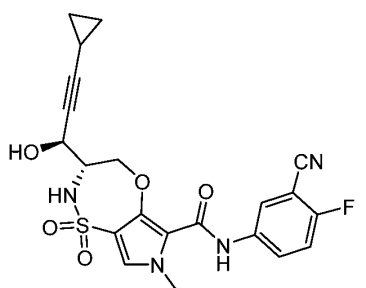
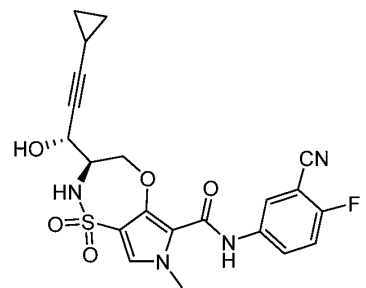
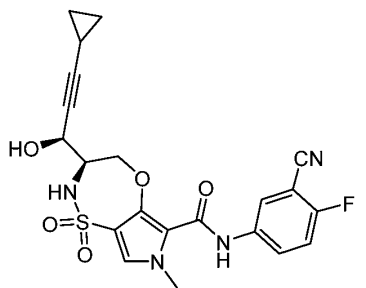
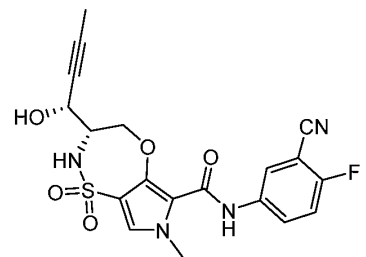
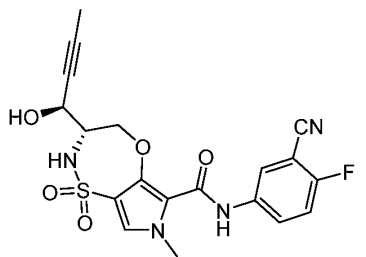
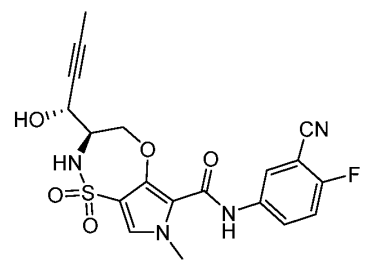
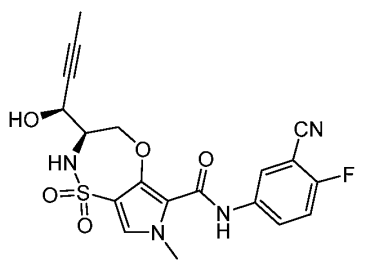
pharmaceutically acceptable salt of any of the foregoing.

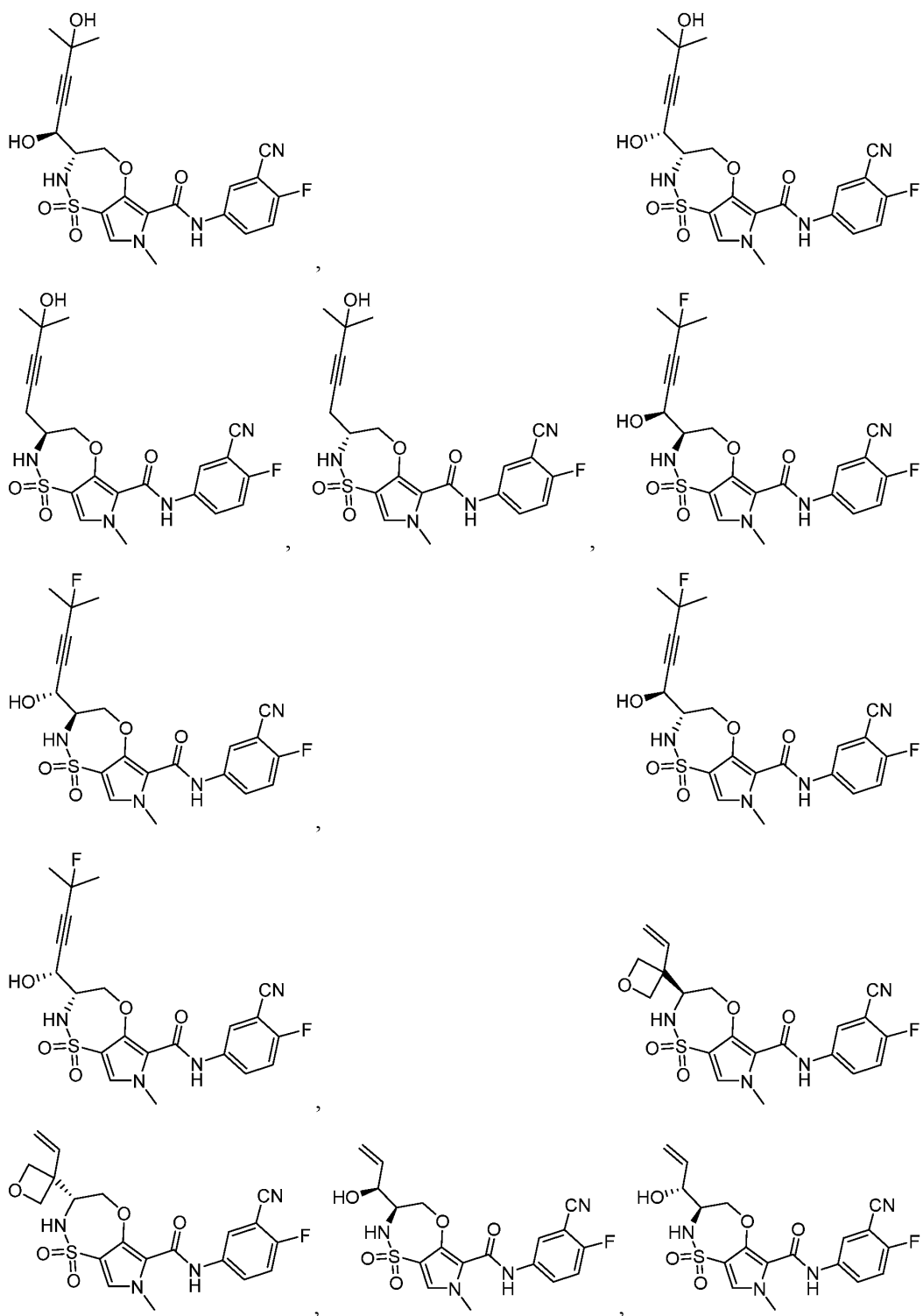
146. The compound of Claim 85, wherein the compound is selected from the group consisting of:

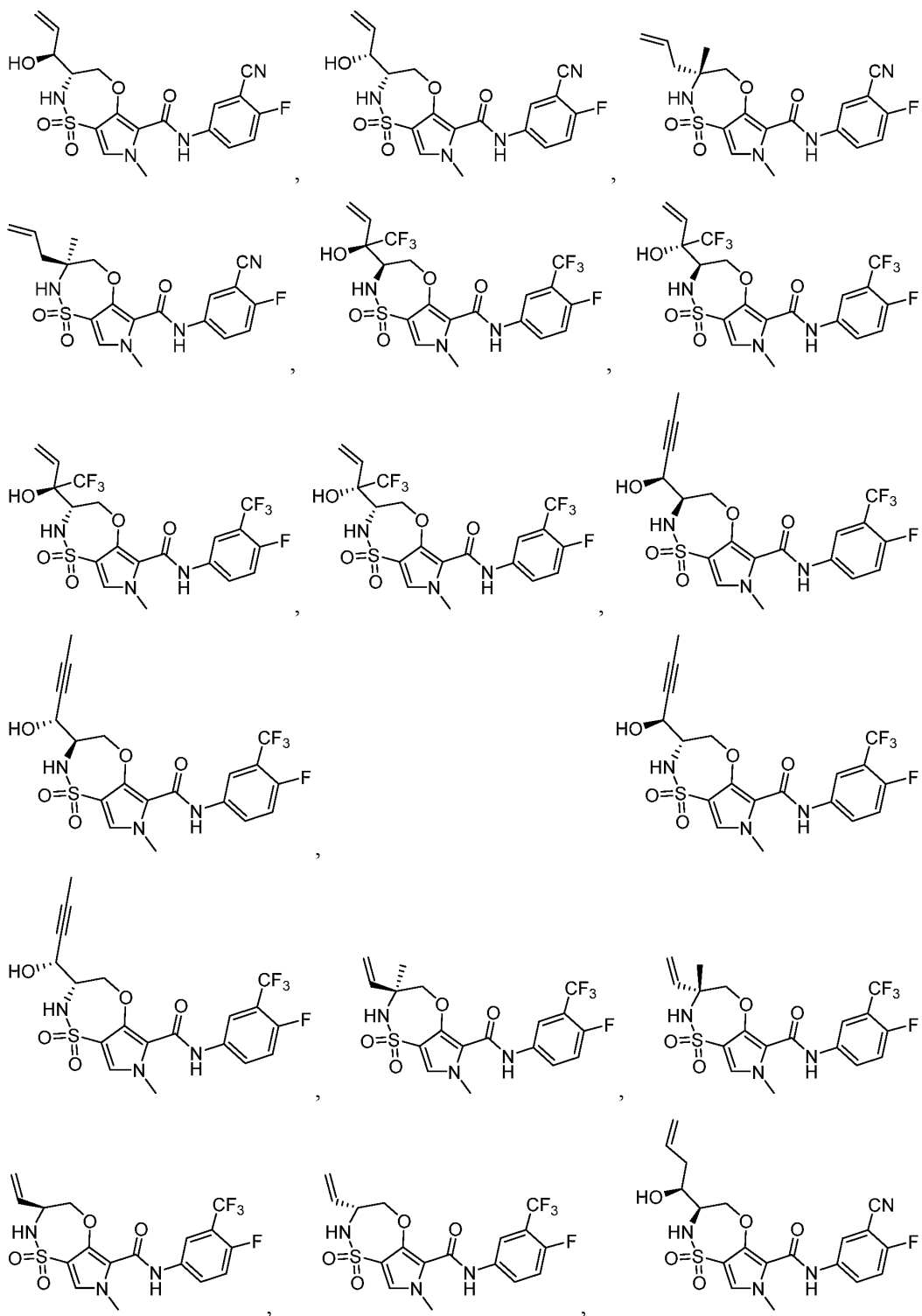


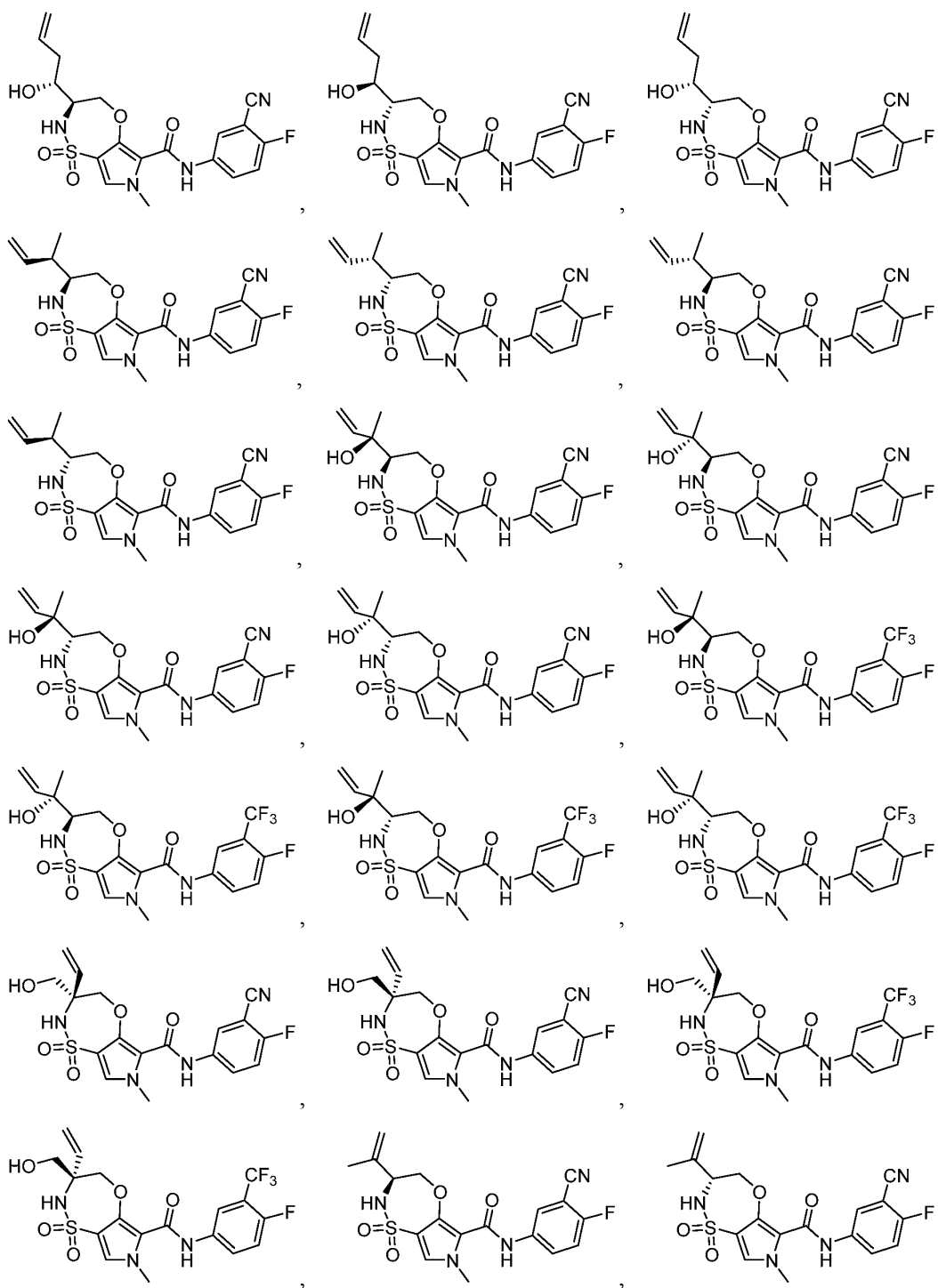


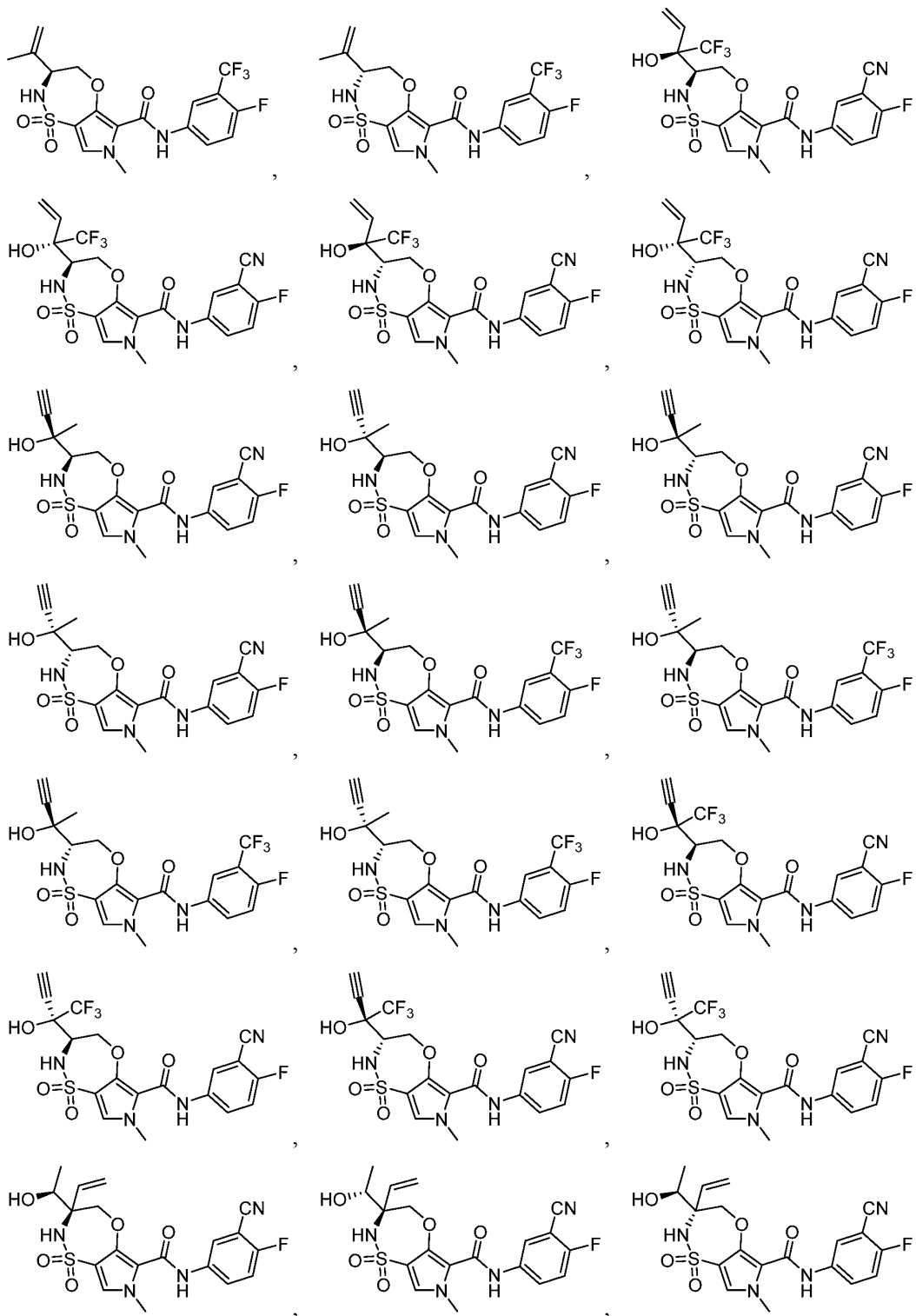


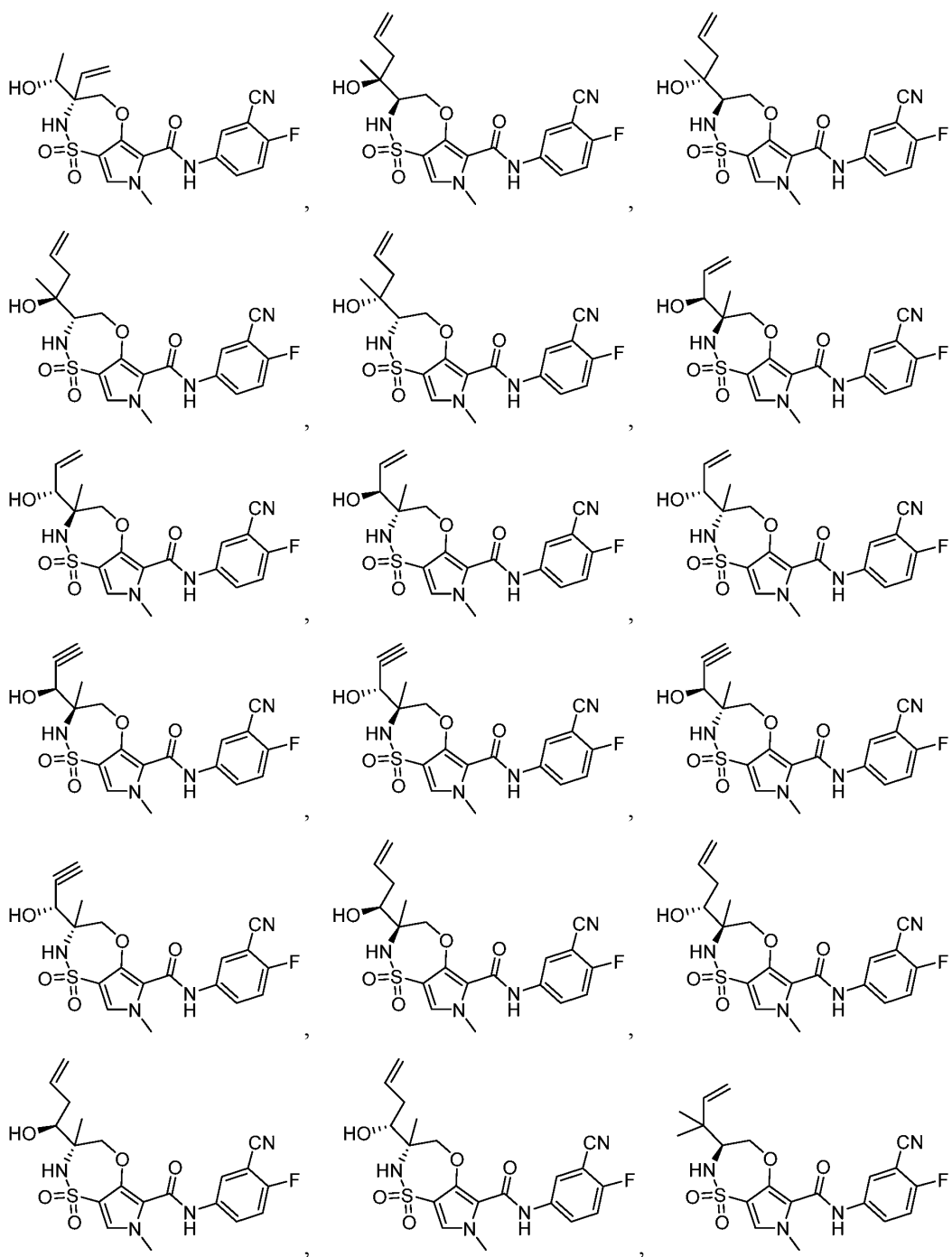


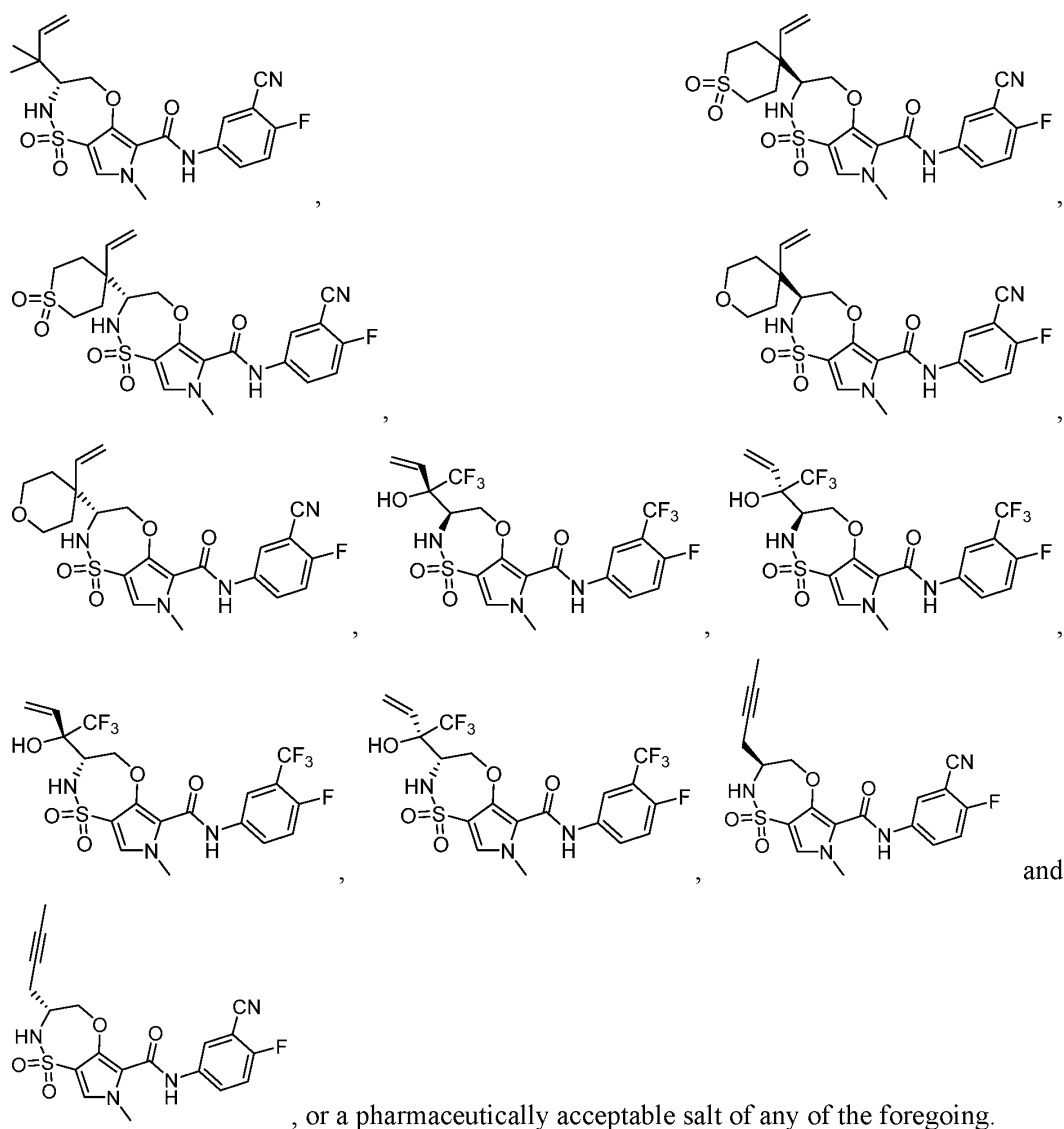












147. A pharmaceutical composition comprising an effective amount of a compound of any one of Claims 1-146, or a pharmaceutically acceptable salt thereof, and excipient.

148. A compound of any one of Claims 1-146 for use in treating hepatitis B.

149. A compound of any one of Claims 1-146 for use in treating hepatitis D.

150. The compound of any one of Claims 148-149, wherein the compound can be used in combination with an additional agent selected from the group consisting of an interferon, a nucleoside analog, a nucleotide analog, a capsid assembly modulator, a

sequence specific oligonucleotide, a nucleic acid polymer, an entry inhibitor and a small molecule immunomodulator.

151. The compound of Claim 150, wherein the additional agent selected from the group consisting of recombinant interferon alpha 2b, IFN- α , PEG-IFN- α -2a, lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide, tenofovir disoproxil, JNJ-6379, GLS4, ABI-HO731, JNJ-440, NZ-4, RG7907, AB-423, AB-506, ABI-H2158, REP 2139 and REP 2165.

152. A method for treating hepatitis B in a subject comprising administering to the subject in need thereof an effective amount of a compound of any one of Claims 1-146 suffering from hepatitis B.

153. A method for treating hepatitis D in a subject comprising administering to the subject in need thereof an effective amount of a compound of any one of Claims 1-146 suffering from hepatitis D.

154. The method of any one of Claims 152-153, further comprising administering an additional agent selected from the group consisting of an interferon, a nucleoside analog, a nucleotide analog, a capsid assembly modulator, a sequence specific oligonucleotide, a nucleic acid polymer, an entry inhibitor and a small molecule immunomodulator.

155. The method of Claim 154, wherein the additional agent selected from the group consisting of recombinant interferon alpha 2b, IFN- α , PEG-IFN- α -2a, lamivudine, telbivudine, adefovir dipivoxil, clevudine, entecavir, tenofovir alafenamide, tenofovir disoproxil, JNJ-6379, GLS4, ABI-HO731, JNJ-440, NZ-4, RG7907, AB-423, AB-506, ABI-H2158, REP 2139 and REP 2165.