

HETEROCYCLIC COMPOUND, INTERMEDIATE, PREPARATION METHOD THEREFOR AND APPLICATION THEREOF

The present application claims priority to the Chinese Patent Application CN201811642319 filed on December 29, 2018, the Chinese Patent Application CN201910440214.3 filed on May 24, 2019, and the Chinese Patent Application CN201911016158.7 filed on October 24, 2019, which are incorporated herein by reference in their entirety.

TECHNICAL FIELD

The present invention relates to a heterocyclic compound, an intermediate, and a preparation method and use thereof.

BACKGROUND

ATP receptors are classified into two main families, the P2Y-purinoreceptors and P2X-purinoreceptors, according to the molecular structure, transduction mechanism and pharmacological properties. The P2X-purinoreceptors are a family of ATP-gated cation channels, and several subtypes have been cloned, including six homomeric receptors (P2X1, P2X2, P2X3, P2X4, P2X5 and P2X7) and three heteromeric receptors (P2X2/3, P2X4/6 and P2X1/5). The P2X3 receptor has been found to be expressed particularly in the primary afferent nerve fibers of "hollow viscera", such as the lower urinary tract and respiratory tract.

Cough is the main symptom of respiratory diseases, and is observed in 70-80% of patients in outpatients of the respiratory department. With increasing prevalence of COPD, IPF, etc., there's also an increasing demand for treatment of cough as it is the main manifestation of most respiratory tract diseases. Cough, as a defensive nervous reflex of the body, is beneficial to the removal of respiratory secretions and harmful factors. However, frequent and violent cough will seriously affect the work, life and social activities of patients.

Few P2X3 antagonists specifically targeting cough have been developed, and one project that advances rapidly at present is the Roche AF-219 project, which, in the newly completed phase II clinical trial, shows a relatively good efficacy for refractory cough but poses serious dysgeusia.

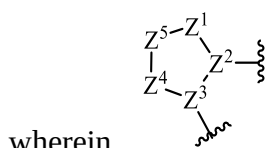
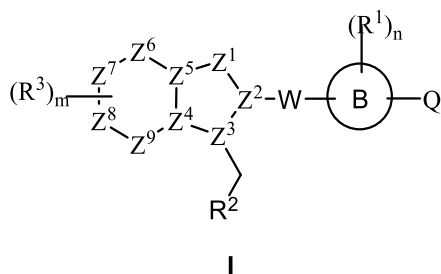
At present, no drug is available on the market that can treat various conditions including chronic cough via the P2X3-inhibiting route. Therefore, the development of new compounds capable of inhibiting the activity of P2X3 is of great significance for treating diseases.

SUMMARY

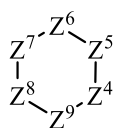
The present invention aims to solve the technical problem of the insufficiency of P2X3 antagonist in the prior art, and thus provides a heterocyclic compound, an intermediate, and a preparation method and use thereof. The heterocyclic compound disclosed herein has high antagonistic activity against P2X3, good selectivity, low toxicity, good metabolic stability and small influence on taste.

The present invention solves the above-mentioned problem by the following technical embodiments.

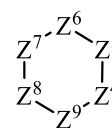
The present invention provides a heterocyclic compound of formula I, or a stereoisomer, a geometric isomer, a tautomer, a nitrogen oxide, a hydrate, a solvate, a metabolite, an ester, a pharmaceutically acceptable salt or a prodrug thereof,



wherein, is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S; and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl; wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S; the 5-6 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S; and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



$Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8$ and Z^9 independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z^6 or Z^9 represents a single bond;

m is 1, 2, 3 or 4;

R^3 is H, halogen, -OH, a C_1 - C_6 haloalkyl, -CN or a C_1 - C_6 alkyl; when m is not 1, R^3 is independently the same or different; or when two adjacent R^3 are present, they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkyl, a C_3 - C_6 cycloalkenyl, a C_3 - C_9 heterocycloalkyl, a C_3 - C_9 heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl; wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C_3 - C_9 heterocycloalkyl, wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

n is 1, 2, 3 or 4;

R^1 is independently H, halogen, -OH, -CN, an unsubstituted or substituted C_1 - C_6 alkyl, or an unsubstituted or substituted C_1 - C_6 alkyl-O-; wherein the substituted C_1 - C_6 alkyl or the substituted C_1 - C_6 alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); when a plurality of substituents are present, they are the same or different; and when n is not 1, R^1 is independently the same or different;

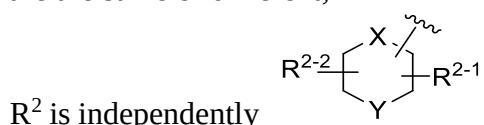
Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)- R^{1-1} , -C(=O)- R^{1-2} , -O-C(=O)-N(R^{1-3} R^{1-4}), -C(=O)-N(R^{1-5} R^{1-6}), -C(=NH)-N(R^{1-7} R^{1-8}), -S(=O)₂-N(R^{1-9} R^{1-10}), -S(=O)-N(R^{1-11} R^{1-12}), -O-C(=O)- R^{1-13} , -N(R^{1-14} R^{1-15}), -NH-S(=O)₂- R^{1-16} , an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_1 - C_6 alkyl-O-, an unsubstituted or substituted C_2 - C_6 alkenyl, an unsubstituted or substituted C_2 - C_6 alkenyl-O-, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, an unsubstituted or substituted C_6 - C_{10} aryl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C_1 - C_6 alkyl, the substituted C_1 - C_6 alkyl-O-, the substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6 alkenyl-O-, the substituted C_3 - C_{10} cycloalkyl, the substituted C_6 - C_{10} aryl, the substituted C_3 - C_9 heterocycloalkyl, the substituted C_3 - C_9 heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or

more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₂-C₆ alkynyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₂-C₆ alkynyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl, the substituted phenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -COOCH₃, -NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d; wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different; or

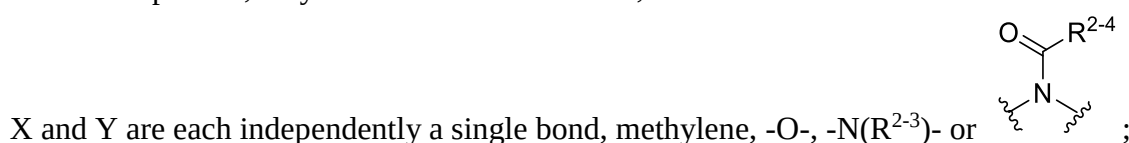
R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C₃-C₉ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O,

S and N in addition to the existing N atom, the 5-10 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b; wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;



5-6 membered heteroaryl; wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CN, an unsubstituted or halogen-substituted C₁-C₆ alkyl, an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-, -C(=O)R²⁻⁷ or -C(=O)N(R²⁻⁸R²⁻⁹); wherein when a plurality of substituents are present, they are the same or different;

R²⁻¹ and R²⁻² are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;



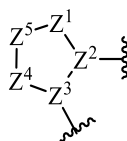
R²⁻³, R²⁻⁴, R²⁻⁵ and R²⁻⁶ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal

hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

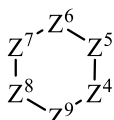
R^{2-7} is independently H, halogen, -OH, -CN, an unsubstituted or halogen-substituted C_1 - C_6 alkyl, or an unsubstituted or halogen-substituted C_1 - C_6 alkyl-O-; and

5 R^{2-8} and R^{2-9} are each independently H, or an unsubstituted or halogen-substituted C_1 - C_6 alkyl.

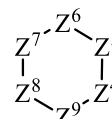
In the present invention, some substituents in the heterocyclic compound of formula I may have the definitions below, and the substituents which are not mentioned have the definitions as described in any one of the above embodiments.



10 In a preferred embodiment of the present invention, is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl; wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



15 is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl; wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



20 $Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8$ and Z^9 independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z^6 or Z^9 represents a single bond;

m is 1, 2, 3 or 4;

25 R^3 is H, halogen, -OH, a C_1 - C_6 haloalkyl, -CN or a C_1 - C_6 alkyl; when m is not 1, R^3 is independently the same or different; or when two adjacent R^3 are present, they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkyl, a C_3 - C_6 cycloalkenyl, a C_3 - C_9 heterocycloalkyl, a C_3 - C_9 heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl; wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N,

and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C₃-C₉ heterocycloalkyl; wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

n is 1, 2, 3 or 4;

R¹ is independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); when a plurality of substituents are present, they are the same or different; and when n is not 1, R¹ is independently the same or different;

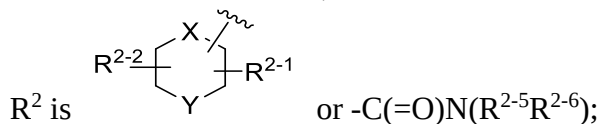
Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₃-C₁₀ cycloalkyl, the substituted C₆-C₁₀ aryl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a; wherein when a plurality of R^a are present, they are the same or different, and each R^a is

halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

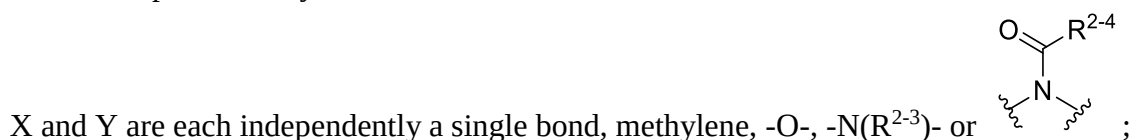
R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₂-C₆ alkynyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₂-C₆ alkynyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl, the substituted phenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -COOCH₃, -NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d, wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different; or

R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C₃-C₉ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the 5-10 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, and the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e.,

two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b, wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;



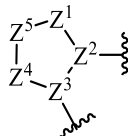
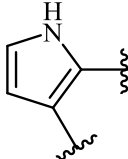
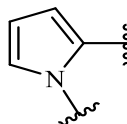
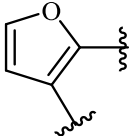
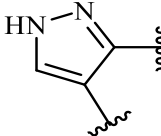
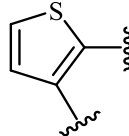
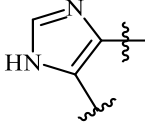
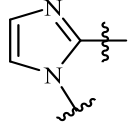
R²⁻¹ and R²⁻² are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

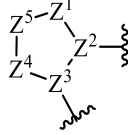
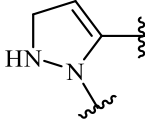


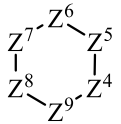
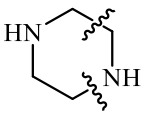
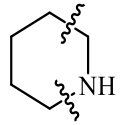
R²⁻³, R²⁻⁴, R²⁻⁵ and R²⁻⁶ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different.

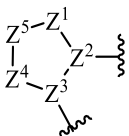
In the heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof disclosed herein, some substituents may have the definitions below, and the substituents which are not mentioned have the definitions as described in any one of the embodiments.

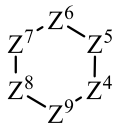
In a preferred embodiment of the present invention, Z⁶ is a single bond and Z⁹ is NH, O or S; or, Z⁹ is a single bond and Z⁶ is NH, O or S.

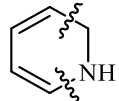
In a preferred embodiment of the present invention, when  is a 5 membered heteroaryl, the 5 membered heteroaryl is pyrrolyl (e.g.,  or ) , furanyl (e.g., ) , pyrazolyl (e.g., ) , thienyl (e.g., ) or imidazolyl (e.g.,  or ) . (In the structural formula, the right-side bond is linked to W)

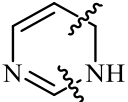
5 In a preferred embodiment of the present invention, when  is a 5 membered heterocycloalkenyl, the 5 membered heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S, and is preferably 2,3-dihydro-1H-pyrazolyl (e.g., ) . (In the structural formula, the right-side bond is linked to W)

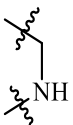
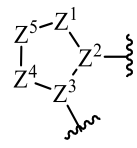
10 In a preferred embodiment of the present invention, when  is a 5-6 membered heterocycloalkyl, the 5-6 membered heterocycloalkyl is a 6 membered heterocycloalkyl; wherein the 6 membered heterocycloalkyl contains 1-3 N heteroatoms; and the 5-6 membered heterocycloalkyl is preferably piperazinyl (e.g., ) or piperidinyl (e.g., ) . (In

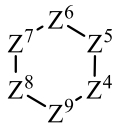
the structural formula, the right-side carbon-nitrogen bond  represents the position fusing to )

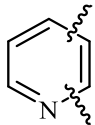
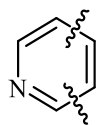
In a preferred embodiment of the present invention, when  is a 5-6 membered heterocycloalkenyl, the 5-6 membered heterocycloalkenyl is a 6 membered heterocycloalkenyl; wherein the 6 membered heterocycloalkenyl contains 1-3 N heteroatoms; and the 5-6 membered

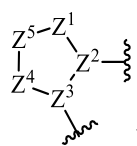
heterocycloalkenyl is preferably 1,2-dihydropyridinyl (e.g., ) or 1,6-

5 dihydropyrimidinyl (e.g., )). (In the structural formula, the right-side carbon-nitrogen

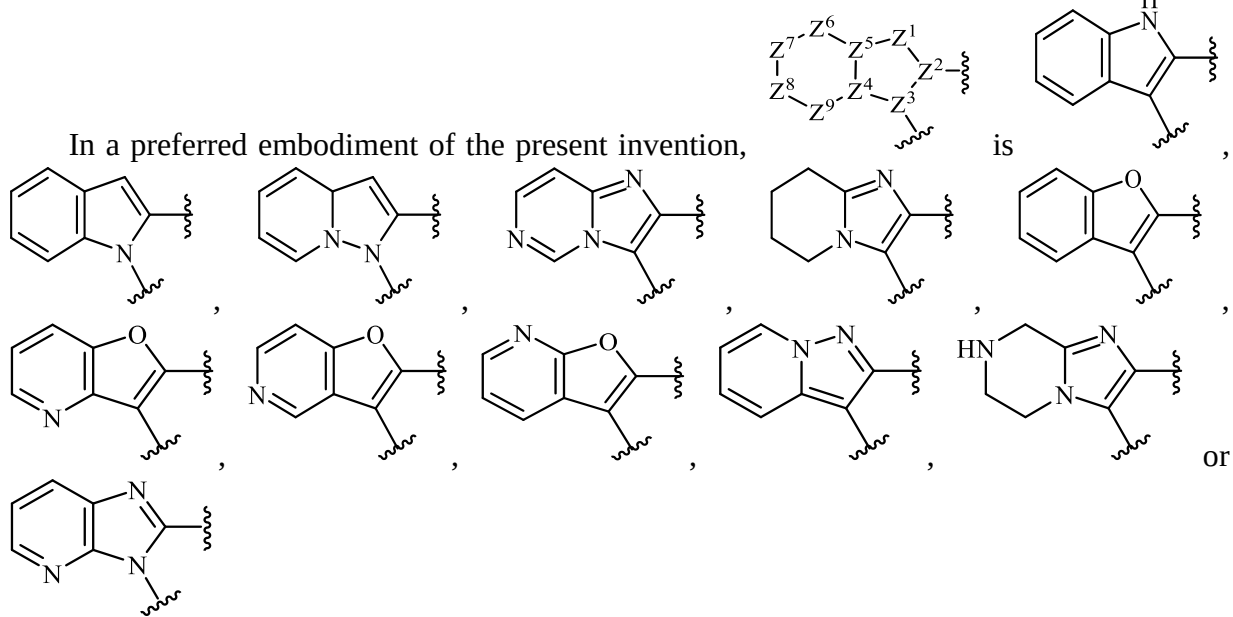
bond  represents the position fusing to )

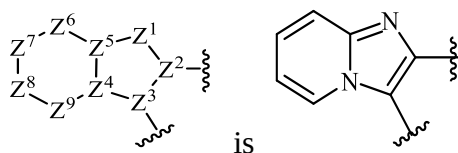
In a preferred embodiment of the present invention, when  is a 5-6 membered heteroaryl, the 5-6 membered heteroaryl is a 6 membered heteroaryl; wherein the 6 membered heteroaryl contains 1-3 N heteroatoms (e.g., pyridinyl or pyrimidinyl); and the 5-6 membered

10 heteroaryl is preferably pyridinyl (e.g.,  or )). (In the structural formula, the

right-side carbon-carbon double bond  represents the position fusing to )

In a preferred embodiment of the present invention,





In a preferred embodiment of the present invention,

In a preferred embodiment of the present invention, m is 1 or 2;

In a preferred embodiment of the present invention, Z⁷ is substituted with R³, Z⁸ is unsubstituted or substituted with R³, Z⁶ and Z⁹ are unsubstituted; when a plurality of R³ are present, they are the same or different; and preferably, Z⁷ is substituted with R³, and Z⁶, Z⁸ and Z⁹ are unsubstituted.

In a preferred embodiment of the present invention, when R³ is halogen, the halogen is fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R³ is a C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl), the C₁-C₆ alkyl is a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl.

In a preferred embodiment of the present invention, when R³ is a C₁-C₆ haloalkyl, the halogen is fluorine, chlorine, bromine or iodine, and preferably chlorine.

In a preferred embodiment of the present invention, when R³ is a C₁-C₆ haloalkyl, it contains one or more (e.g., 1, 2, 3, 4 or 5) halogens, and preferably 1, 2 or 3 halogens.

In a preferred embodiment of the present invention, R³ is -CF₃, -CH₂F or -CHF₂.

In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₆ cycloalkyl, the C₃-C₆ cycloalkyl is cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl.

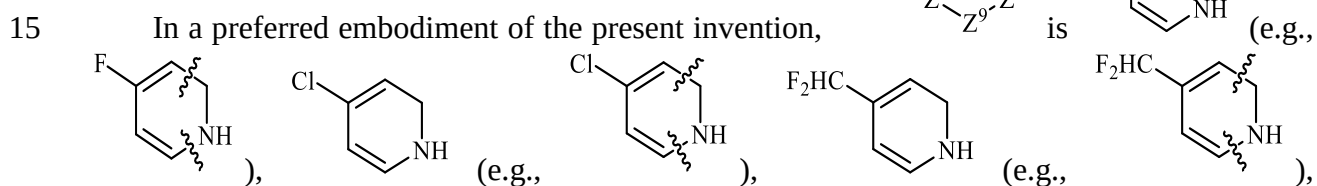
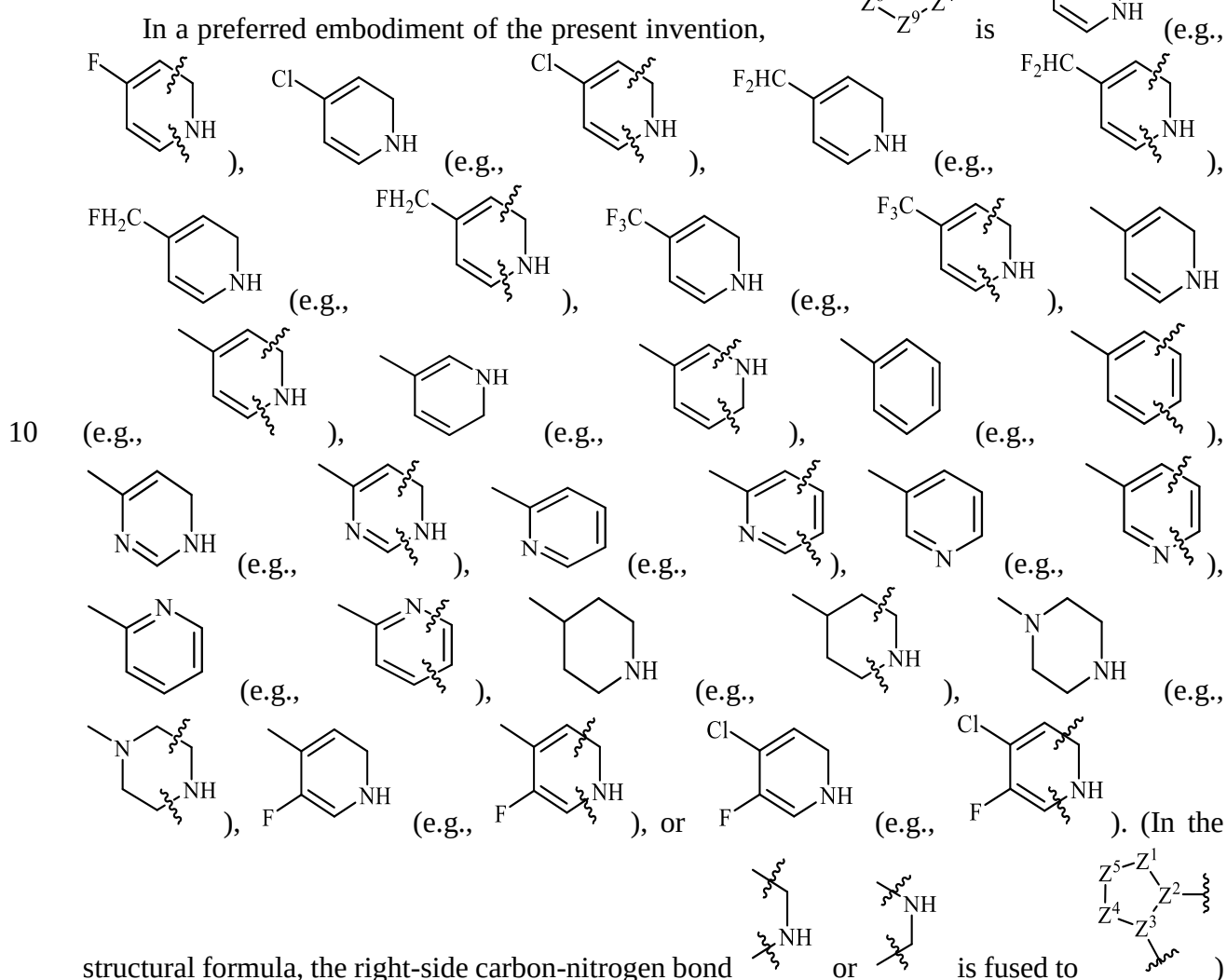
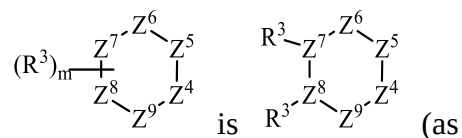
In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₆ cycloalkenyl, the C₃-C₆ cycloalkenyl is cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl or cyclohexadienyl.

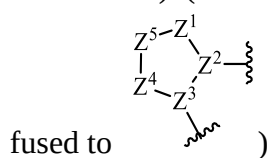
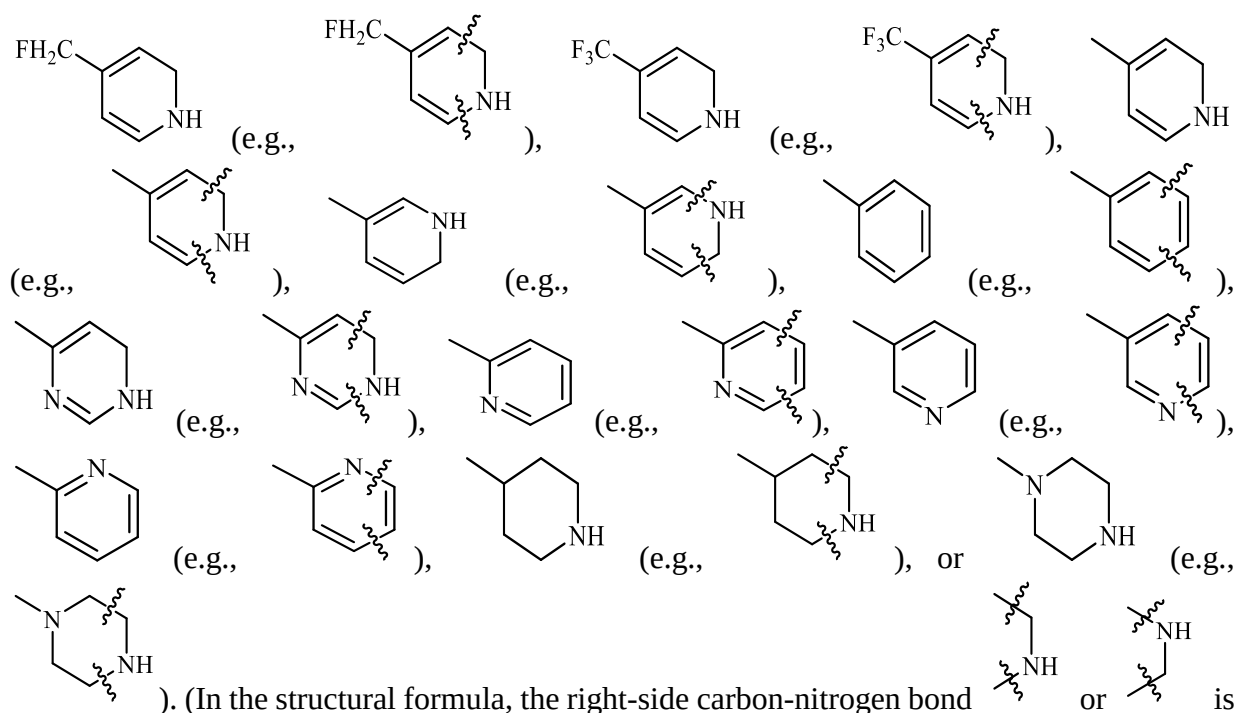
In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₄-C₅ heterocycloalkyl; wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl; wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S.

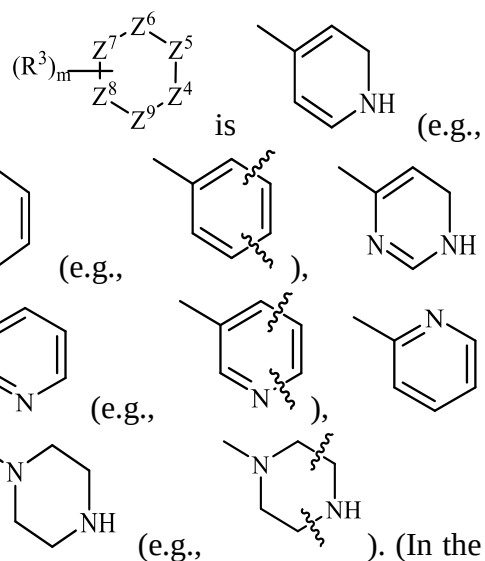
In a preferred embodiment of the present invention, when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a 5-6 membered heteroaryl, the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S.

5 In a preferred embodiment of the present invention, described above, R^3 is the same or different).

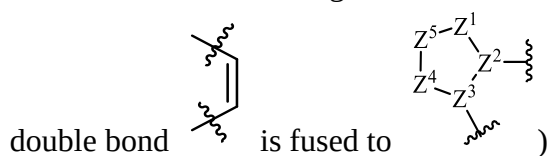




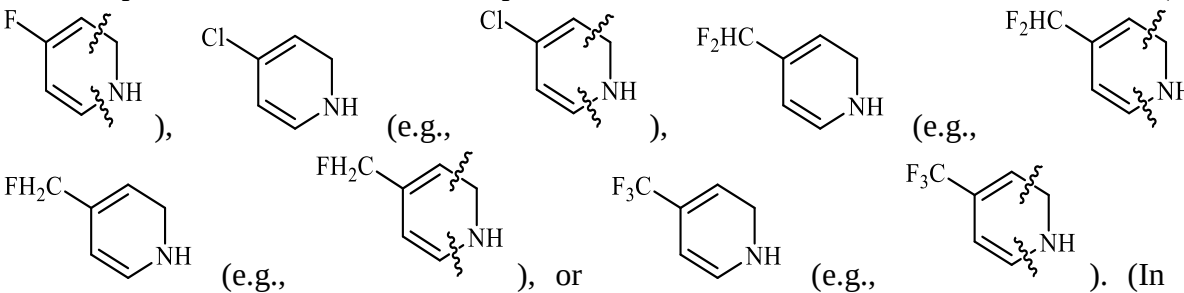
In a preferred embodiment of the present invention,

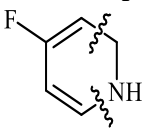
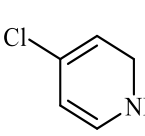
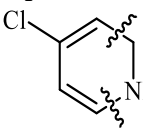
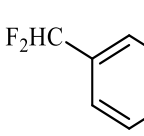
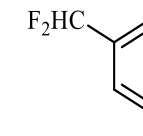


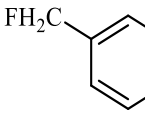
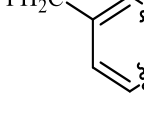
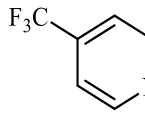
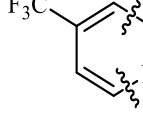
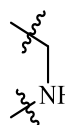
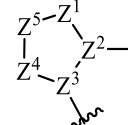
structural formula, the right-side carbon-nitrogen bond



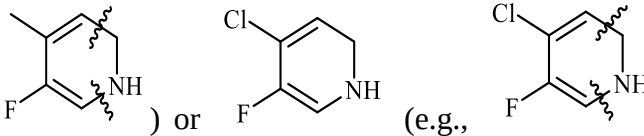
In a preferred embodiment of the present invention,

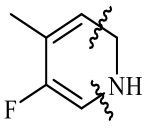
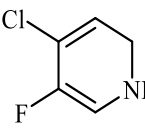
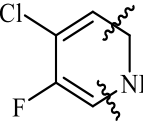
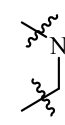
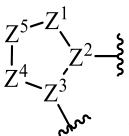


(e.g., ) , (e.g., ) , (e.g., ) , (e.g., ) , (e.g., ) ,

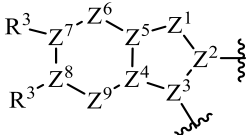
(e.g., ) , (e.g., ) , or (e.g., ) , (e.g., ) . (In the structural formula, the right-side carbon-nitrogen bond  or  is fused to )

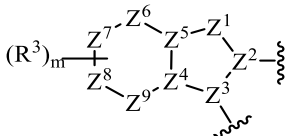
5 In a preferred embodiment of the present invention,



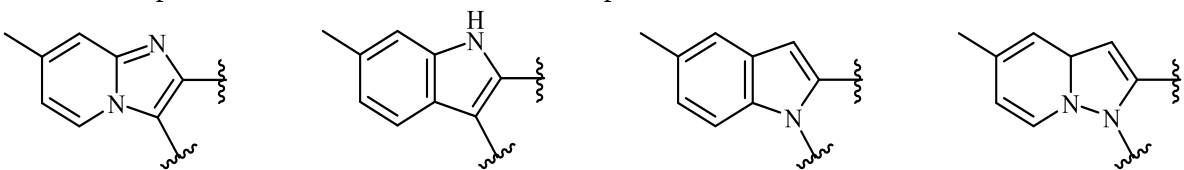
(e.g., ) or (e.g., ) , (e.g., ) . (In the structural formula, the right-side carbon-nitrogen bond  or  or the carbon-carbon double bond  is fused to )

10 In a preferred embodiment of the present invention,

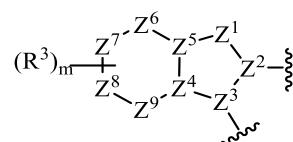
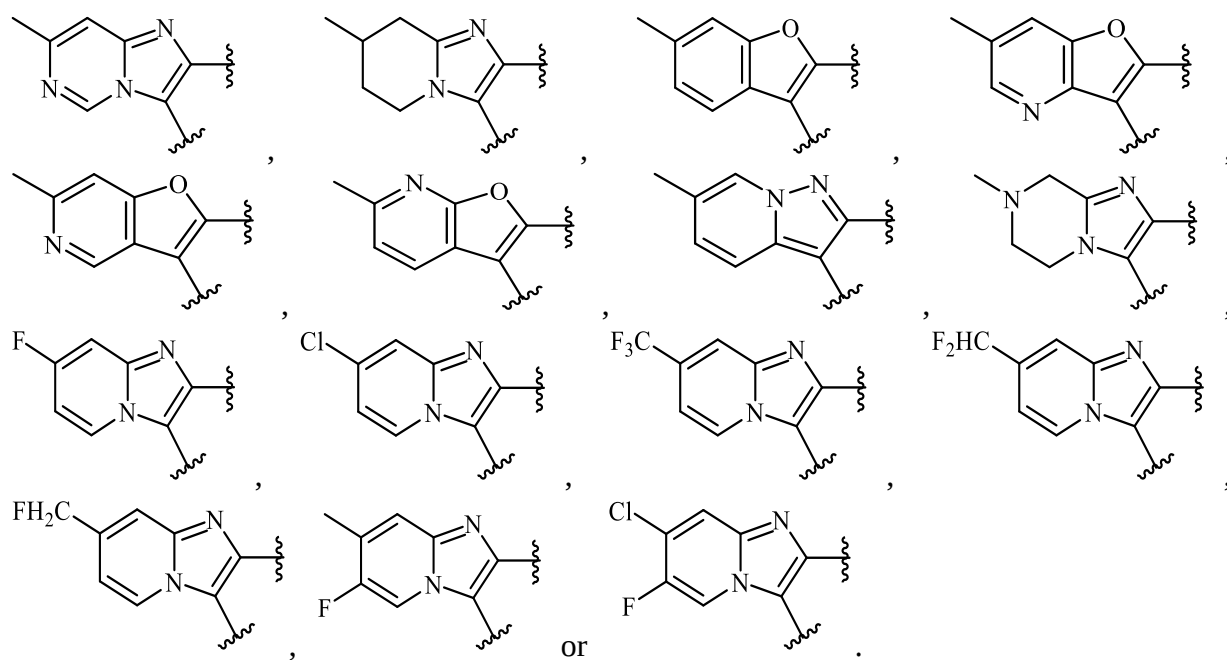


 is

In a preferred embodiment of the present invention,

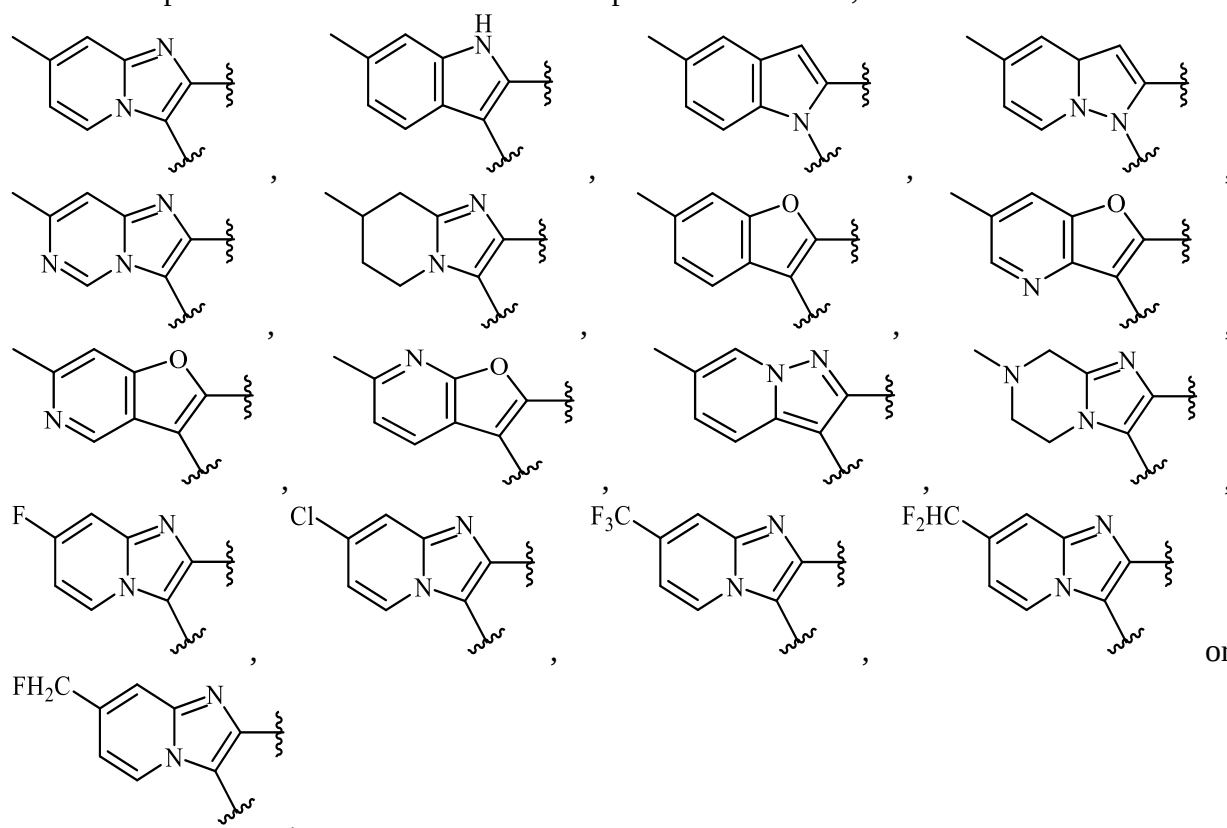


, , , ,

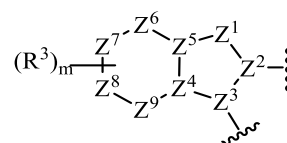


5 In a preferred embodiment of the present invention,

is

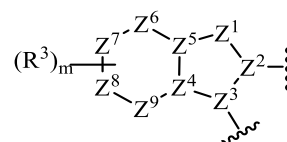
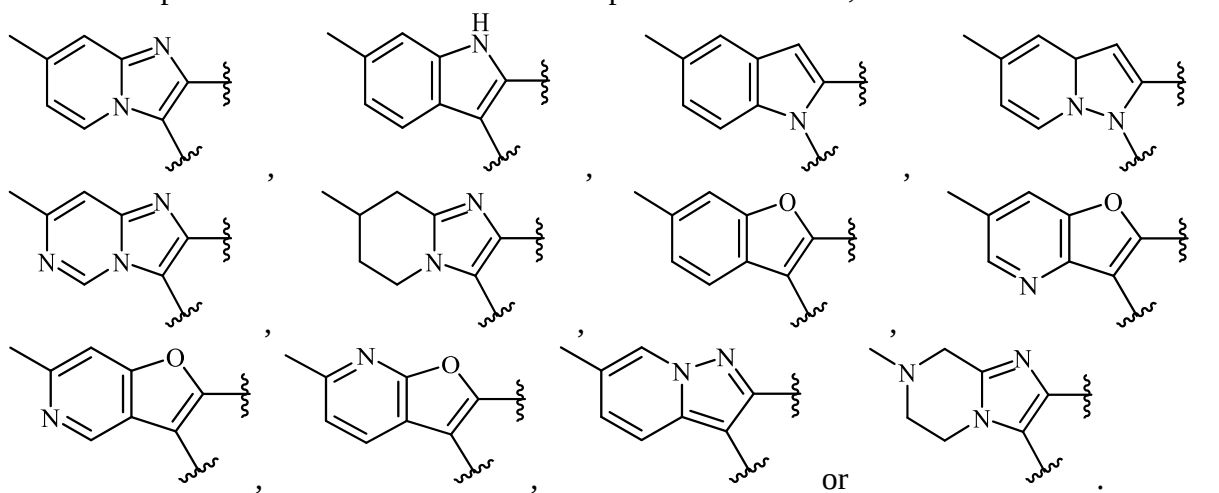


10



In a preferred embodiment of the present invention,

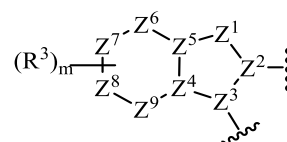
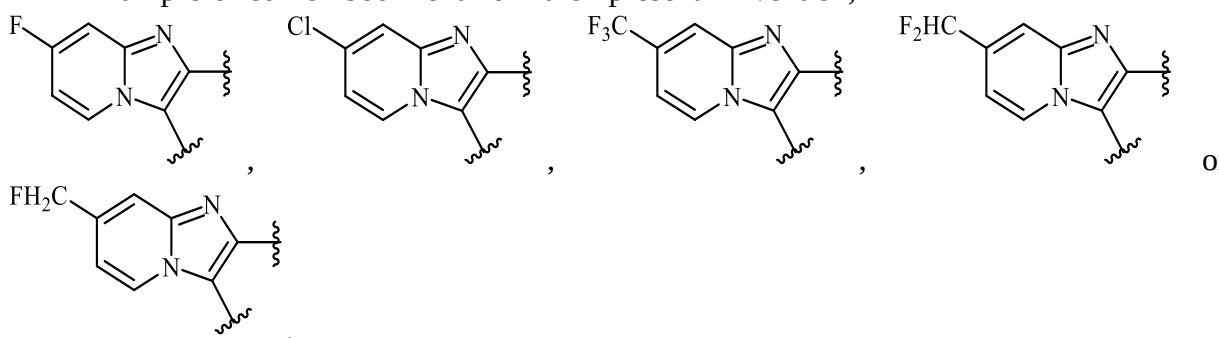
is



5

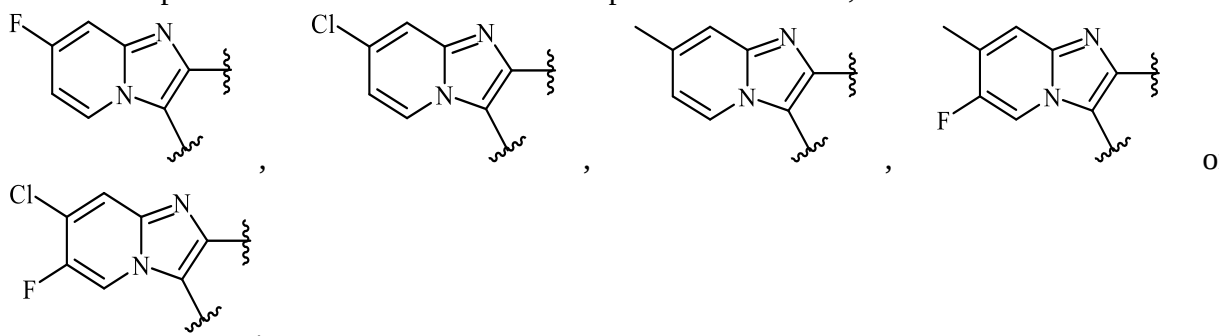
In a preferred embodiment of the present invention,

is



In a preferred embodiment of the present invention,

is



10

In a preferred embodiment of the present invention, n is 1 or 2.

In a preferred embodiment of the present invention, each R^1 is independently located at position α (the ortho-position of Z^2) or position β (the meta-position of Z^2) of the linkage between  and Z^2 .

In a preferred embodiment of the present invention, at least one R^1 is located at position α of the linkage between  and Z^2 ; and preferably, when n is 2, each R^1 is independently located at position α of the linkage between  and Z^2 .

In a preferred embodiment of the present invention, when R^1 is halogen, the halogen is fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R^1 is an unsubstituted or substituted C_1 - C_6 alkyl, or an unsubstituted or substituted C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl.

In a preferred embodiment of the present invention, when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, and the substituent is each independently halogen, the halogen is fluorine, chlorine, bromine or iodine.

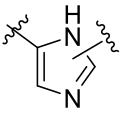
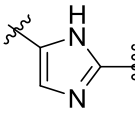
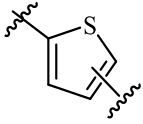
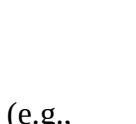
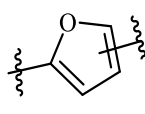
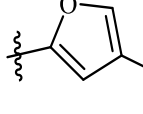
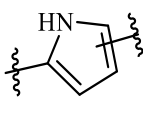
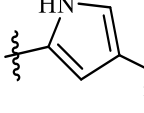
In a preferred embodiment of the present invention, when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, the substituent is independently a C_1 - C_6 alkyl or a C_1 - C_6 alkyl-O-, and the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the substituent is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl.

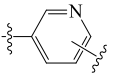
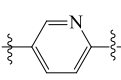
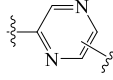
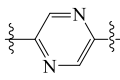
In a preferred embodiment of the present invention, when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, the number of the substituents is independently 1, 2 or 3.

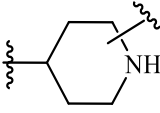
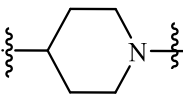
In a preferred embodiment of the present invention, when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O- and the substituents are each independently halogen, the substituted C_1 - C_6 alkyl in the substituted C_1 - C_6 alkyl or the substituted C_1 - C_6 alkyl-O- is independently - CF_3 .

In a preferred embodiment of the present invention, Q is located at position β or γ of the linkage between  and W .

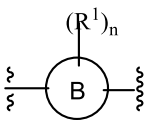
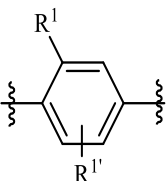
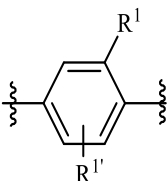
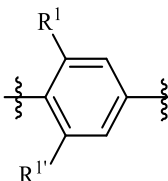
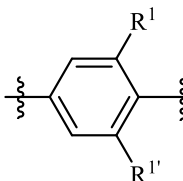
In a preferred embodiment of the present invention, when  is a 5-6 membered heteroaryl, the 5-6 membered heteroaryl is a 5 membered heteroaryl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and is preferably

imidazolyl (e.g.,  (e.g., )), thienyl (e.g.,  (e.g., ), furanyl (e.g.,  (e.g., )) or pyrrolyl (e.g.,  (e.g., ). (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)

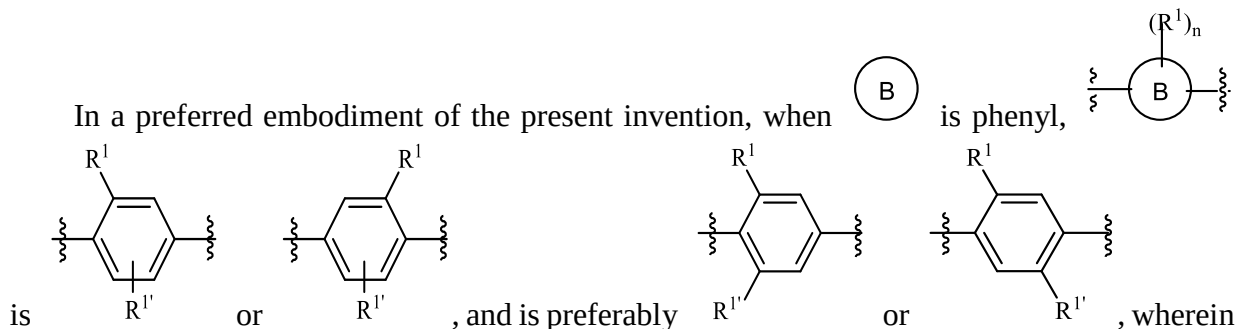
5 In a preferred embodiment of the present invention, when  is a 5-6 membered heteroaryl, the 5-6 membered heteroaryl is a 6 membered heteroaryl; wherein the 6 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and is preferably pyridinyl (e.g.,  (e.g., )) or pyrazinyl (e.g.,  (e.g., ). (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)

10 In a preferred embodiment of the present invention, when  is a C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S; the C₃-C₉ heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl is preferably piperidinyl (e.g.,  (e.g., ). (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)

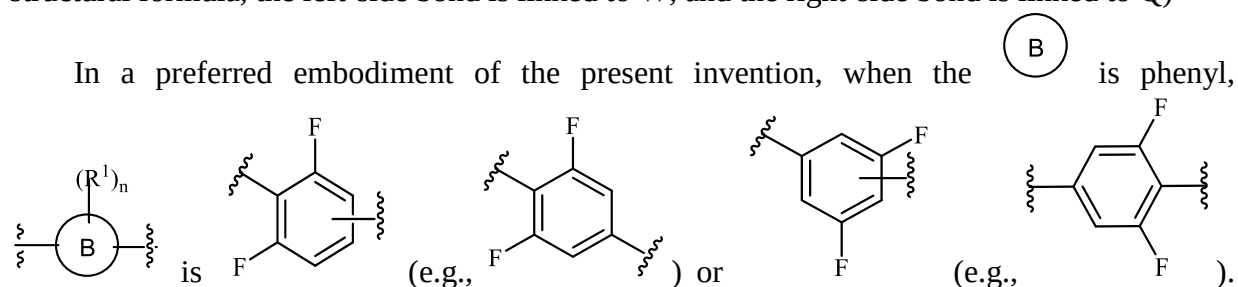
In a preferred embodiment of the present invention, R¹ is located at position β of the linkage between  and Z². Preferably, when n is 2, each R¹ is independently located at position β of the linkage between  and Z².

20 In a preferred embodiment of the present invention, when  is phenyl,  is  or  , and is preferably  or  , wherein

$R^{1'}$ has the definition as described for R^1 , and $R^{1'}$ is the same as or different from R^1 . (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)

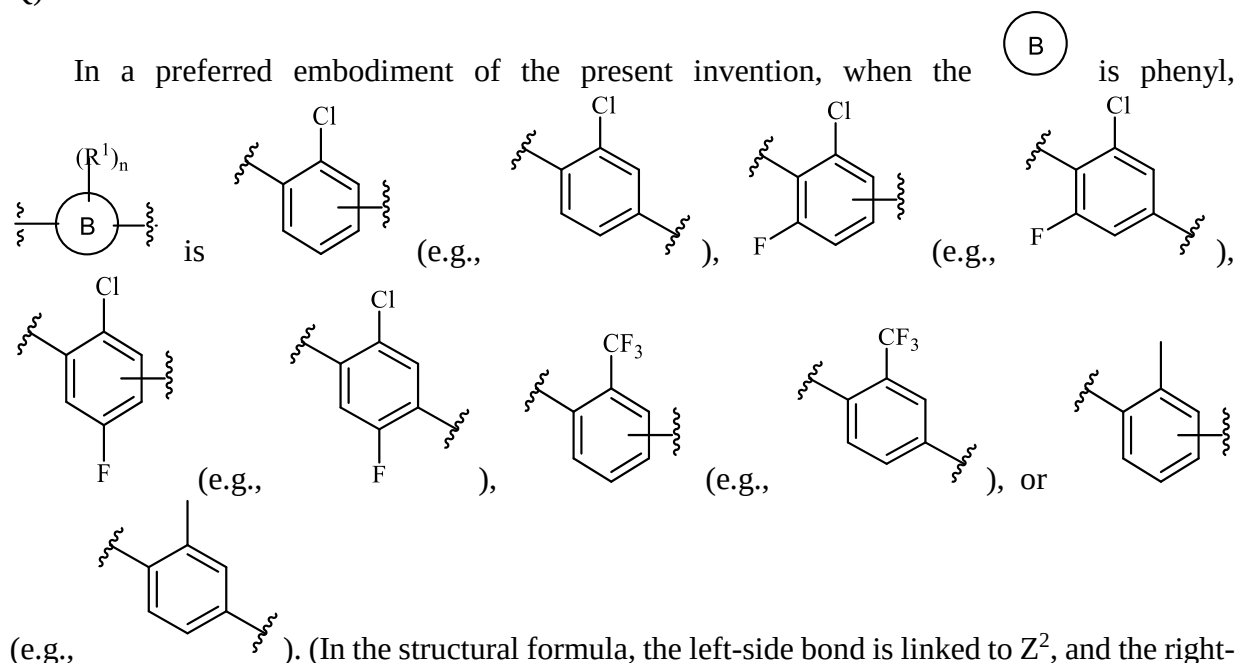


5 $R^{1'}$ has the definition as described for R^1 , and $R^{1'}$ is the same as or different from R^1 . (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)



(In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to

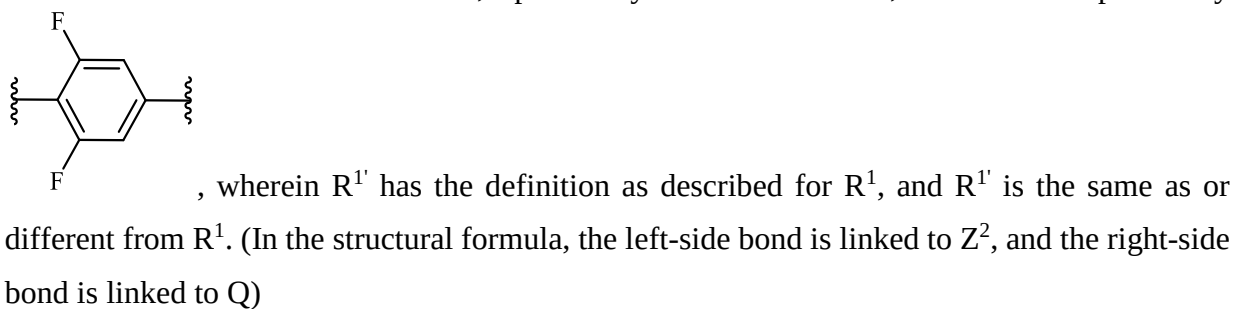
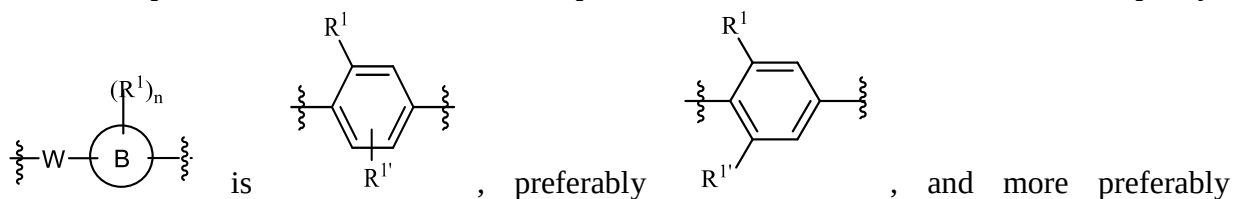
10 Q)



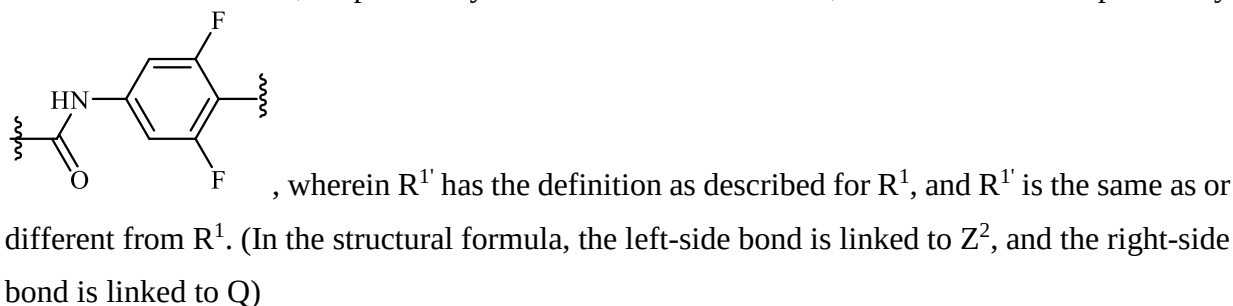
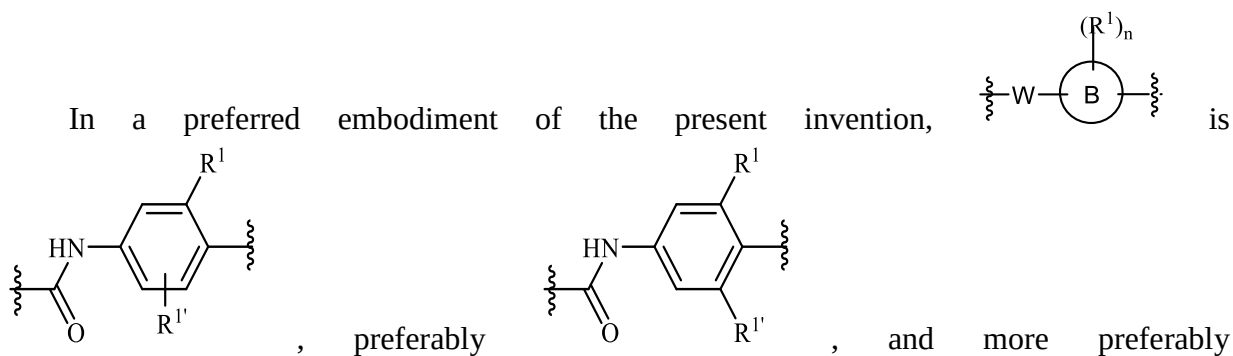
(In the structural formula, the left-side bond is linked to Z^2 , and the right-side bond is linked to Q)

15

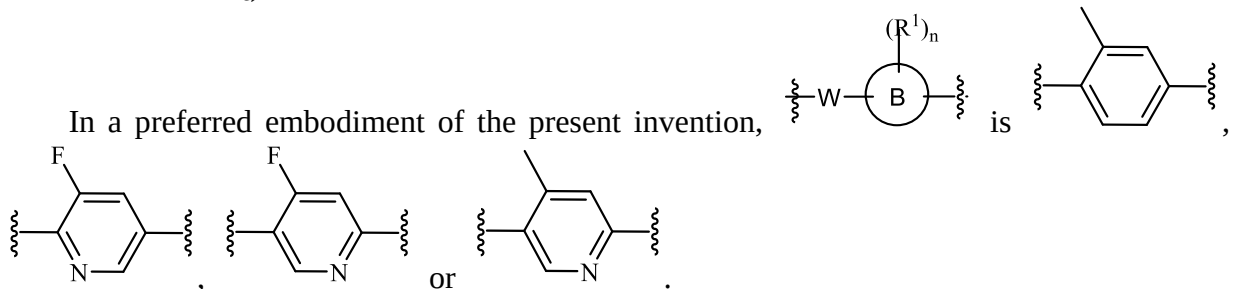
In a preferred embodiment of the present invention, when the  is phenyl,

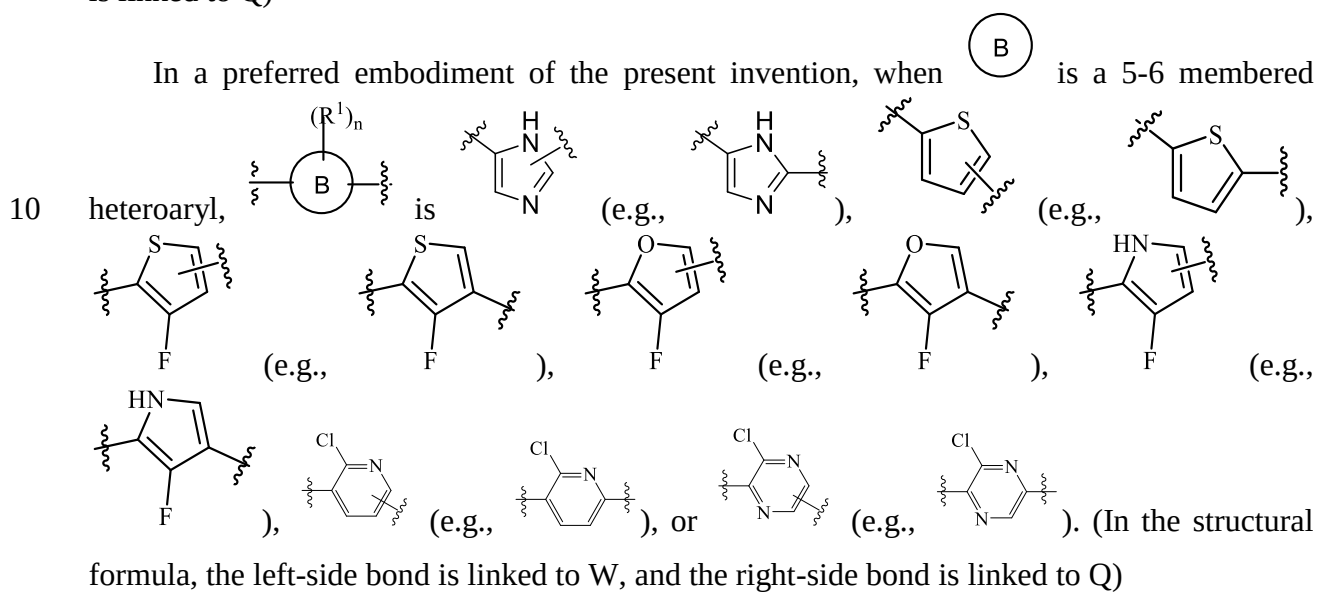
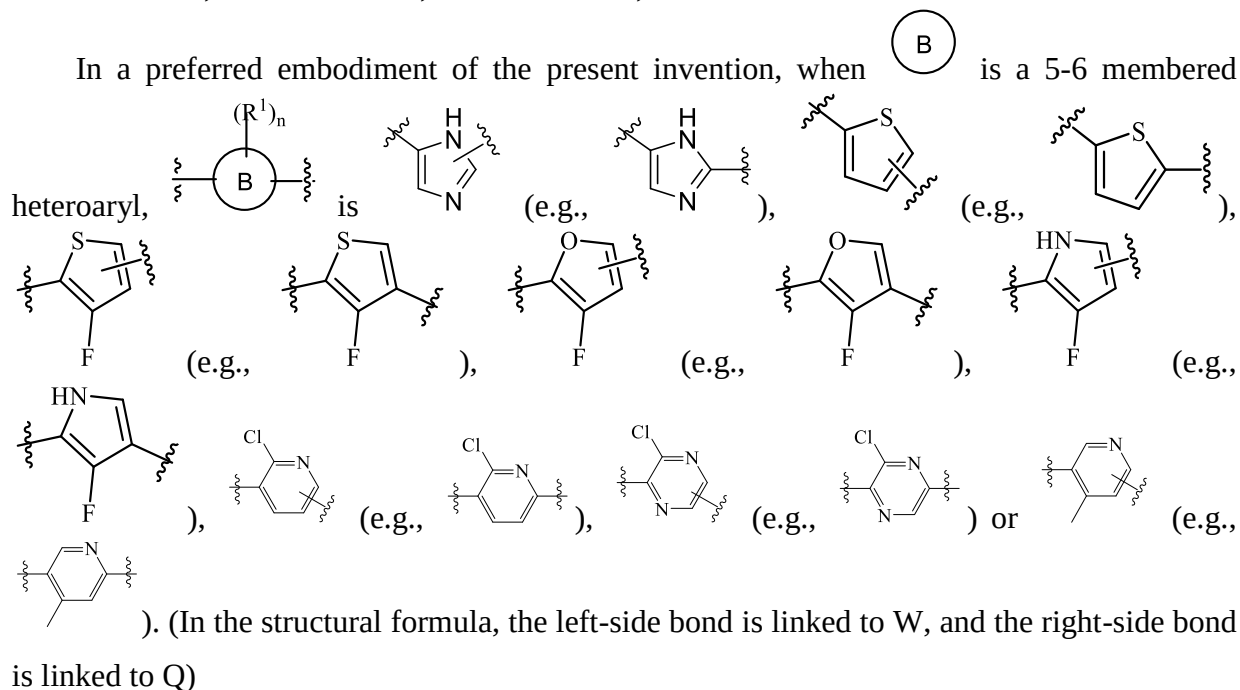
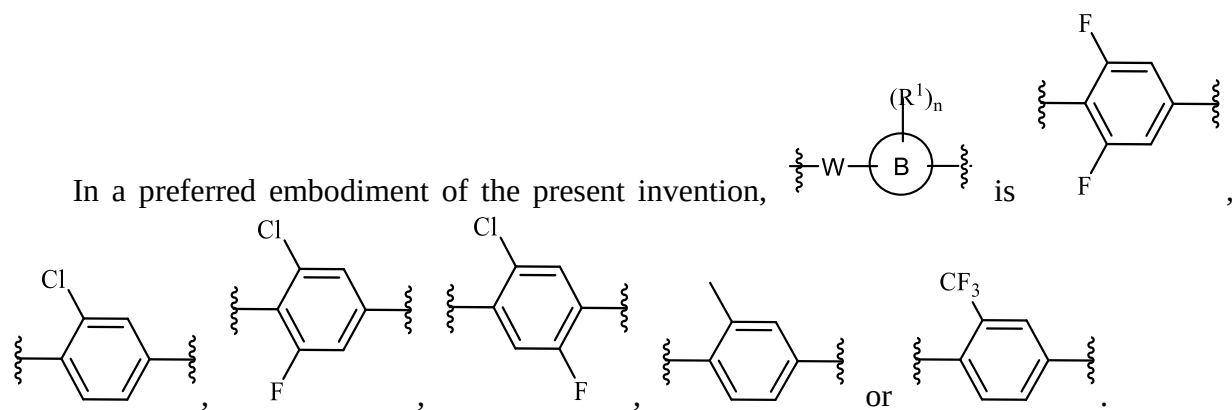


In a preferred embodiment of the present invention,

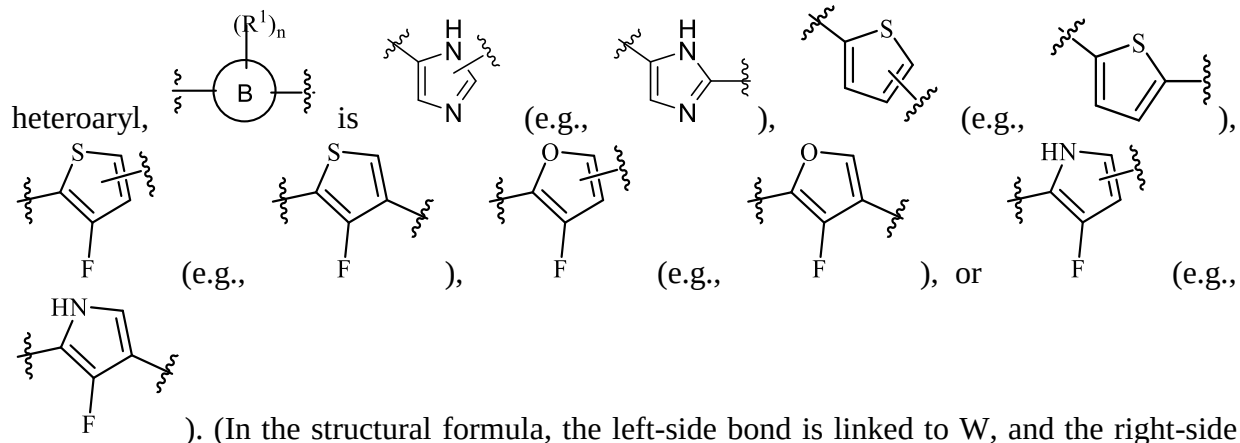


In a preferred embodiment of the present invention,



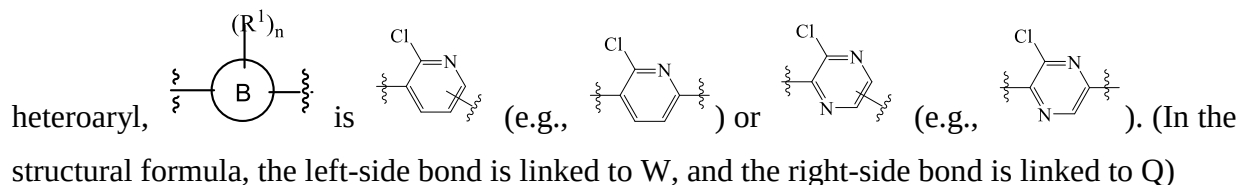


In a preferred embodiment of the present invention, when  is a 5-6 membered

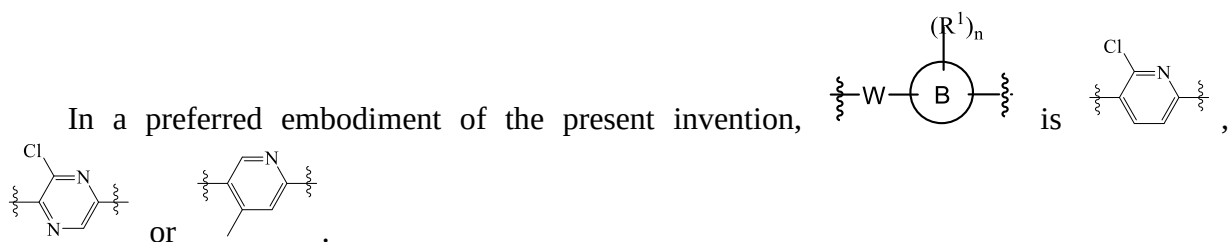


5 bond is linked to Q)

In a preferred embodiment of the present invention, when  is a 5-6 membered

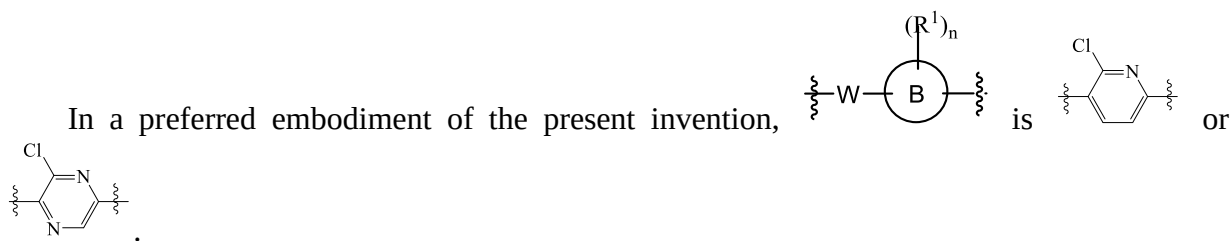


In a preferred embodiment of the present invention,

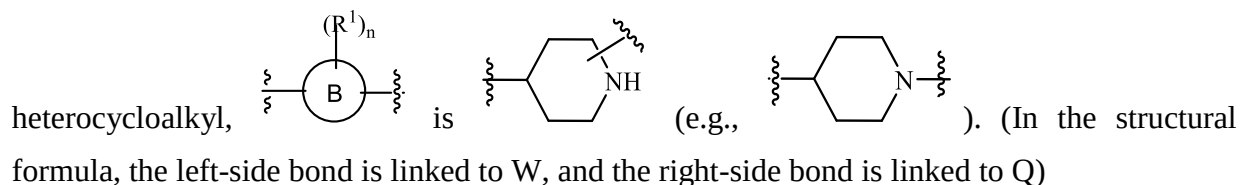


10

In a preferred embodiment of the present invention,



In a preferred embodiment of the present invention, when  is a C₃-C₅



15

In a preferred embodiment of the present invention, when Q is halogen, the halogen is fluorine, chlorine, bromine or iodine.

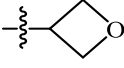
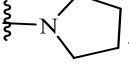
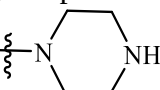
In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl (e.g.,

methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

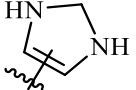
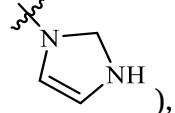
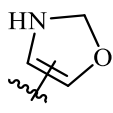
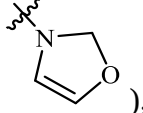
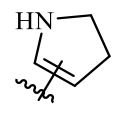
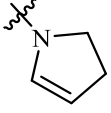
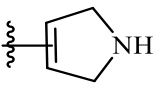
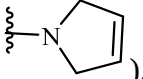
In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₂-C₆ alkenyl, or an unsubstituted or substituted C₂-C₆ alkenyl-O-, the C₂-C₆ alkenyl is a C₂-C₄ alkenyl (e.g., ethenyl, propenyl (e.g., 1-propenyl or 2-propenyl), or butenyl (e.g., 2-butenyl, 1-butenyl, or butadienyl)).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₃-C₁₀ cycloalkyl, the C₃-C₁₀ cycloalkyl is cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl.

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₃-

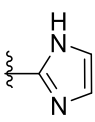
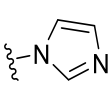
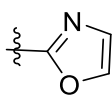
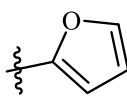
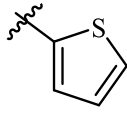
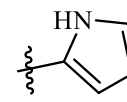
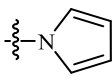
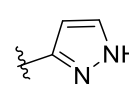
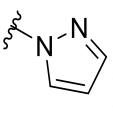
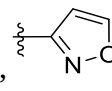
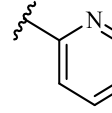
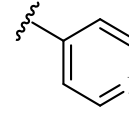
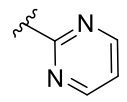
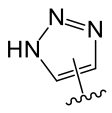
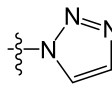
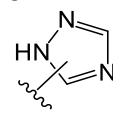
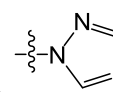
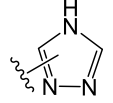
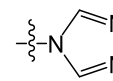
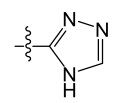
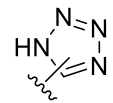
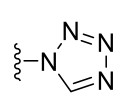
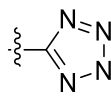
C₉ heterocycloalkyl is preferably glycidyl (e.g., , pyrrolidinyl (e.g., ) or piperazinyl (e.g., )).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl is a C₃-C₅ heterocycloalkenyl, wherein the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkenyl is preferably a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected

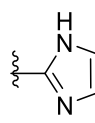
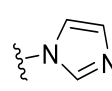
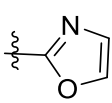
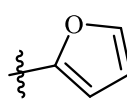
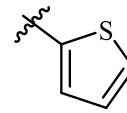
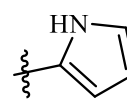
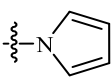
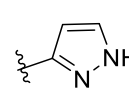
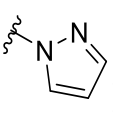
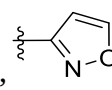
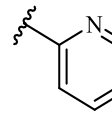
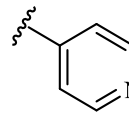
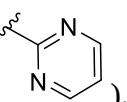
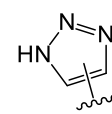
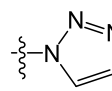
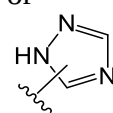
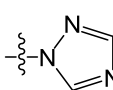
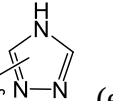
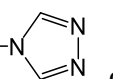
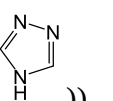
from one or more of N, O and S; and the C₃-C₉ heterocycloalkenyl is preferably  (e.g., , ) (e.g., , ) (e.g., ) or  (e.g., )).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₆-C₁₀ aryl, the C₆-C₁₀ aryl is a phenyl.

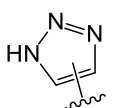
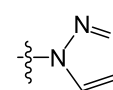
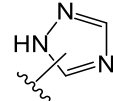
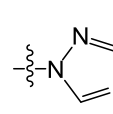
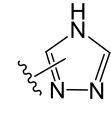
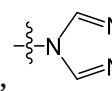
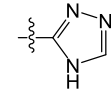
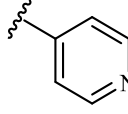
In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered

heteroaryl, and preferably imidazolyl (e.g.,  or , oxazolyl (e.g., , furanyl (e.g., , thienyl (e.g., , pyrrolyl (e.g.,  or , pyrazolyl (e.g.,  or , isoxazolyl (e.g., , pyridinyl (e.g., , or , pyrimidinyl (e.g., , triazolyl (e.g.,  (e.g., , or  (e.g., , or  (e.g.,  or , or  (e.g.,  or ).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered

heteroaryl, and preferably imidazolyl (e.g.,  or , oxazolyl (e.g., , furanyl (e.g., , thienyl (e.g., , pyrrolyl (e.g.,  or , pyrazolyl (e.g.,  or , isoxazolyl (e.g., , pyridinyl (e.g., , or , pyrimidinyl (e.g., , or triazolyl (e.g.,  (e.g., , or  (e.g., , or  (e.g.,  or )).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered

heteroaryl, and preferably triazolyl (e.g.,  (e.g., , or  (e.g., ,  (e.g.,  or )) or .

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₃-C₁₀ cycloalkyl, a substituted C₆-C₁₀ aryl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4.

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₃-C₁₀ cycloalkyl, a substituted C₆-C₁₀ aryl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine, or iodine, and preferably fluorine.

In a preferred embodiment of the present invention, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the C₁-C₆ alkyl or C₁-C₆ alkyl-O- that is unsubstituted or substituted with one or more R^a is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl or isopropyl.

In a preferred embodiment of the present invention, the C₃-C₆ cycloalkyl in the C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a is independently cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl.

In a preferred embodiment of the present invention, when R^a is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl) or *tert*-pentyl (1,1-dimethyl-propyl).

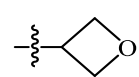
In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₂-C₆ alkenyl, or an unsubstituted or substituted C₂-C₆ alkenyl-O-, the C₂-C₆ alkenyl is independently a C₂-C₄ alkenyl (e.g. ethenyl, propenyl (e.g., 1-propenyl or 2-propenyl), or butenyl (e.g., 2-butenyl 1-butenyl or butadienyl)), and preferably ethenyl.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and

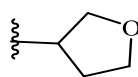
R¹⁻²⁷ are each an unsubstituted or substituted C₂-C₆ alkynyl, the C₂-C₆ alkynyl is a C₂-C₄ alkynyl (e.g., ethynyl, propynyl (e.g., 1-propynyl or 2-propynyl), or butynyl (e.g., 2-butynyl, 1-butynyl or butadiynyl)), and preferably ethynyl.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₁₀ cycloalkyl, the C₃-C₁₀ cycloalkyl is a C₃-C₆ cycloalkyl and preferably cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

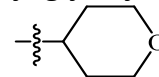
In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl is preferably glycidyl (e.g.,



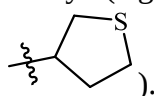
tetrahydrofuranyl (e.g.,



tetrahydropyranyl (e.g.,

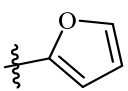
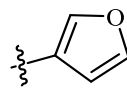
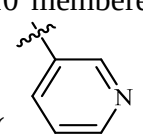
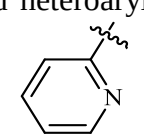
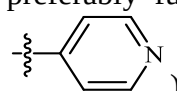


or tetrahydrothienyl (e.g.,



In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl is a C₃-C₅ heterocycloalkenyl, wherein the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the C₃-C₉ heterocycloalkenyl is preferably a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered heteroaryl, wherein the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or two of N, O and S; and the 5-10 membered heteroaryl is

preferably furanyl (e.g.,  or ) or pyridinyl (,  or ).

In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each a substituted C_1 - C_6 alkyl, a substituted C_1 - C_6 alkyl-O-, a substituted C_2 - C_6 alkenyl, a substituted C_2 - C_6 alkenyl-O-, a substituted C_2 - C_6 alkynyl, a substituted C_3 - C_{10} cycloalkyl, a substituted phenyl, a substituted C_3 - C_9 heterocycloalkyl, a substituted C_3 - C_9 heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine.

In a preferred embodiment of the present invention, when R^d is independently halogen, the halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine.

In a preferred embodiment of the present invention, the number of R^d is independently 1, 2, 3, or 4, and preferably 1 or 2.

In a preferred embodiment of the present invention, the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the C_1 - C_6 alkyl or C_1 - C_6 alkyl-O- that is unsubstituted or substituted with one or more R^d is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

In a preferred embodiment of the present invention, the C_3 - C_6 cycloalkyl in the C_3 - C_6 cycloalkyl that is unsubstituted or substituted with one or more R^d is independently cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl.

In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each a substituted C_1 - C_6 alkyl, a substituted C_1 - C_6 alkyl-O-, a substituted C_2 - C_6 alkenyl, a substituted C_2 - C_6 alkenyl-O-, a substituted C_2 - C_6 alkynyl, a substituted C_3 - C_{10} cycloalkyl, a substituted phenyl, a substituted C_3 - C_9 heterocycloalkyl, a substituted C_3 - C_9 heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , R^{1-11} and R^{1-12} , R^{1-14} and R^{1-15} , R^{1-21} and R^{1-22} , R^{1-23} and R^{1-24} , or R^{1-25} and R^{1-26} , each independently together with the N atom to which they directly link, form an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, the C_3 - C_9 heterocycloalkyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a C_3 - C_5 heterocycloalkyl.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , R^{1-11} and R^{1-12} , R^{1-14} and R^{1-15} , R^{1-21} and R^{1-22} , R^{1-23} and R^{1-24} , or R^{1-25} and R^{1-26} , each independently together with the N atom to which they directly link, form an

unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a C₃-C₅ heterocycloalkenyl.

In a preferred embodiment of the present invention, when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, each independently together with the N atom to which they directly link, form an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a 5-6 membered heteroaryl.

In a preferred embodiment of the present invention, when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, each independently together with the N atom to which they directly link, form a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

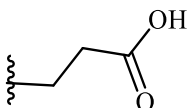
In a preferred embodiment of the present invention, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the C₁-C₆ alkyl or C₁-C₆ alkyl-O- that is unsubstituted or substituted with one or more R^b is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

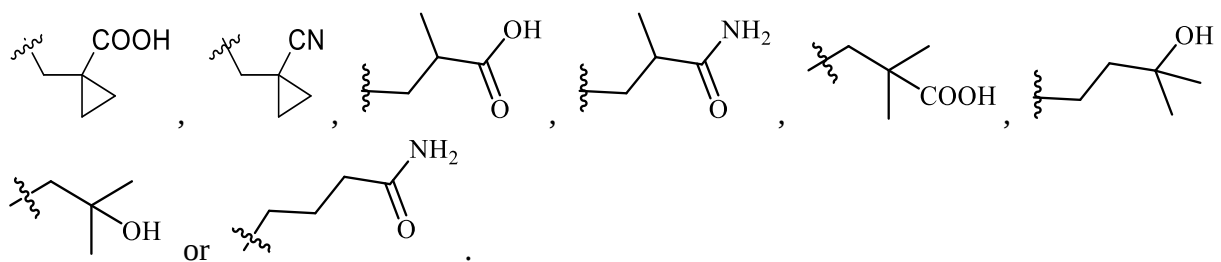
In a preferred embodiment of the present invention, the C₃-C₆ cycloalkyl in the C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b is independently cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

In a preferred embodiment of the present invention, when R^b is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, each independently together with the N atom to which they directly link, form a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2.

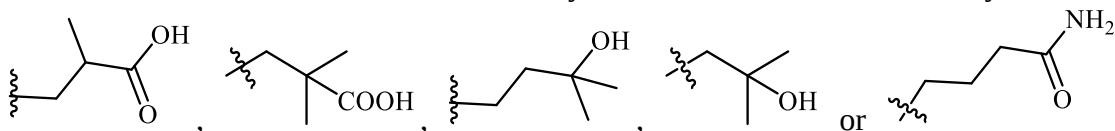
In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and

R¹⁻²⁷ are each a substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl is ,
*CCCC(=O)O



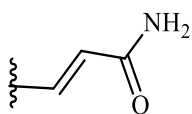
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and

5 R^{1-27} are each a substituted C_1 - C_6 alkyl, the substituted C_1 - C_6 alkyl is



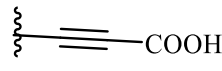
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and

R^{1-27} are each a substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6 alkenyl is



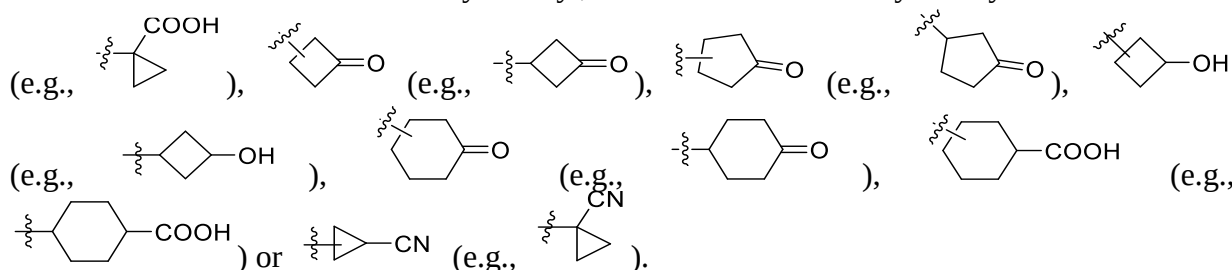
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and

R^{1-27} are each a substituted C_2 - C_6 alkynyl, the substituted C_2 - C_6 alkynyl is

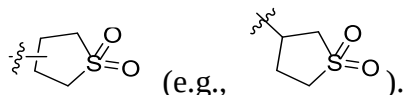


In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and

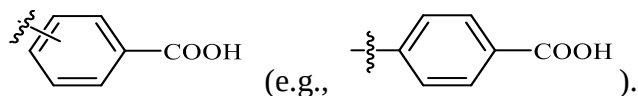
R^{1-27} are each a substituted C_3 - C_{10} cycloalkyl, the substituted C_3 - C_{10} cycloalkyl is



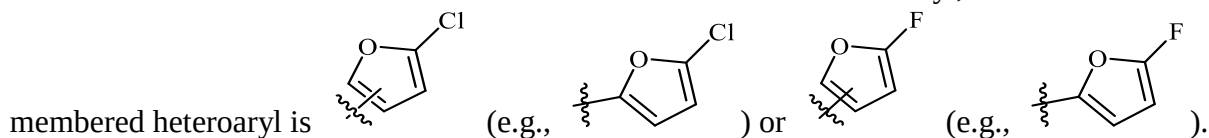
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each a substituted C_3 - C_9 heterocycloalkyl, the substituted C_3 - C_9 heterocycloalkyl is



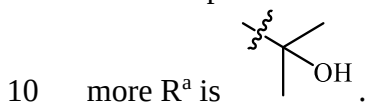
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each an unsubstituted or substituted phenyl, the substituted phenyl is



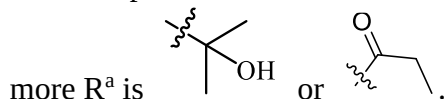
5 In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each an unsubstituted or substituted 5-10 membered heteroaryl, the substituted 5-10



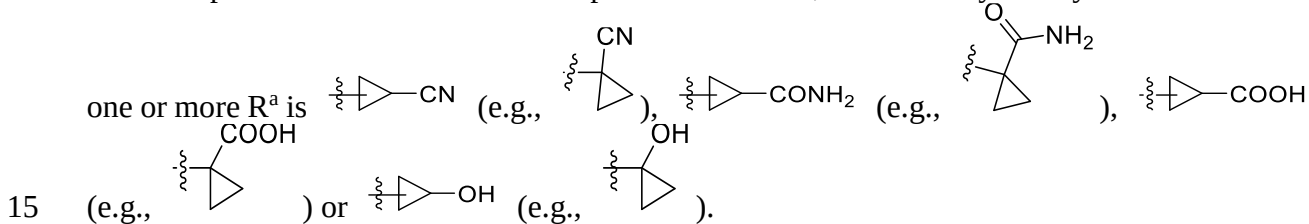
In a preferred embodiment of the present invention, the C_1 - C_6 alkyl substituted with one or



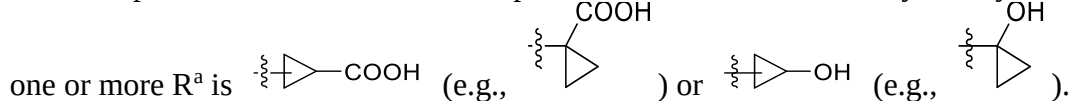
In a preferred embodiment of the present invention, the C_1 - C_6 alkyl substituted with one or



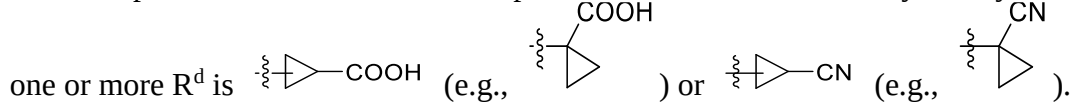
In a preferred embodiment of the present invention, the C_3 - C_6 cycloalkyl substituted with



In a preferred embodiment of the present invention, the C_3 - C_6 cycloalkyl substituted with



In a preferred embodiment of the present invention, the C_3 - C_6 cycloalkyl substituted with



20 In a preferred embodiment of the present invention, $-N(R^{1-21}R^{1-22})$ is $-NHCH_3$ or $-NH_2$.

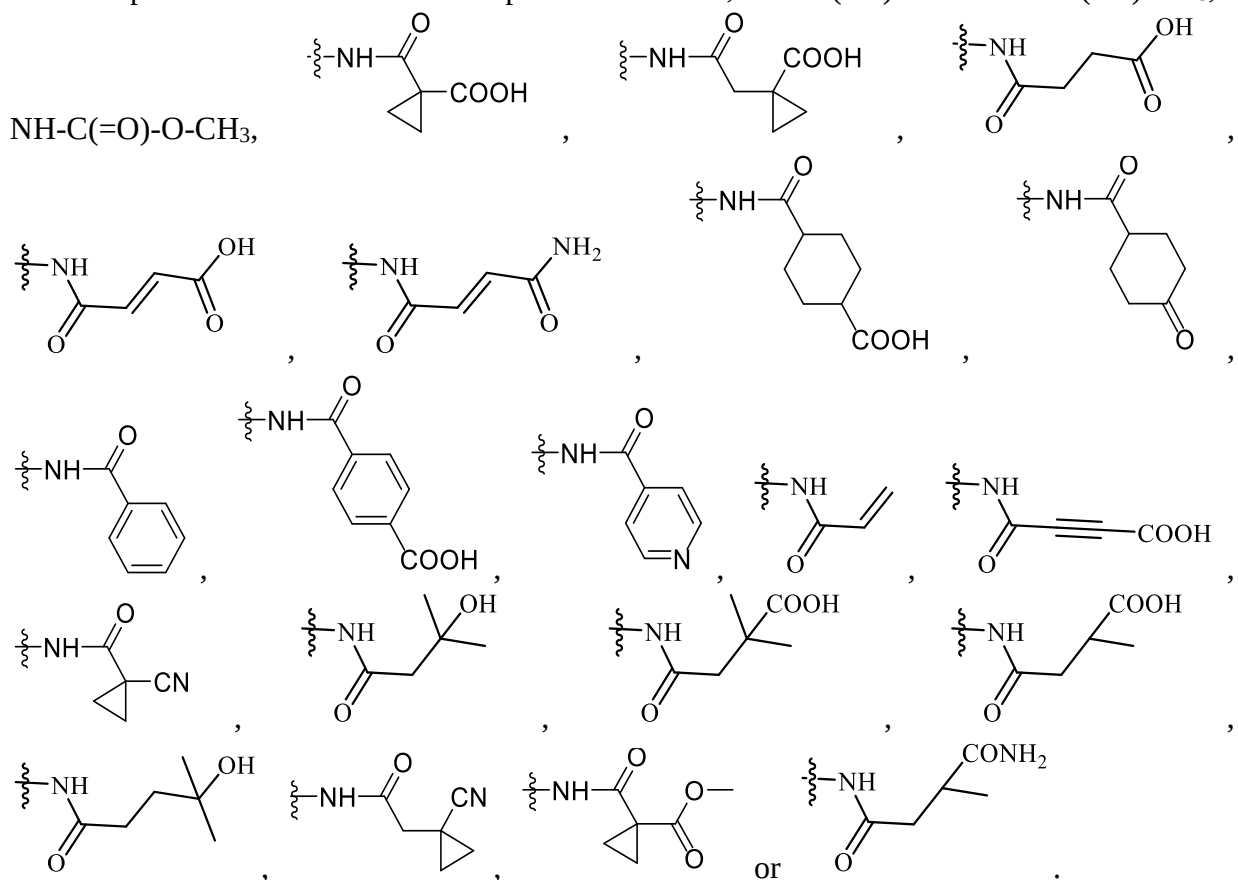
In a preferred embodiment of the present invention, $-C(=O)-N(R^{1-23}R^{1-24})$ is $-C(=O)-NH_2$ or $-C(=O)-NHCH_3$.

In a preferred embodiment of the present invention, $-S(=O)_2-N(R^{1-25}R^{1-26})$ is $-S(=O)_2-NH_2$.

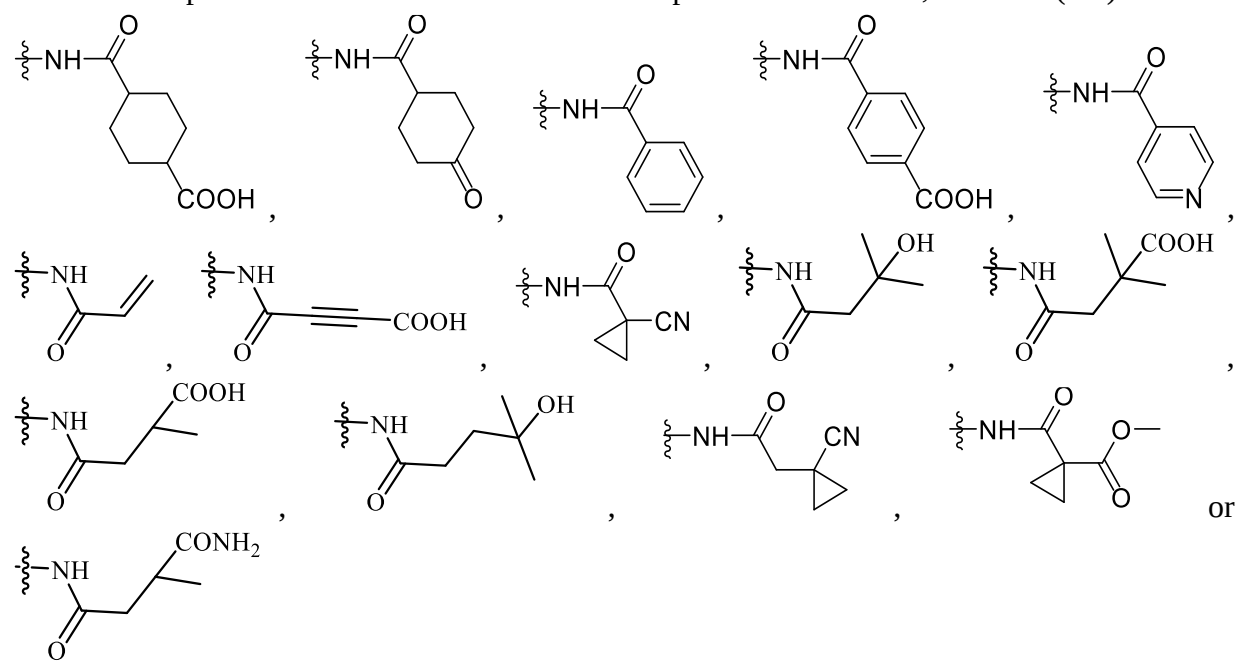
25 In a preferred embodiment of the present invention, $-C(=O)-O-R^{1-27}$ is $-COOH$, $-COOCH_3$ or $-COOC_2H_5$.

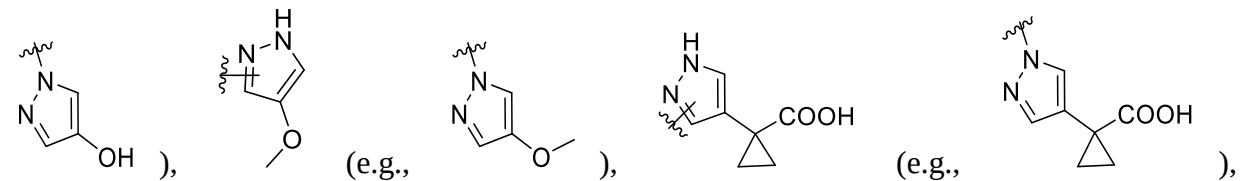
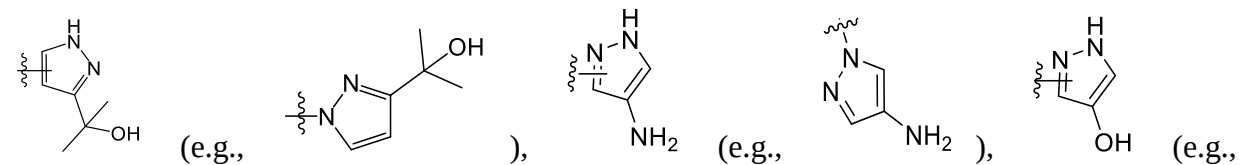
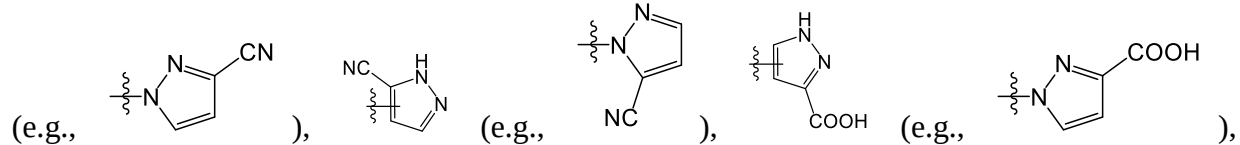
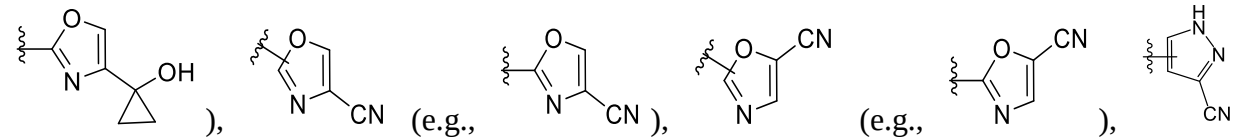
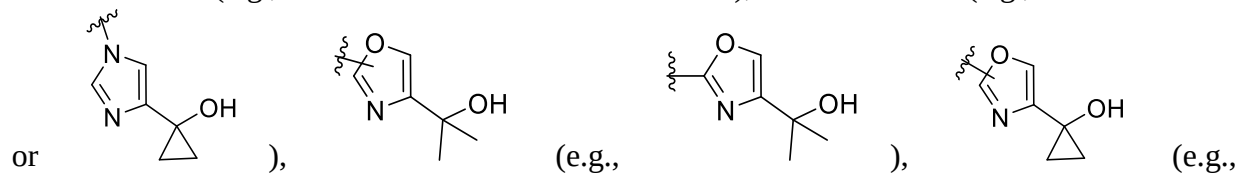
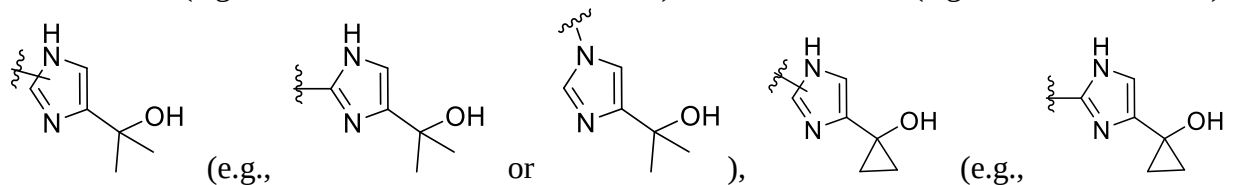
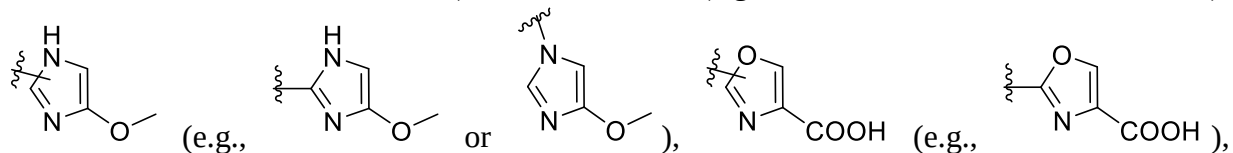
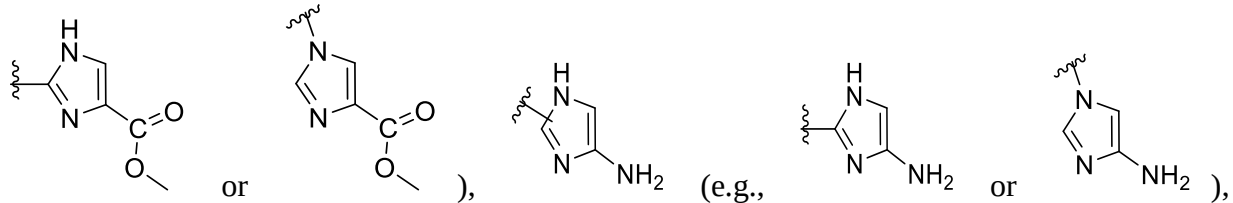
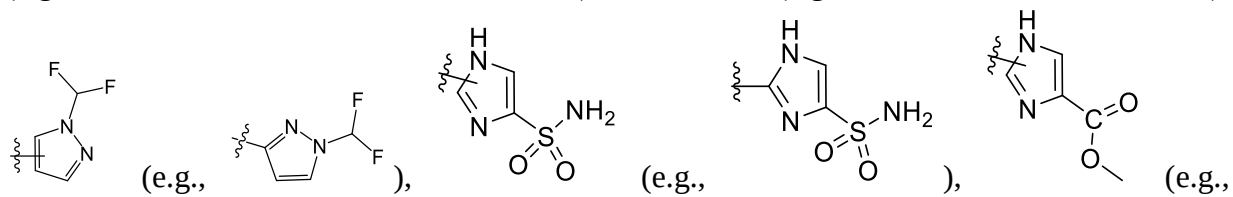
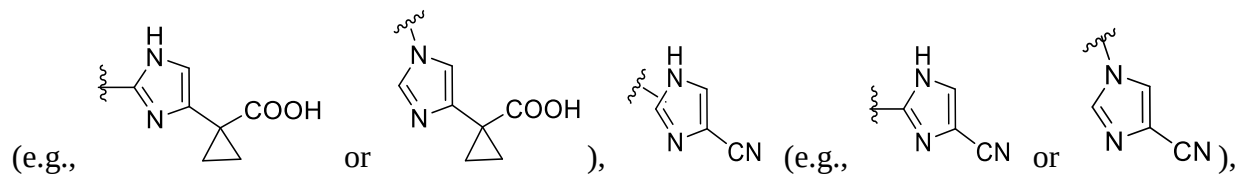
In a preferred embodiment of the present invention, $-C(=O)-O-R^{1-27}$ is $-COOH$ or $-COOCH_3$.

In a preferred embodiment of the present invention, $-NH-C(=O)-R^{1-1}$ is $-NH-C(=O)-CH_3$, $-$

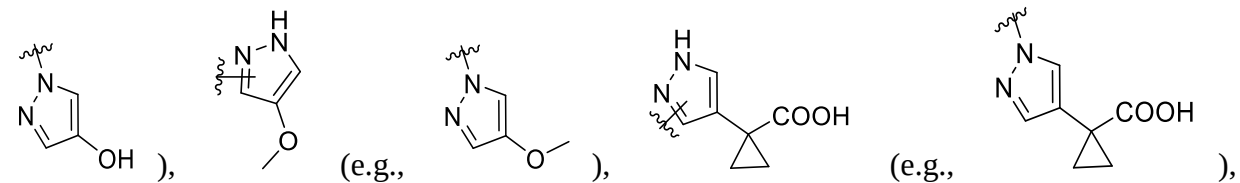
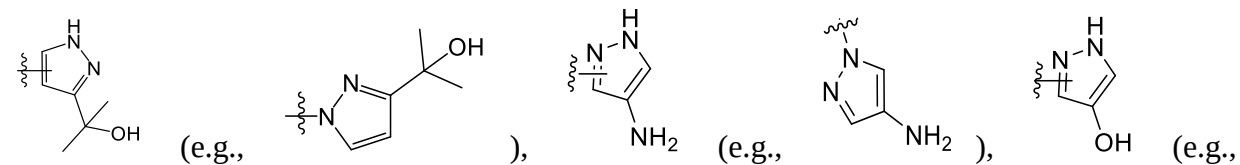
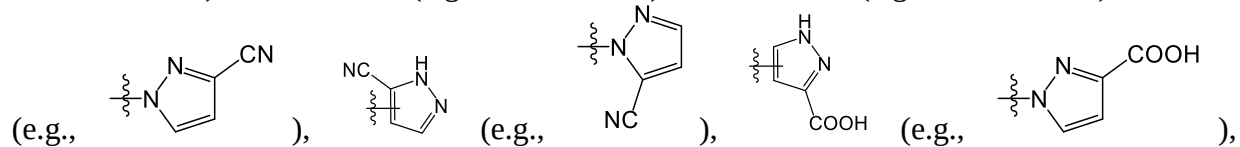
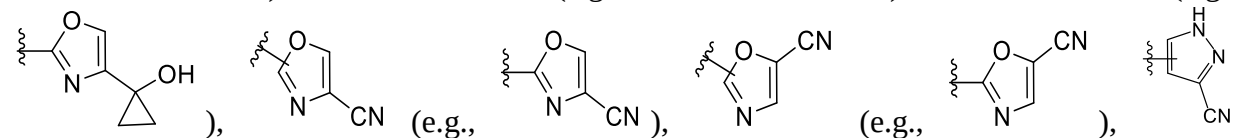
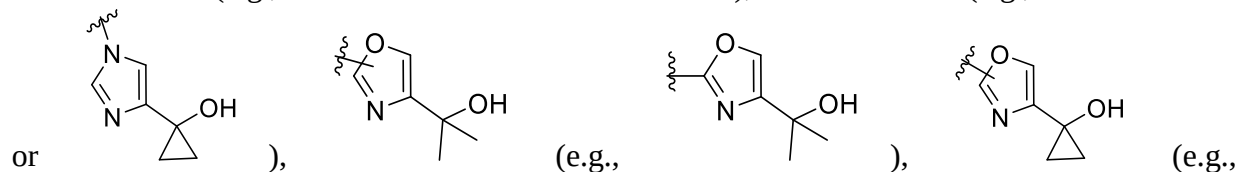
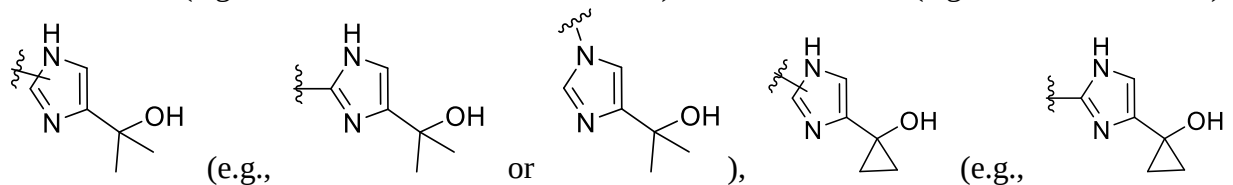
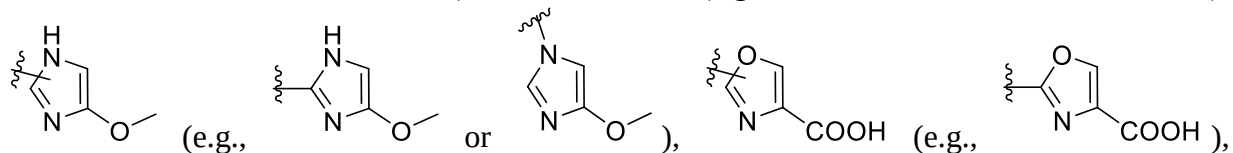
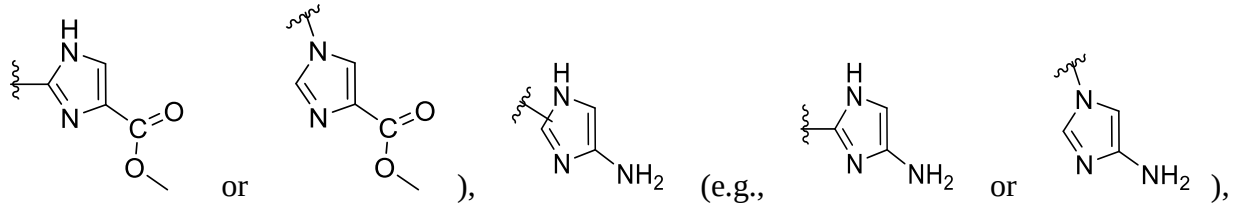
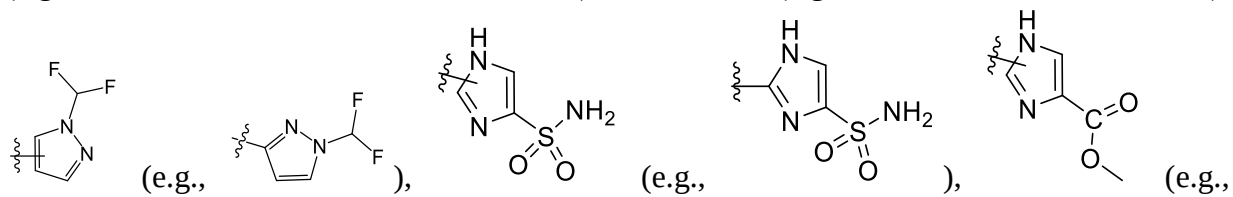
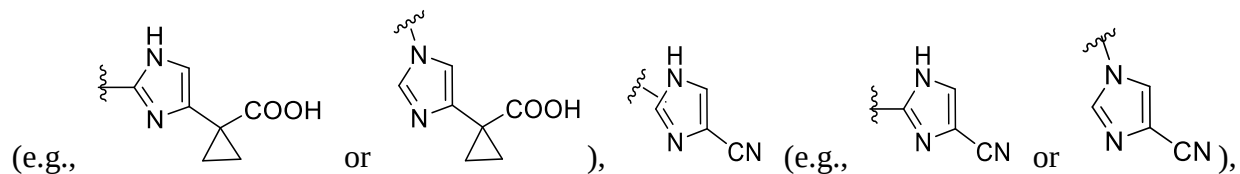


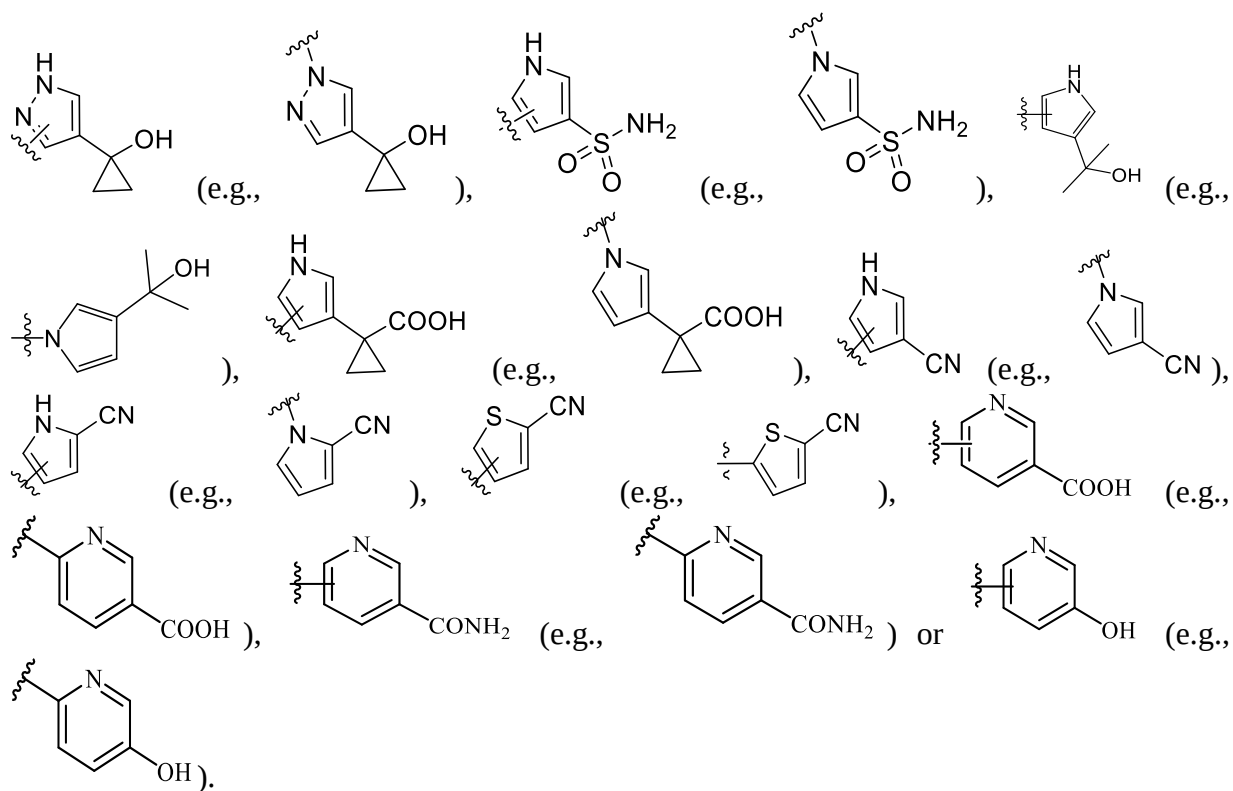
In a preferred embodiment of the present invention, $-NH-C(=O)-R^{1-1}$ is





10

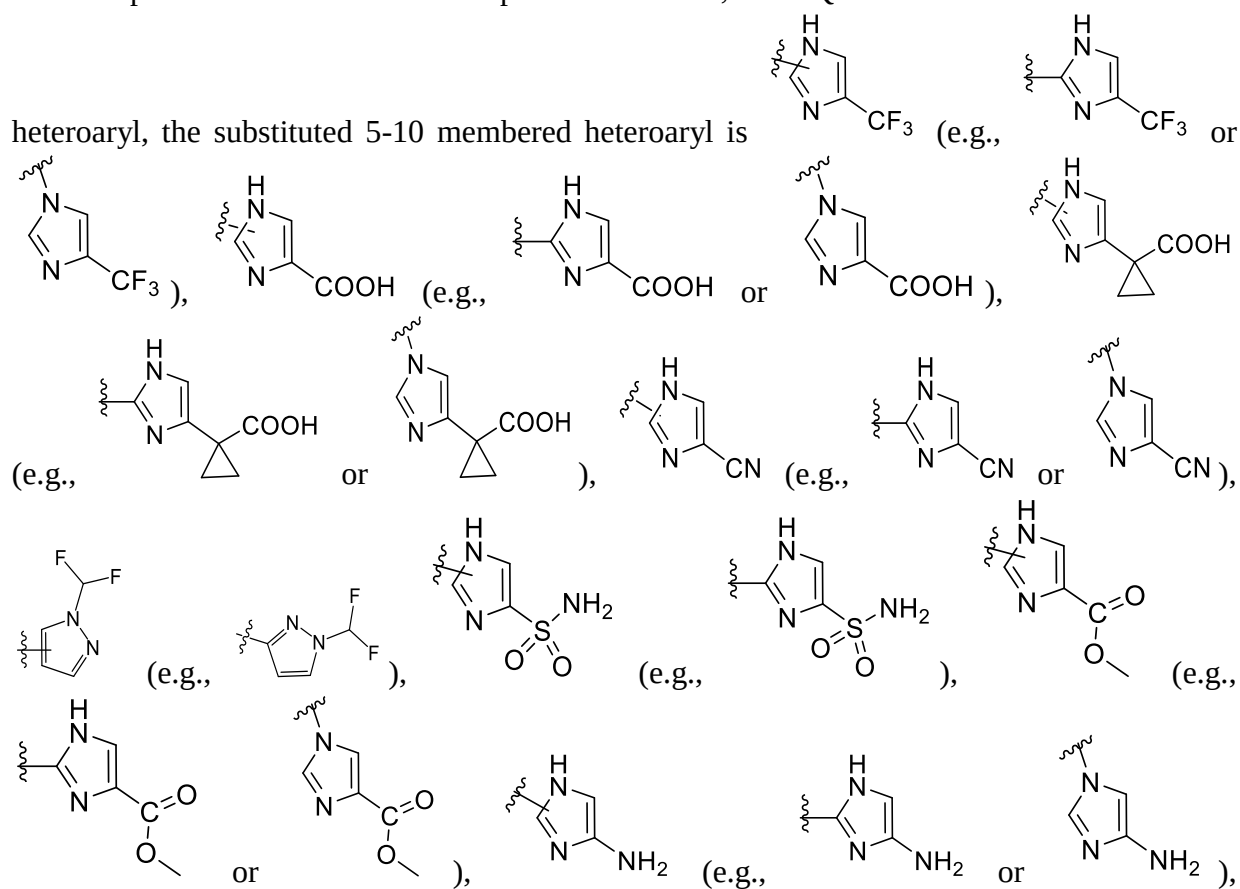




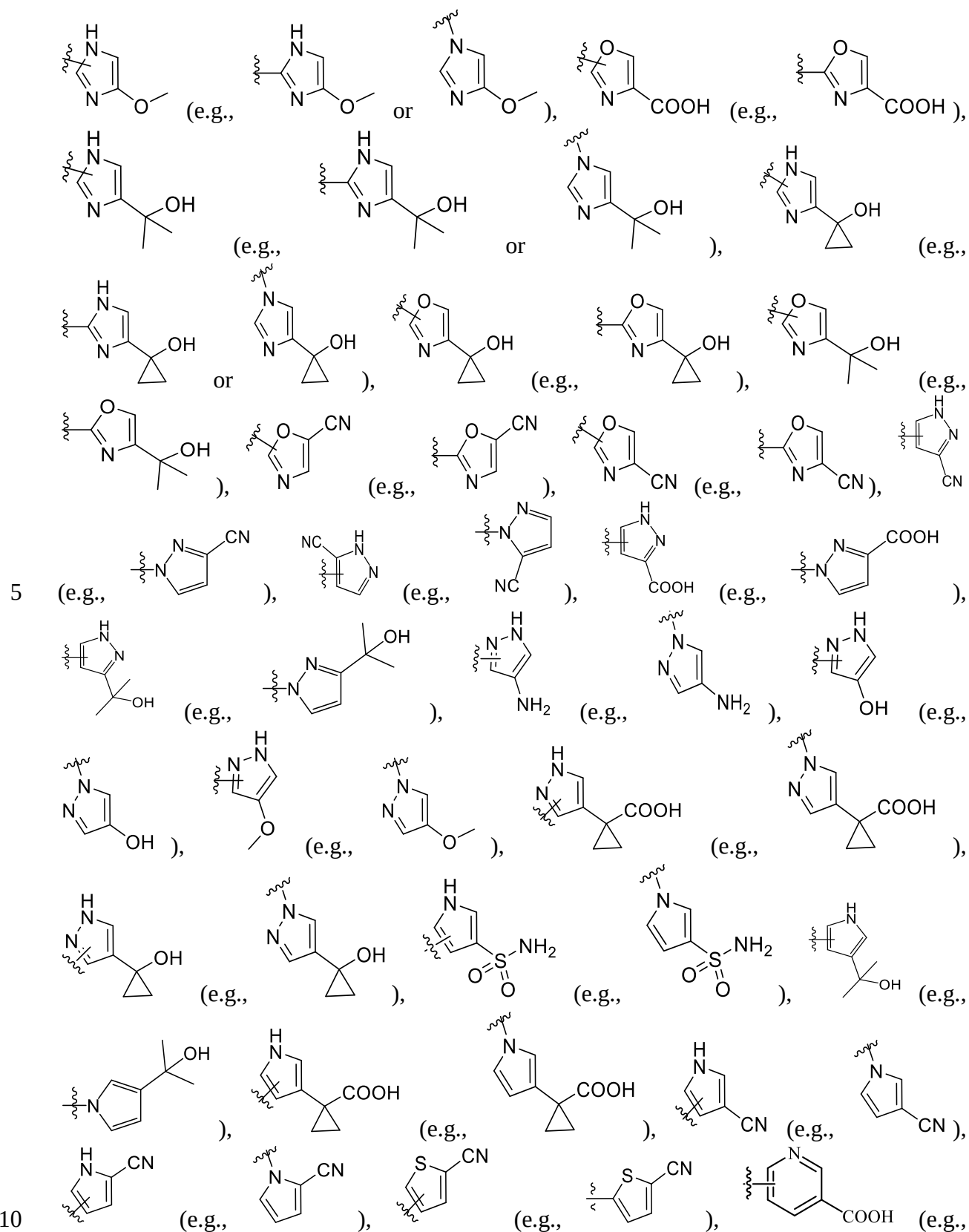
5

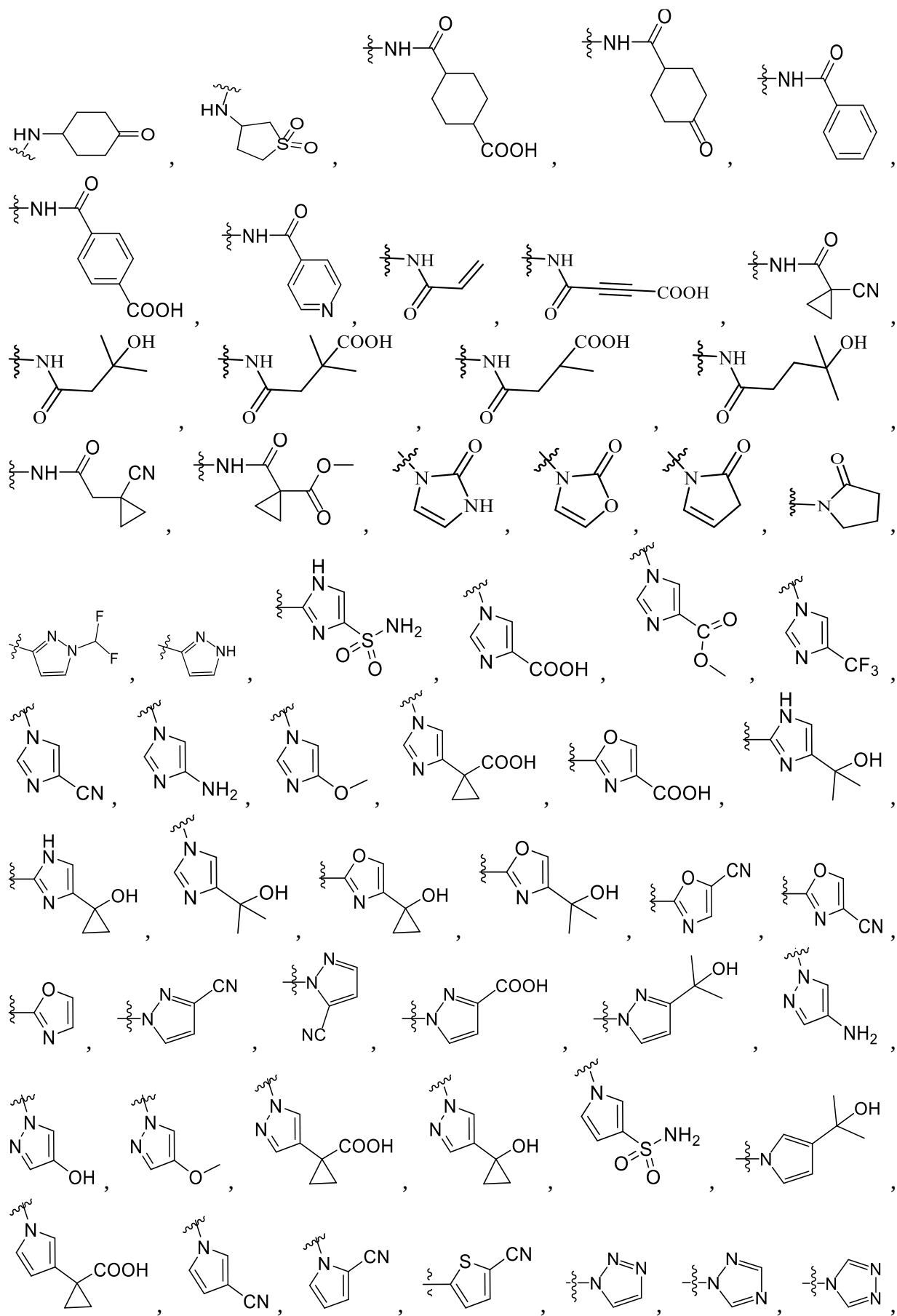
In a preferred embodiment of the present invention, when Q is a substituted 5-10 membered

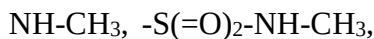
heteroaryl, the substituted 5-10 membered heteroaryl is



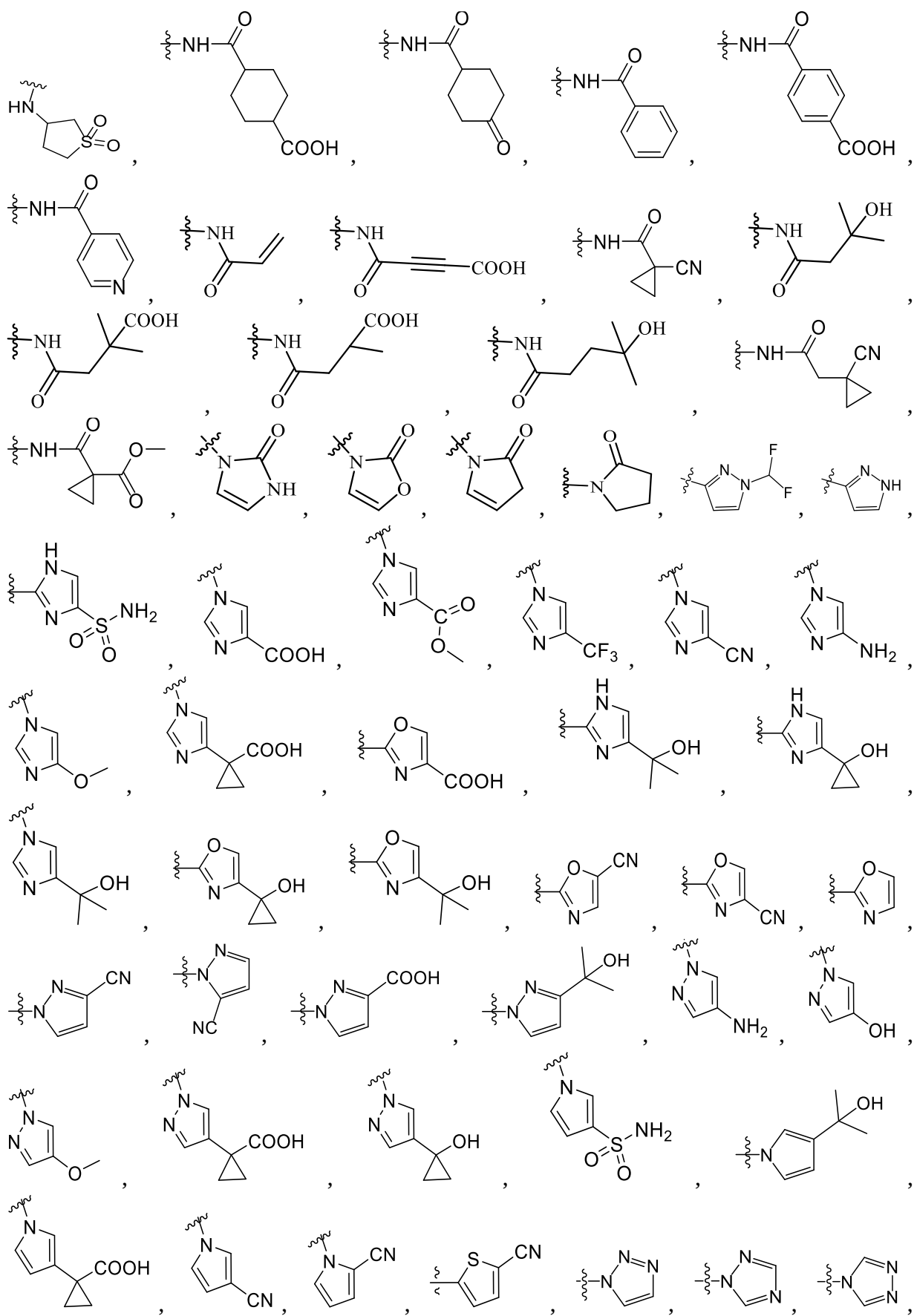
10

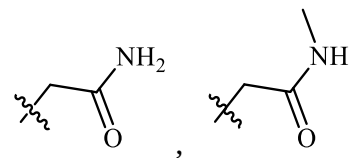




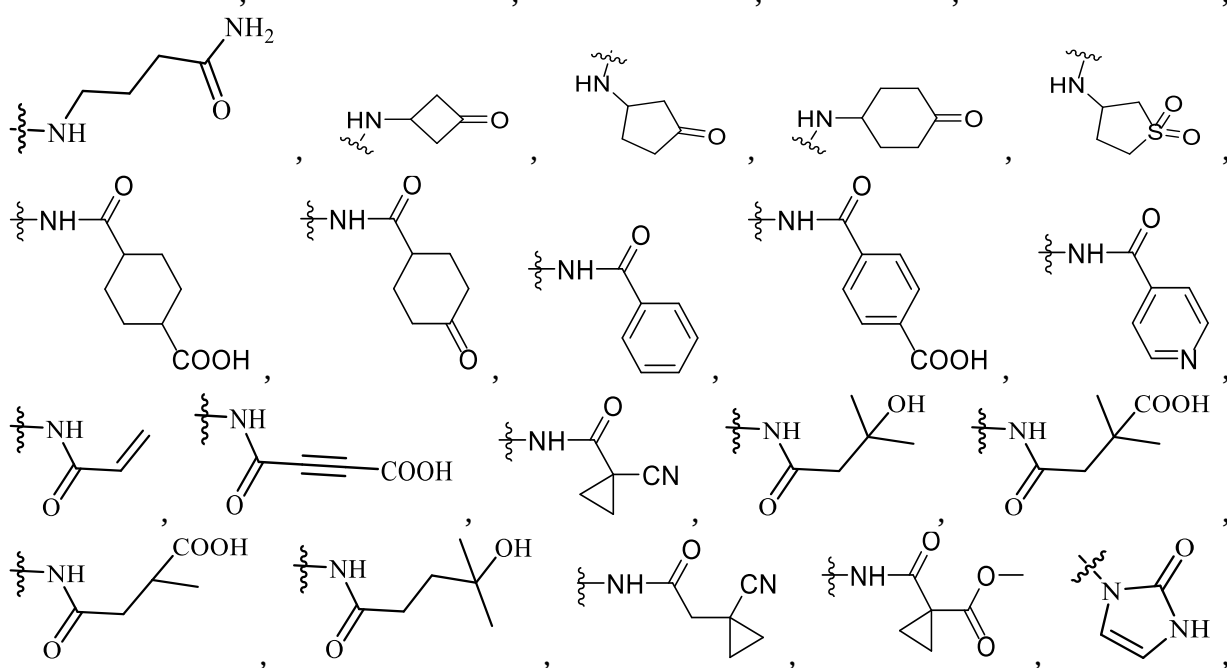
 $\text{CH}_3, \text{-NH-C(=O)-O-CH}_3,$ 

10

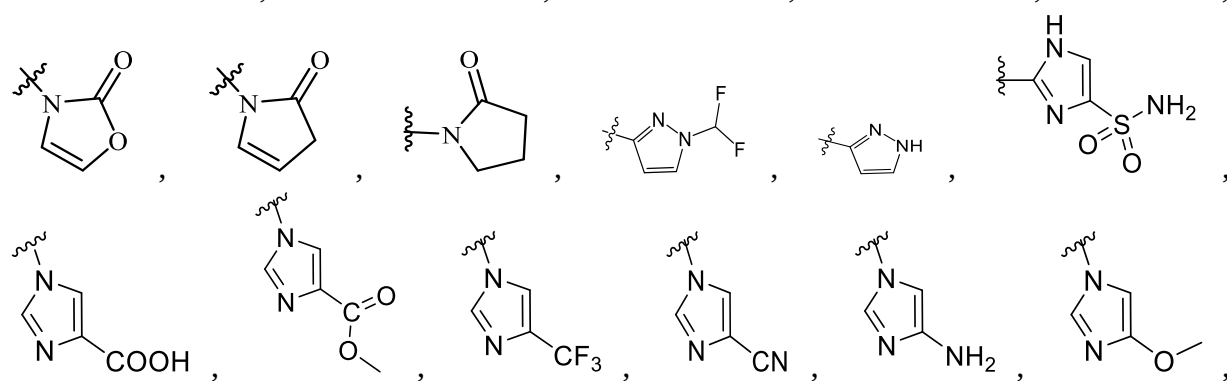


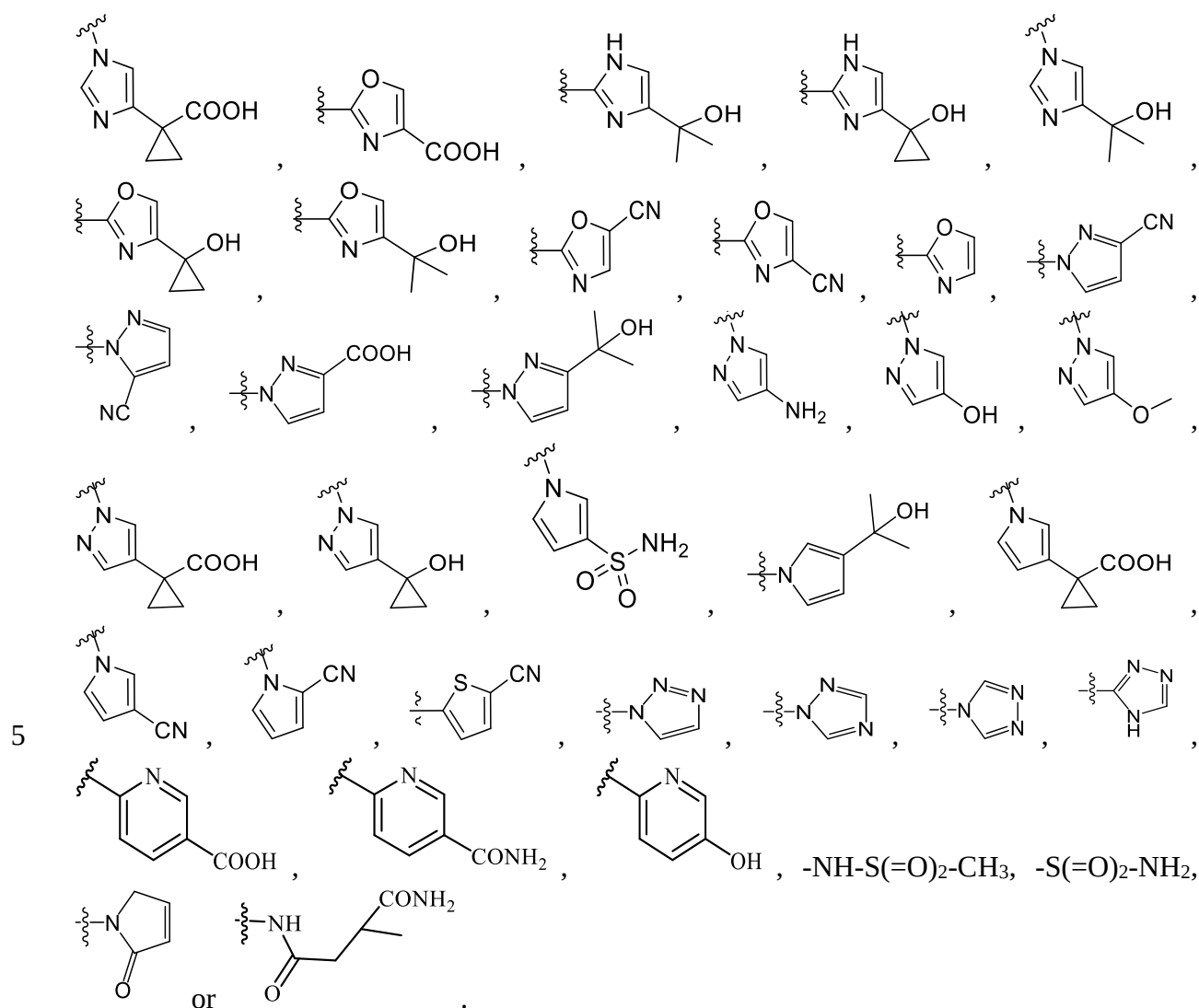
[illegible]

5



10





In a preferred embodiment of the present invention, when R^{2-1} or R^{2-2} is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

10 In a preferred embodiment of the present invention, when R^{2-1} or R^{2-2} is an unsubstituted or substituted C_1 - C_6 alkyl, or an unsubstituted or substituted C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

15 In a preferred embodiment of the present invention, when R^{2-1} or R^{2-2} is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

20 In a preferred embodiment of the present invention, when R^{2-1} or R^{2-2} is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, and the substituent is a C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the substituent is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

In a preferred embodiment of the present invention, when R^{2-1} or R^{2-2} is a substituted C_1-C_6 alkyl or a substituted C_1-C_6 alkyl-O-, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2.

In a preferred embodiment of the present invention, when R^{2-3} , R^{2-4} , R^{2-5} or R^{2-6} is an unsubstituted or substituted C_1-C_6 alkyl, or an unsubstituted or substituted C_1-C_6 alkyl-O-, the C_1-C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C_1-C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

In a preferred embodiment of the present invention, when R^{2-3} , R^{2-4} , R^{2-5} or R^{2-6} is an unsubstituted or substituted C_3-C_8 cycloalkyl, or an unsubstituted or substituted C_3-C_8 cycloalkyl-O-, the C_3-C_8 cycloalkyl is a C_3-C_6 cycloalkyl (e.g., cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl).

In a preferred embodiment of the present invention, when R^{2-3} , R^{2-4} , R^{2-5} or R^{2-6} is a substituted C_1-C_6 alkyl, a substituted C_3-C_8 cycloalkyl, a substituted C_3-C_8 cycloalkyl-O-, or a substituted C_1-C_6 alkyl-O-, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R^{2-3} , R^{2-4} , R^{2-5} or R^{2-6} is a substituted C_1-C_6 alkyl, a substituted C_3-C_8 cycloalkyl, a substituted C_3-C_8 cycloalkyl-O-, or a substituted C_1-C_6 alkyl-O-, and the substituent is a C_1-C_6 alkyl or a substituted C_1-C_6 alkyl-O-, the C_1-C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the substituent is independently a C_1-C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl or ethyl.

In a preferred embodiment of the present invention, when R^{2-3} , R^{2-4} , R^{2-5} or R^{2-6} is a substituted C_1-C_6 alkyl, a substituted C_3-C_8 cycloalkyl, a substituted C_3-C_8 cycloalkyl-O- or a substituted C_1-C_6 alkyl-O-, and the substituent is a C_1-C_6 alkyl or a substituted C_1-C_6 alkyl-O-, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2.

In a preferred embodiment of the present invention, when R^{2-7} is independently halogen, R^{2-7} is independently a halogen-substituted C_1-C_6 alkyl, or R^{2-7} is independently a halogen-substituted C_1-C_6 alkyl-O-, the halogen therein is independently fluorine, chlorine, bromine or iodine.

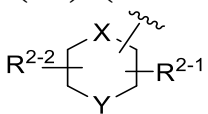
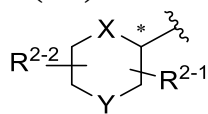
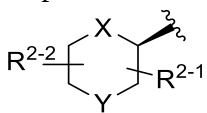
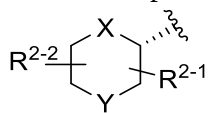
In a preferred embodiment of the present invention, when R^{2-7} is independently an unsubstituted or substituted C_1-C_6 alkyl, or R^{2-7} is independently an unsubstituted or substituted C_1-C_6 alkyl-O-, the C_1-C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) therein is independently a C_1-C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

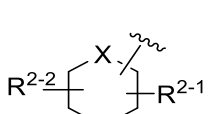
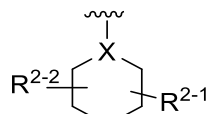
In a preferred embodiment of the present invention, when R^{2-8} or R^{2-9} is independently a halogen-substituted C_1 - C_6 alkyl, the halogen therein is independently fluorine, chlorine, bromine or iodine.

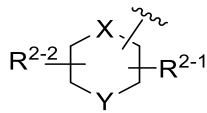
In a preferred embodiment of the present invention, when R^{2-8} or R^{2-9} is independently an unsubstituted or halogen-substituted C_1 - C_6 alkyl, the C_1 - C_6 alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) therein is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

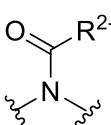
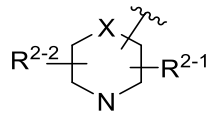
In a preferred embodiment of the present invention, $-C(=O)N(R^{2-5}R^{2-6})$ is $-C(=O)NH_2$.

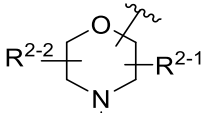
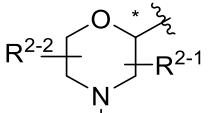
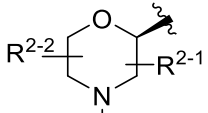
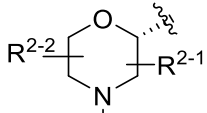
In a preferred embodiment of the present invention, $-C(=O)N(R^{2-8}R^{2-9})$ is $-C(=O)NHCH_3$.

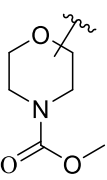
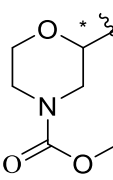
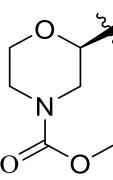
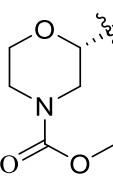
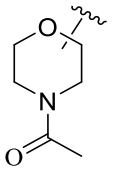
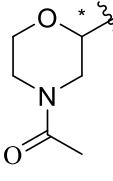
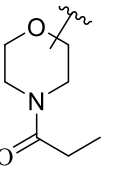
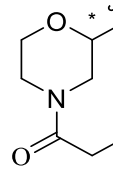
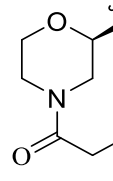
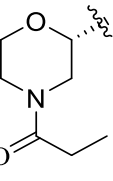
In a preferred embodiment of the present invention,  is  (e.g.,  or , wherein, the carbon atom labeled with "*" is a chiral carbon atom in S configuration, R configuration or a mixture thereof; and preferably, the carbon atom labeled with "*" is a chiral carbon atom in a form of a single (*R*) or (*S*) enantiomer or in a form enriched with one enantiomer.

In a preferred embodiment of the present invention,  is  (wherein X is methylene or -NH-).

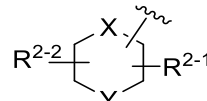
In a preferred embodiment of the present invention, when R^2 is , one of X

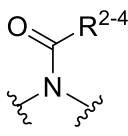
and Y is , and the other is a single bond, methylene or -O- (e.g., , wherein X is a single bond, methylene or -O-); preferably X is -O-, namely, R^2 is

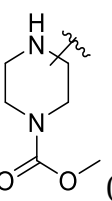
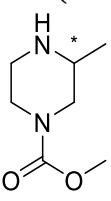
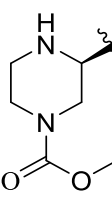
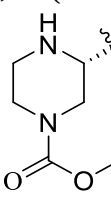
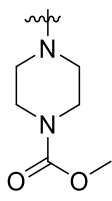
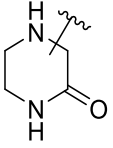
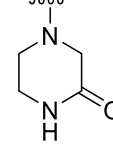
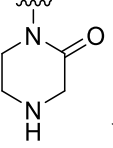
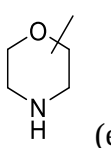
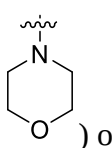
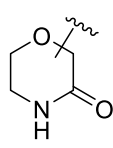
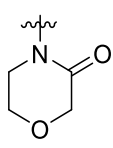
 (e.g.,  (e.g.,  or )); and

more preferably, R^2 is  (e.g.,  (e.g.,  or ),
 (e.g., ) or  (e.g.,  (e.g.,  or ).

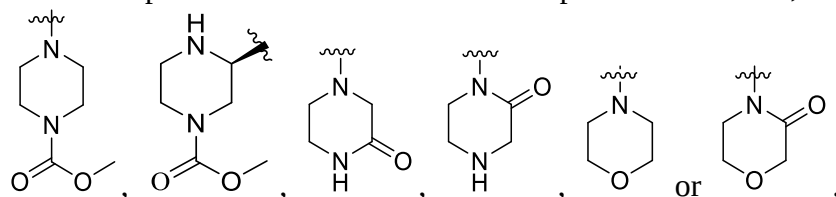
In a preferred embodiment of the present invention, when R^2 is



, X or Y is  R^{2-4} ;
 $-\text{N}(\text{R}^{2-3})-$; preferably, one of X and Y is $-\text{N}(\text{R}^{2-3})-$, and the other is $-\text{O}-$, $-\text{N}(\text{R}^{2-3})-$ or

5 more preferably, R^2 is  (e.g.,  (e.g.,  or ) or ),
 (e.g.,  or ),
 (e.g.,  or  (e.g., ).

In a preferred embodiment of the present invention, R^2 is

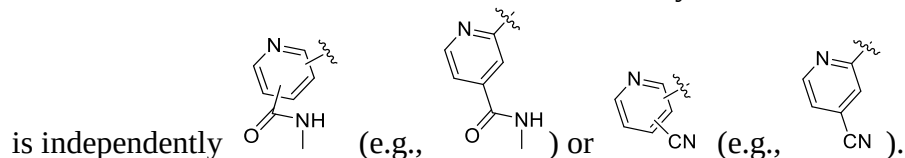


10 In a preferred embodiment of the present invention, when R^2 is independently an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl is independently pyridinyl (e.g.,  (e.g., ).

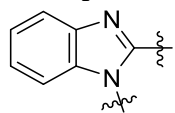
In a preferred embodiment of the present invention, when R^2 is independently a substituted 5-6 membered heteroaryl that may be substituted with halogen, a halogen-substituted $\text{C}_1\text{-C}_6$ alkyl or a halogen-substituted $\text{C}_1\text{-C}_6$ alkyl-O-, the halogen in the substituents is independently fluorine,
 15 chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R² is independently a substituted 5-6 membered heteroaryl that may be substituted with an unsubstituted or halogen-substituted C₁-C₆ alkyl, or an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the substituents is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

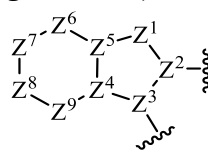
In a preferred embodiment of the present invention, when R² is independently an unsubstituted or substituted 5-6 membered heteroaryl, the substituted 5-6 membered heteroaryl



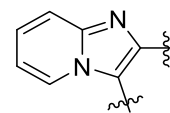
In a preferred embodiment of the present invention,



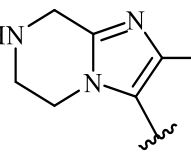
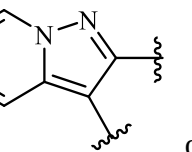
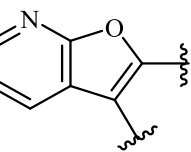
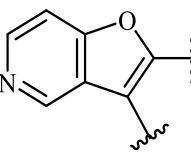
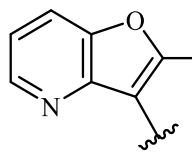
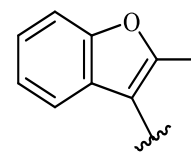
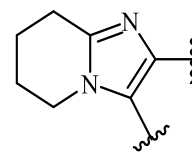
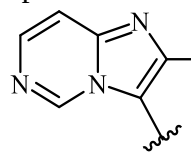
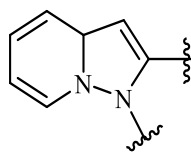
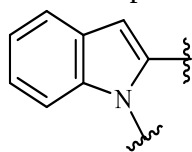
10 or



is not



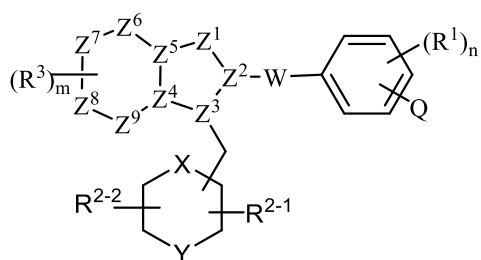
In a preferred embodiment of the present invention,



or

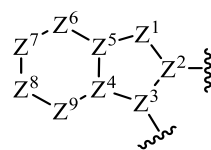
In a preferred embodiment of the present invention, the heterocyclic compound of formula

15 I is of formula I-A as shown below:

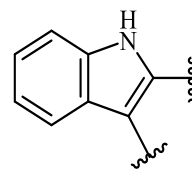


I-A

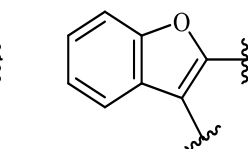
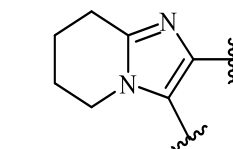
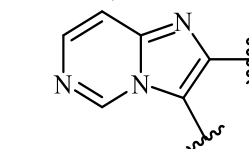
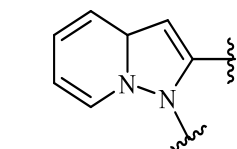
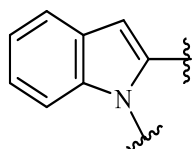
, wherein



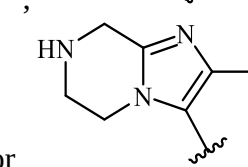
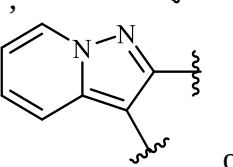
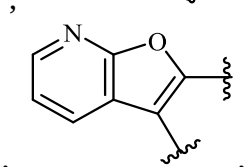
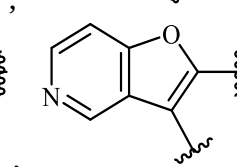
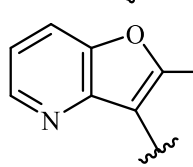
is



,



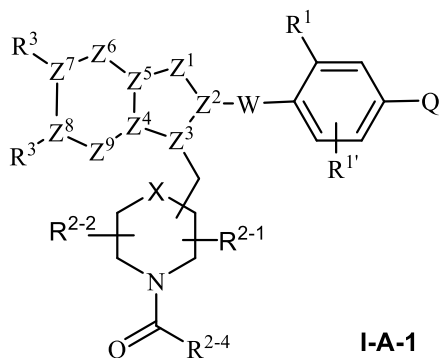
,



;

or

preferably, the heterocyclic compound of formula I is of formula I-A-1 as shown below:

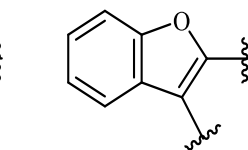
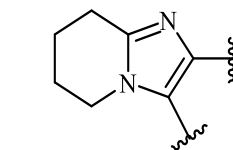
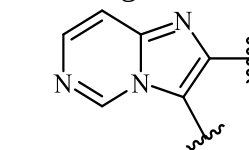
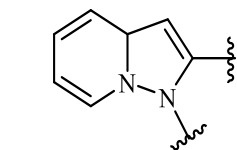
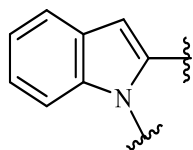


I-A-1

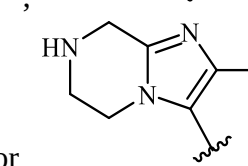
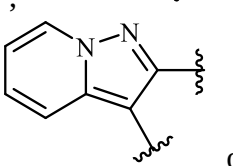
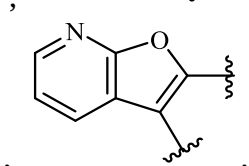
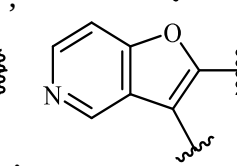
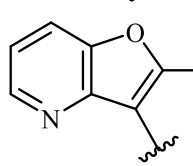
, wherein $R^{1'}$ has the definition as described for R^1 , and $R^{1'}$ is

5

the same as or different from R^1 ; W is a single bond; and



,

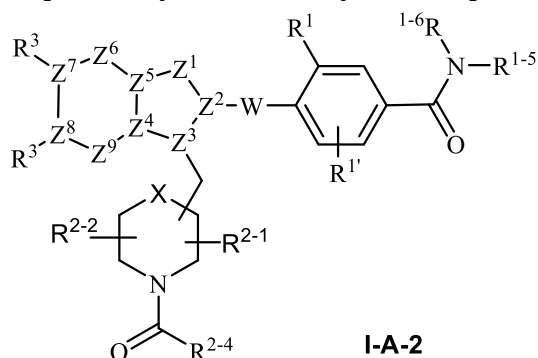


;

or

and

more preferably, the heterocyclic compound of formula I is of formula I-A-2 as shown

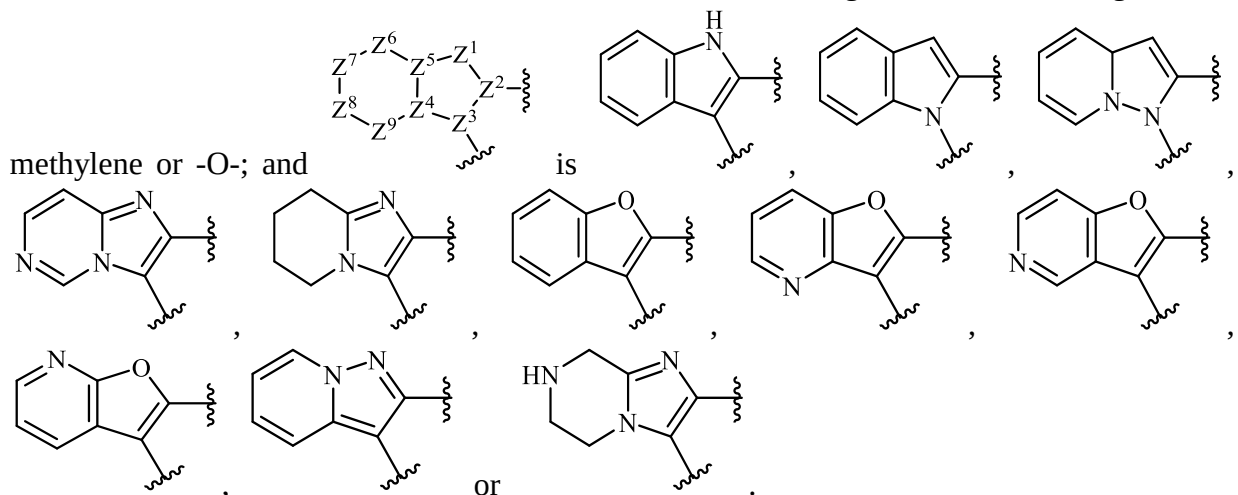


below:

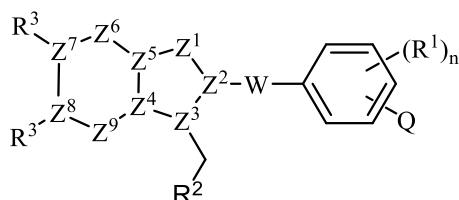
I-A-2

, wherein $R^{1'}$ has the definition as described

for R^1 , and $R^{1'}$ is the same as or different from R^1 ; W is a single bond; X is a single bond,

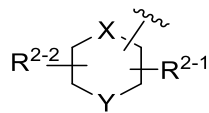


In a preferred embodiment of the present invention, the heterocyclic compound of formula I is of formula I-B as shown below:



I-B

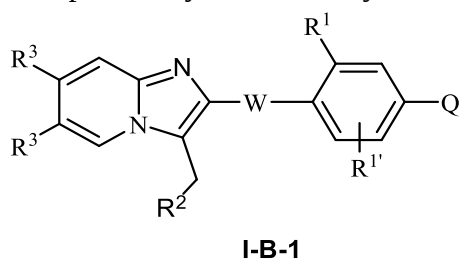
, wherein R^2 is



or $-C(=O)N(R^{2-5}R^{2-6})$

10 ⁶), X or Y is $-N(R^{2-3})-$;

preferably, the heterocyclic compound of formula I is of formula I-B-1 as shown below:

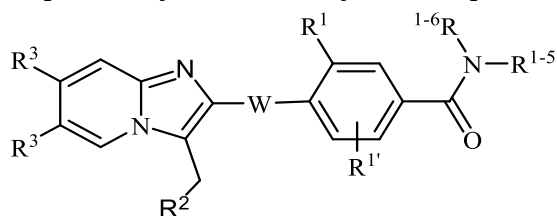


I-B-1

, wherein $R^{1'}$ has the definition as described for R^1 , and $R^{1'}$

is the same as or different from R^1 ; W is a single bond; and R^2 is $C(=O)N(R^{2-5}R^{2-6})$, wherein X or Y is $-N(R^{2-3})-$; and

more preferably, the heterocyclic compound of formula I is of formula I-B-2 as shown



below:

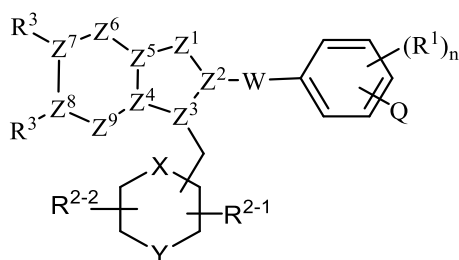
I-B-2

, wherein $R^{1'}$ has the definition as described

for R^1 , and $R^{1'}$ is the same as or different from R^1 ; W is a single bond; and R^2 is or $-C(=O)N(R^{2-5}R^{2-6})$, wherein X or Y is $-N(R^{2-3})-$.

In a preferred embodiment of the present invention, in the heterocyclic compound of formula I, Q is H, halogen, $-OH$, $-CF_3$, $-CN$, $-NH-C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-S(=O)-N(R^{1-11}R^{1-12})$, $-O-C(=O)-R^{1-13}$, $-N(R^{1-14}R^{1-15})$, $-NH-S(=O)_2-R^{1-16}$, an unsubstituted or substituted C_1-C_6 alkyl, an unsubstituted or substituted C_1-C_6 alkyl-O-, an unsubstituted or substituted C_2-6 alkenyl, an unsubstituted or substituted C_2-6 alkenyl-O-, an unsubstituted or substituted C_3-C_{10} cycloalkyl, an unsubstituted or substituted C_3-C_9 heterocycloalkyl, an unsubstituted or substituted C_3-C_9 heterocycloalkenyl, an unsubstituted or substituted C_6-C_{10} aryl, or an unsubstituted or substituted 5-10 membered heteroaryl.

In a preferred embodiment of the present invention, the heterocyclic compound of formula I is of formula I-C as shown below:

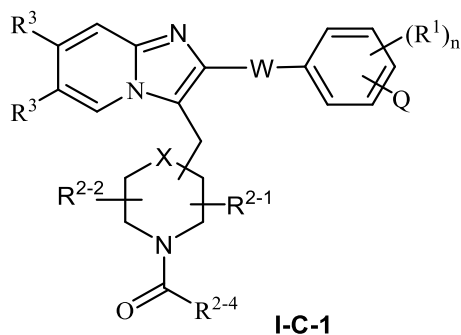


I-C

, wherein Q is H, halogen, $-OH$, $-CF_3$, $-CN$, $-NH-C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-S(=O)-N(R^{1-11}R^{1-12})$, $-O-C(=O)-R^{1-13}$, $-N(R^{1-14}R^{1-15})$, $-NH-S(=O)_2-R^{1-16}$, an unsubstituted or substituted C_1-C_6 alkyl, an unsubstituted or substituted C_1-C_6 alkyl-O-, an unsubstituted or substituted C_2-6 alkenyl, an unsubstituted or substituted C_2-6 alkenyl-O-, an unsubstituted or substituted C_3-C_{10} cycloalkyl, an unsubstituted or substituted C_3-C_9 heterocycloalkyl, an

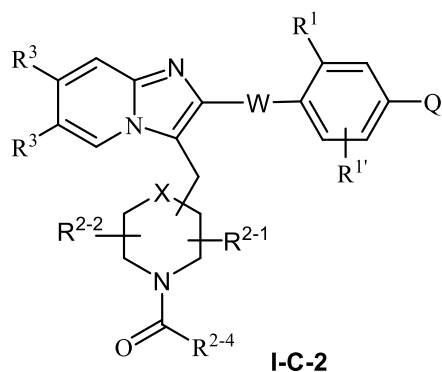
unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an unsubstituted or substituted 5-10 membered heteroaryl;

preferably, the heterocyclic compound of formula I is of formula I-C-1 as shown below:



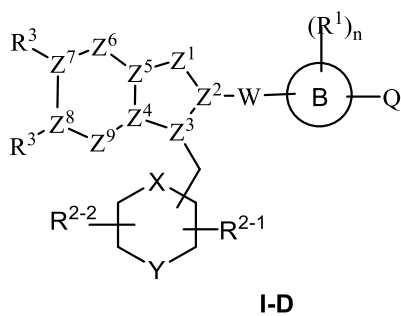
; and

5 more preferably, the heterocyclic compound of formula I is of formula I-C-2 as shown



below: , wherein R¹' has the definition as described for R¹, R¹' is the same as or different from R¹, and X is a single bond, methylene or -O-.

In a preferred embodiment of the present invention, the heterocyclic compound of formula I is of formula I-D as shown below:

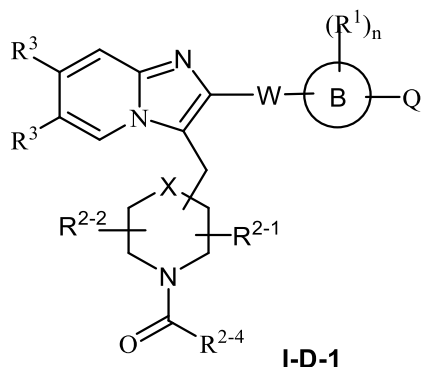


10

, wherein is a 5-6 membered heteroaryl or a C₃-C₅

heterocycloalkyl;

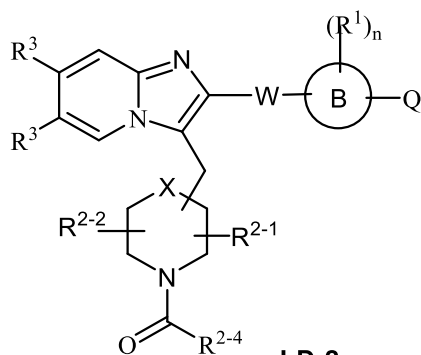
preferably, the heterocyclic compound of formula I is of formula I-D-1 as shown below:



I-D-1

, wherein $\textcircled{\text{B}}$ is a 5-6 membered heteroaryl or a C₃-C₅ heterocycloalkyl; and

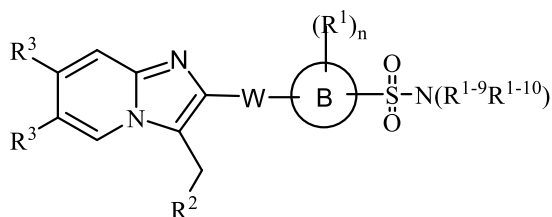
more preferably, the heterocyclic compound of formula I is of formula I-D-2 as shown



I-D-2

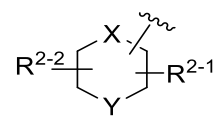
below: $\textcircled{\text{B}}$ is a 5-6 membered heteroaryl or a C₃-C₅ heterocycloalkyl.

In a preferred embodiment of the present invention, the heterocyclic compound of formula I is of formula I-E as shown below:

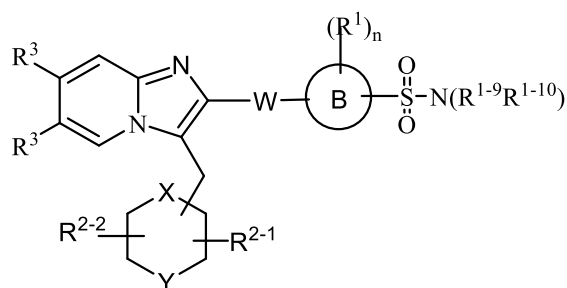


I-E

; wherein R² is independently



or an unsubstituted or substituted 5-6 membered heteroaryl; for example, W is a single bond; preferably, the heterocyclic compound of formula I is of formula I-E-1 as shown below:

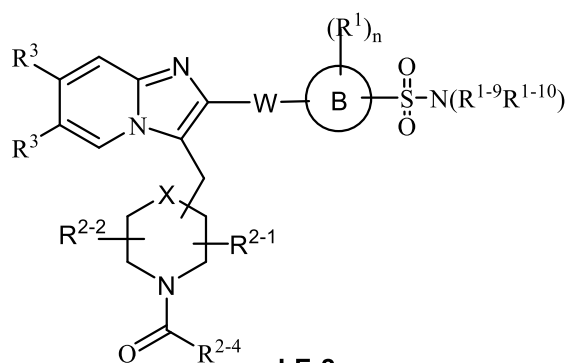


I-E-1

; for example,  is phenyl or 6 membered

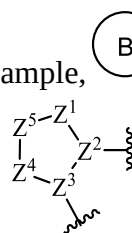
heteroaryl; and

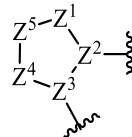
more preferably, the heterocyclic compound of formula I of formula I-E-2 as shown below:

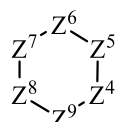


I-E-2

; for example,  is phenyl.



5 In a preferred embodiment of the present invention,  is a 5 membered heteroaryl;



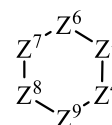
is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl,

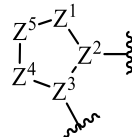
or a 5-6 membered heteroaryl, wherein the 5-6 membered heterocycloalkyl contains 1-3

heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl

contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered

10 heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



$Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8$ and Z^9 independently represent a ring atom; and when  is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z^6 or Z^9 represents a single bond;

m is 1, 2, 3 or 4;

15 R^3 is H, halogen, -OH, a C_1 - C_6 haloalkyl or a C_1 - C_6 alkyl;

W is a single bond or $-C(=O)-NH-$;

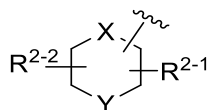


is phenyl, a 5-6 membered heteroaryl or a C₃-C₉ heterocycloalkyl;

n is 1, 2, 3 or 4;

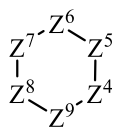
R¹ is independently H, halogen, -OH, -CN or an unsubstituted or substituted C₁-C₆ alkyl;

Q is H, halogen, -OH, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl;



R² is independently , -C(=O)N(R²⁻⁵R²⁻⁶) or an unsubstituted or substituted

5-6 membered heteroaryl;



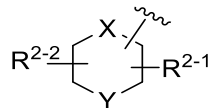
preferably, is phenyl, a 6 membered heterocycloalkyl, a 6 membered heterocycloalkenyl, or a 6 membered heteroaryl;

m is 1 or 2;

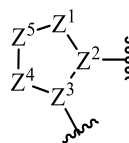
the substituent in R¹ is independently one or more selected from the following substituents:

halogen, -OH, -CN or -C(=O)NH₂; and preferably halogen.

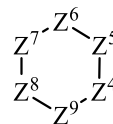
R³ is H, halogen, -OH, -CF₃ or a C₁-C₆ alkyl;



R² is independently or -C(=O)N(R²⁻⁵R²⁻⁶).



In a preferred embodiment of the present invention, is a 5 membered heteroaryl.



In a preferred embodiment of the present invention, is phenyl, a 6 membered

heterocycloalkyl, a 6 membered heterocycloalkenyl or a 6 membered heteroaryl.

In a preferred embodiment of the present invention, m is 1 or 2.

In a preferred embodiment of the present invention, R³ is H, halogen, -OH, a C₁-C₆ haloalkyl or a C₁-C₆ alkyl; and preferably, R³ is H, halogen, -OH, -CF₃ or a C₁-C₆ alkyl.

In a preferred embodiment of the present invention, W is a single bond.

In a preferred embodiment of the present invention, n is 1 or 2.

In a preferred embodiment of the present invention, R^1 is independently H, halogen, -OH, -CN, or an unsubstituted or substituted C_1 - C_6 alkyl, and preferably H, halogen, or an unsubstituted or substituted C_1 - C_6 alkyl.

In a preferred embodiment of the present invention, the substituent in R^1 is independently one or more selected from the following substituents: halogen, -OH, -CN or $-C(=O)NH_2$; and preferably halogen.

In a preferred embodiment of the present invention, Q is H, halogen, -OH, -CN, -NH- $C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=O)-N(R^{1-5}R^{1-6})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-N(R^{1-14}R^{1-15})$, $-NH-S(=O)_2-R^{1-16}$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl.

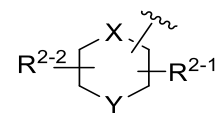
In a preferred embodiment of the present invention, Q is H, halogen, -OH, $-CF_3$, -CN, -NH- $C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=O)-N(R^{1-5}R^{1-6})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-S(=O)-N(R^{1-11}R^{1-12})$, $-O-C(=O)-R^{1-13}$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_1 - C_6 alkyl-O-, an unsubstituted or substituted C_2 - C_6 alkenyl, an unsubstituted or substituted C_2 - C_6 alkenyl-O-, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, an unsubstituted or substituted C_6 - C_{10} aryl, or an unsubstituted or substituted 5-6 membered heteroaryl; and

preferably, Q is H, halogen, -OH, -CN, -NH- $C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=O)-N(R^{1-5}R^{1-6})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl.

In a preferred embodiment of the present invention, the substituted C_1 - C_6 alkyl, the substituted C_1 - C_6 alkyl-O-, the substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6 alkenyl-O-, the substituted C_6 - C_{10} aryl, the substituted C_3 - C_5 heterocycloalkyl, the substituted C_3 - C_5 heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, $-CF_3$, -CN, -COOH, $-C(=O)NH_2$, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or a C_3 - C_6 cycloalkyl that is unsubstituted or substituted with one or more R^a , wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, $-CF_3$, -CN, -COOH or $-C(=O)NH_2$; and when a plurality of substituents are present, they are the same or different.

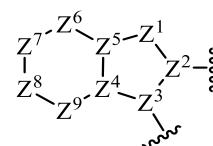
In a preferred embodiment of the present invention, R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} and R^{1-13} are each independently H, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_1 - C_6 alkyl-O-, an unsubstituted or substituted C_2 - C_6 alkenyl, an unsubstituted or substituted C_2 - C_6 alkenyl-O-, an unsubstituted or substituted C_3 - C_6 cycloalkyl, an unsubstituted or substituted C_3 - C_5 heterocycloalkyl, an unsubstituted or substituted C_3 - C_5 heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein the C_3 - C_5 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_5 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C_1 - C_6 alkyl, the substituted C_1 - C_6 alkyl-O-, the substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6 alkenyl-O-, the substituted C_3 - C_6 cycloalkyl, the substituted C_3 - C_5 heterocycloalkyl, the substituted C_3 - C_5 heterocycloalkenyl, the substituted phenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, - CF_3 , -CN, a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O-, a C_3 - C_6 cycloalkyl, -COOH or =O, wherein when a plurality of substituents are present, they are the same or different.

In a preferred embodiment of the present invention, R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form an unsubstituted or substituted C_3 - C_5 heterocycloalkyl, an unsubstituted or substituted C_3 - C_5 heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein the C_3 - C_5 heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C_3 - C_5 heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the 5-6 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C_3 - C_5 heterocycloalkyl, the substituted C_3 - C_5 heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, - CF_3 , -CN, a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O-, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or a C_3 - C_6 cycloalkyl that is unsubstituted or substituted with one or more R^a , wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, - CF_3 , -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different.

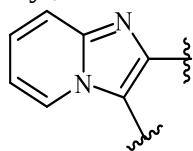


In a preferred embodiment of the present invention, R^2 is independently or $-C(=O)N(R^{2-5}R^{2-6})$.

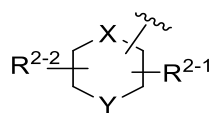
In a preferred embodiment of the present invention, when Q is H, halogen, $-OH$, $-CF_3$, $-CN$, $-NH-C(=O)-R^{1-1}$, $-C(=O)-R^{1-2}$, $-O-C(=O)-N(R^{1-3}R^{1-4})$, $-C(=NH)-N(R^{1-7}R^{1-8})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-S(=O)-N(R^{1-11}R^{1-12})$, $-O-C(=O)-R^{1-13}$, $-N(R^{1-14}R^{1-15})$, $-NH-S(=O)_2-R^{1-16}$, an unsubstituted or substituted C_1-C_6 alkyl, an unsubstituted or substituted C_1-C_6 alkyl-O-, an unsubstituted or substituted C_2-6 alkenyl, an unsubstituted or substituted C_2-C_6 alkenyl-O-, an unsubstituted or substituted C_3-C_{10} cycloalkyl, an unsubstituted or substituted C_3-C_9 heterocycloalkyl, an unsubstituted or substituted C_3-C_9 heterocycloalkenyl, an unsubstituted or substituted C_6-C_{10}



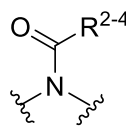
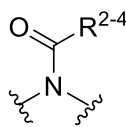
10 aryl, or an unsubstituted or substituted 5-10 membered heteroaryl,



, or R^2 is

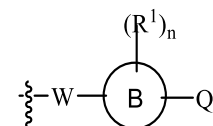


, wherein X and Y are each independently a single

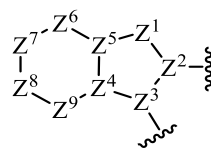
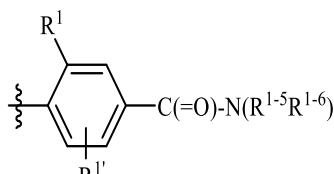


bond, methylene, $-O-$ or

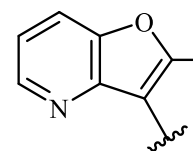
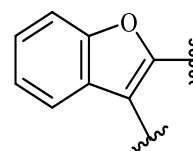
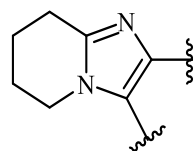
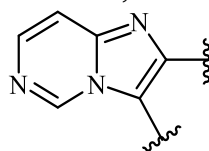
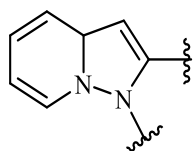
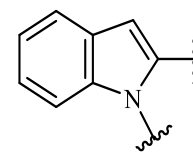
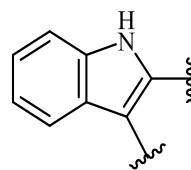
, e.g., .



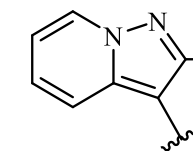
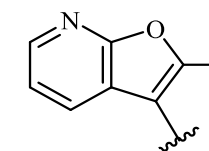
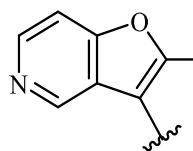
In a preferred embodiment of the present invention, when



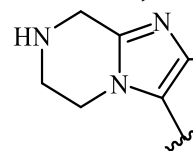
is



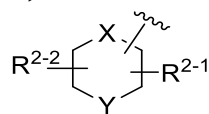
15



or



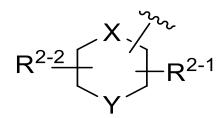
, or R^2 is



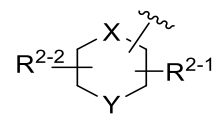
independently

, $-C(=O)N(R^{2-5}R^{2-6})$ or an unsubstituted or substituted 5-6 membered heteroaryl, wherein X or Y is $-N(R^{2-3})-$.

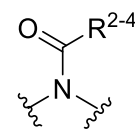
In a preferred embodiment of the present invention, R^2 is independently $-C(=O)N(R^{2-5}R^{2-6})$, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein X or Y is $-N(R^{2-3})-$.



In a preferred embodiment of the present invention, R^2 is independently

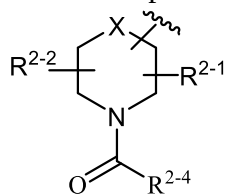


preferably, X and Y are each independently a single bond, methylene, $-O-$, or

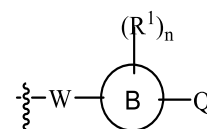


; and

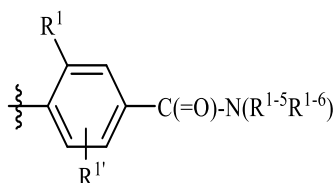
more preferably, R^2 is



In a preferred embodiment of the present invention,



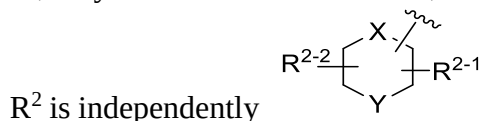
is



In a preferred embodiment of the present invention, R^3 is halogen, a C_1 - C_6 haloalkyl or a C_1 - C_6 alkyl;

Q is $-NH-C(=O)-R^{1-1}$, $-C(=O)-N(R^{1-5}R^{1-6})$, $-S(=O)_2-N(R^{1-9}R^{1-10})$, $-N(R^{1-14}R^{1-15})$, $-NH-S(=O)_2-R^{1-16}$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C_1 - C_6 alkyl, the substituted C_3 - C_{10} cycloalkyl, the substituted C_3 - C_9 heterocycloalkyl, the substituted C_3 - C_9 heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: $-OH$, $-CF_3$, $-CN$, $-N(R^{1-21}R^{1-22})$, $-C(=O)-N(R^{1-23}R^{1-24})$, $-S(=O)_2-N(R^{1-25}R^{1-26})$, $-C(=O)-O-R^{1-27}$, $=O$ (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or a C_3 - C_6 cycloalkyl that is unsubstituted

or substituted with one or more R^a , wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH or -COOH; and when a plurality of substituents are present, they are the same or different;

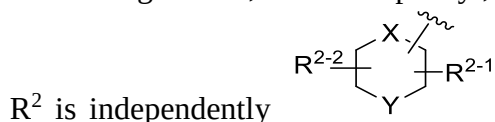


, -C(=O)N(R²⁻⁵R²⁻⁶) or an unsubstituted or substituted

5 5-6 membered heteroaryl; wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CN, an unsubstituted or halogen-substituted C₁-C₆ alkyl, an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-, -C(=O)R²⁻⁷ or -C(=O)N(R²⁻⁸R²⁻⁹), wherein when a plurality of
10 substituents are present, they are the same or different;

preferably, Q is -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰);

W is a single bond;  is phenyl; and



, or an unsubstituted or substituted 5-6 membered heteroaryl.

15 In a preferred embodiment of the present invention, R³ is halogen, a C₁-C₆ haloalkyl or a C₁-C₆ alkyl;

Q is -NH-C(=O)-R¹⁻¹, -C(=O)-N(R¹⁻⁵R¹⁻⁶), -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉
20 heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉
25 heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: -OH, -CF₃, -CN, -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted
30 with one or more R^a , wherein when a plurality of R^a are present, they are the same or different,

and each R^a is halogen, -OH or -COOH; and when a plurality of substituents are present, they are the same or different;

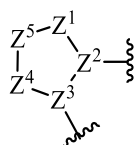
R^{1-1} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each independently H, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_2 - C_6 alkenyl, an unsubstituted or substituted C_2 - C_6 alkynyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_5 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C_1 - C_6 alkyl, the substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6 alkynyl, the substituted C_3 - C_{10} cycloalkyl, the substituted C_3 - C_9 heterocycloalkyl, the substituted phenyl, or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, -COOCH₃, -NH₂, =O, and a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or a C_3 - C_6 cycloalkyl that is unsubstituted or substituted with one or more R^d , wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different.

In a preferred embodiment of the present invention, R^3 is halogen, -CHF₂, -CH₂F or a C_1 - C_6 alkyl;

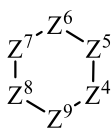
Q is -NH-C(=O)- R^{1-1} , -N(R^{1-14} R^{1-15}), -NH-S(=O)₂- R^{1-16} , an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C_1 - C_6 alkyl, the substituted C_3 - C_{10} cycloalkyl, the substituted C_3 - C_9 heterocycloalkyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: -N(R^{1-21} R^{1-22}), -C(=O)-N(R^{1-23} R^{1-24}), -S(=O)₂-N(R^{1-25} R^{1-26}) or -C(=O)-O- R^{1-27} , wherein when a plurality of substituents are present, they are the same or different;

R^{1-1} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each independently H, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_2 - C_6 alkynyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the

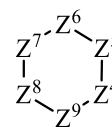
5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₂-C₆ alkynyl, the substituted C₃-C₉ heterocycloalkyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CN, =O, -COOH, -C(=O)NH₂, -COOCH₃, -NH₂, and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d, wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different.



In a preferred embodiment of the present invention, is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl, wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



Z¹, Z², Z³, Z⁴, Z⁵, Z⁶, Z⁷, Z⁸ and Z⁹ independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z⁶ or Z⁹ represents a single bond;

m is 1, 2, 3 or 4;

R³ is H, halogen, -OH, -CF₃, -CN or a C₁-C₆ alkyl; when m is not 1, R³ is independently the same or different; or when two adjacent R³ are present, they, together with the ring atom to which they directly link, form a C₃-C₆ cycloalkyl, a C₃-C₆ cycloalkenyl, a C₃-C₅ heterocycloalkyl, a C₃-C₅ heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C₃-C₅ heterocycloalkyl, wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

5 n is 1, 2, 3 or 4;

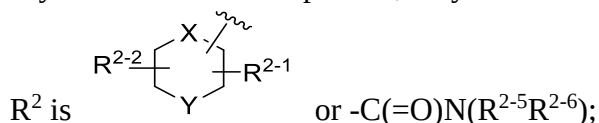
R¹ is independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); when a plurality of substituents are present, they are the same or different; and when n is not 1, R¹ is independently the same or different;

Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl; wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₆-C₁₀ aryl, the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² and R¹⁻¹³ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆

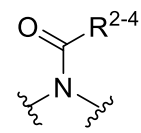
alkenyl-O-, an unsubstituted or substituted C₃-C₆ cycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-6 membered heteroaryl; wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₃-C₆ cycloalkyl, the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl, the substituted phenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, a C₃-C₆ cycloalkyl, -COOH or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), wherein when a plurality of substituents are present, they are the same or different; or

R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, or R¹⁻¹¹ and R¹⁻¹², independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl; wherein the C₃-C₅ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C₃-C₅ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, and the 5-6 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b, wherein when a plurality of R^b are present, they are the same or different and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;



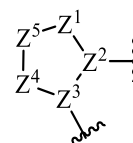
R²⁻¹ and R²⁻² are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens

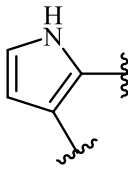
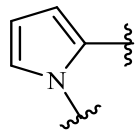
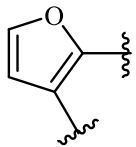
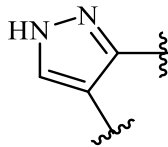
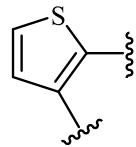
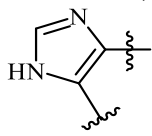
on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

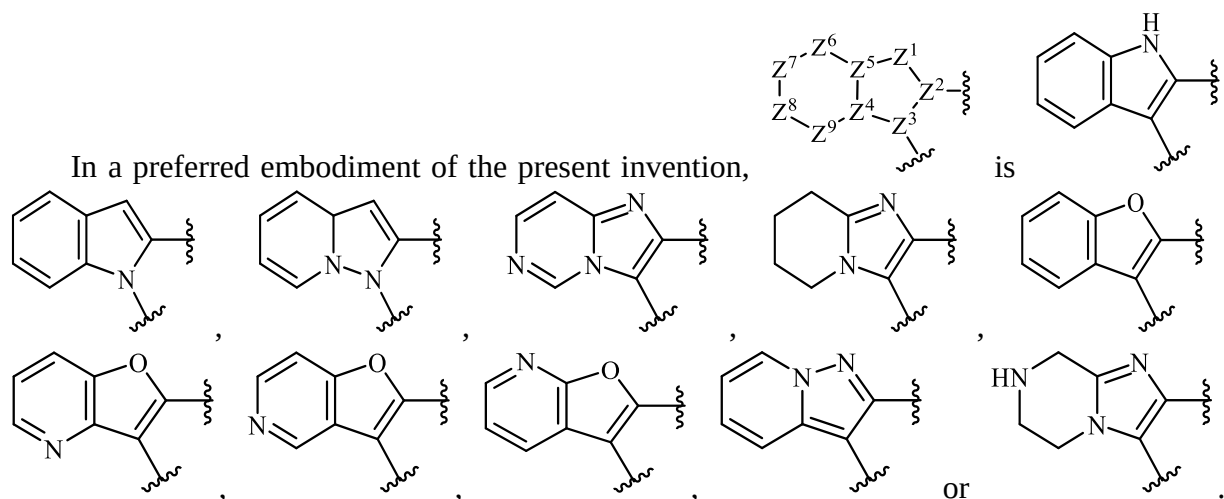


X and Y are each independently a single bond, methylene, -O-, -N(R²⁻³)- or

R²⁻³, R²⁻⁴, R²⁻⁵ and R²⁻⁶ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different.



In a preferred embodiment of the present invention, when is a 5 membered heteroaryl, the 5 membered heteroaryl is pyrrolyl (e.g., , or , furanyl (e.g., , pyrazolyl (e.g., , thienyl (e.g., ) or imidazolyl (e.g., ). (In the structural formula, the right-side bond is linked to W)



In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when  is a C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S, and is

preferably piperidinyI (e.g.,  (e.g., )). (In the structural formula, the left-side bond is linked to W, and the right-side bond is linked to Q)

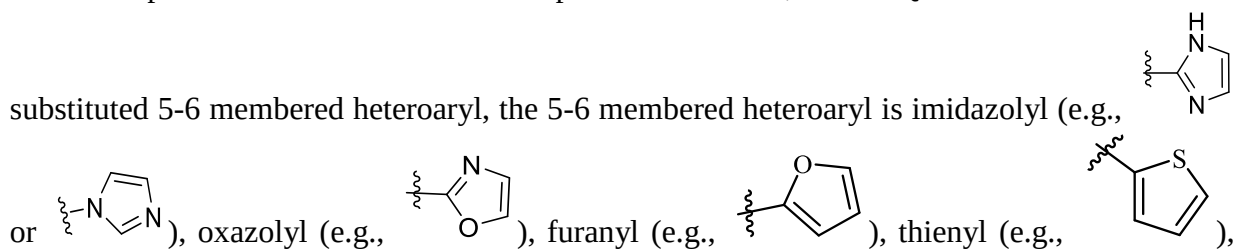
In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O

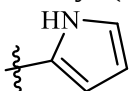
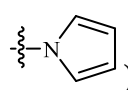
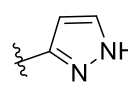
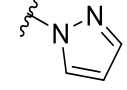
and S, and is preferably pyrrolidinyl (e.g.,) or piperazinyl (e.g.,)

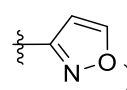
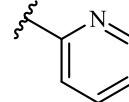
In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₄-C₅ heterocycloalkenyl is preferably 2,5-dihydro-

1*H*-pyrrolyl (e.g., ).

In a preferred embodiment of the present invention, when Q is an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl is imidazolyl (e.g.,



pyrrolyl (e.g.,  or , pyrazolyl (e.g.,  or , isoxazolyl

5 (e.g., , pyridinyl (e.g., ) or pyrimidinyl (e.g., 

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, the number of the substituents is

10 independently 1, 2, 3 or 4.

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is halogen, the

15 halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine.

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₁-C₆ alkyl

20 or a C₁-C₆ alkyl-O- substituted with one or more R^a, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) in the substituent is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl.

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₃-C₆ cycloalkyl substituted with one or more R^a, the C₃-C₆ cycloalkyl in the substituent (e.g., cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl) is cyclopropyl.

25

In a preferred embodiment of the present invention, when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, the substituent is a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, or a C₃-C₆ cycloalkyl substituted with one or more R^a, and R^a is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl), and preferably methyl or ethyl.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₂-C₆ alkenyl, or an unsubstituted or substituted C₂-C₆ alkenyl-O-, the C₂-C₆ alkenyl is a C₂-C₄ alkenyl {e.g., vinyl, propenyl (e.g., 1-propenyl or 2-propenyl), or butenyl (e.g., 2-butenyl, 1-butenyl or butadienyl)}, and preferably ethenyl.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or two of N, O and S.

In a preferred embodiment of the present invention, when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted phenyl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6-membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine, or iodine, and preferably fluorine.

In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C_1 - C_6 alkyl, a substituted C_1 - C_6 alkyl-O-, a substituted C_2 - C_6 alkenyl, a substituted C_2 - C_6 alkenyl-O-, a substituted phenyl, a substituted C_3 - C_5 heterocycloalkyl, a substituted C_3 - C_5 heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C_1 - C_6 alkyl or C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl in the substituent (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C_1 - C_4 alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C_1 - C_6 alkyl, a substituted C_1 - C_6 alkyl-O-, a substituted C_2 - C_6 alkenyl, a substituted C_2 - C_6 alkenyl-O-, a substituted phenyl, a substituted C_3 - C_5 heterocycloalkyl, a substituted C_3 - C_5 heterocycloalkenyl or a substituted 5-6 membered heteroaryl, and the substituent is a C_3 - C_6 cycloalkyl, the C_3 - C_6 cycloalkyl in the substituent (e.g., cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl) is cyclopropyl.

In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C_1 - C_6 alkyl, a substituted C_1 - C_6 alkyl-O-, a substituted C_2 - C_6 alkenyl, a substituted C_2 - C_6 alkenyl-O-, a substituted phenyl, a substituted C_3 - C_5 heterocycloalkyl, a substituted C_3 - C_5 heterocycloalkenyl or a substituted 5-6 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form an unsubstituted or substituted C_3 - C_5 heterocycloalkyl, the C_3 - C_5 heterocycloalkyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form an unsubstituted or substituted C_3 - C_5 heterocycloalkenyl, the C_3 - C_5 heterocycloalkenyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form an unsubstituted or substituted 5-6 membered heteroaryl, the unsubstituted or substituted 5-6 membered heteroaryl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine, or iodine.

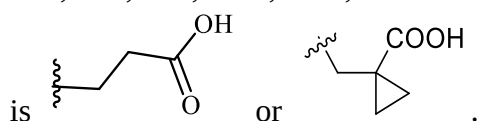
In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₁-C₆ alkyl or a C₁-C₆ alkyl-O- substituted with one or more R^b , the C₁-C₆ alkyl in the substituent (e.g., methyl, ethyl, propyl, butyl, pentyl or hexyl) is independently a C₁-C₄ alkyl (e.g., methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl).

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₃-C₆ cycloalkyl substituted with one or more R^b , the C₃-C₆ cycloalkyl in the substituent is cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl.

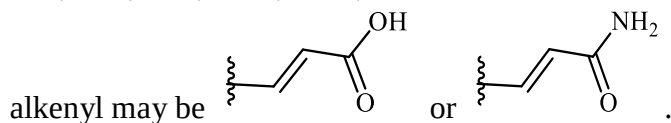
In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, the substituent is a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl substituted with one or more R^b , wherein when R^b is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

In a preferred embodiment of the present invention, when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl or a substituted 5-6 membered heteroaryl, the number of the substituent is 1, 2, 3, or 4 independently, and preferably 1 or 2.

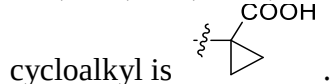
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl



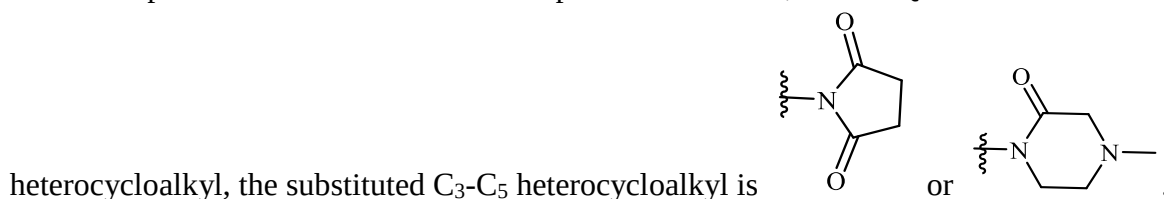
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C_2 - C_6 alkenyl, the substituted C_2 - C_6



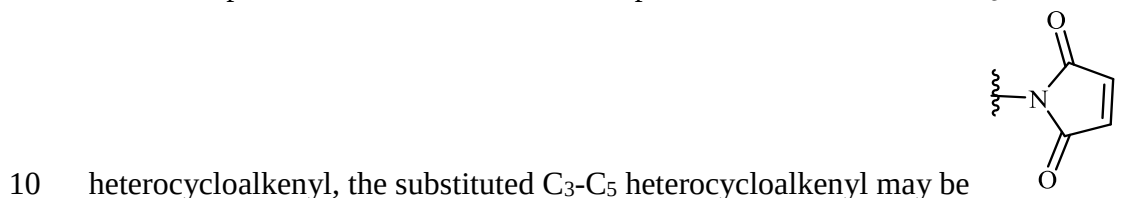
In a preferred embodiment of the present invention, when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C_3 - C_6 cycloalkyl, the substituted C_3 - C_6



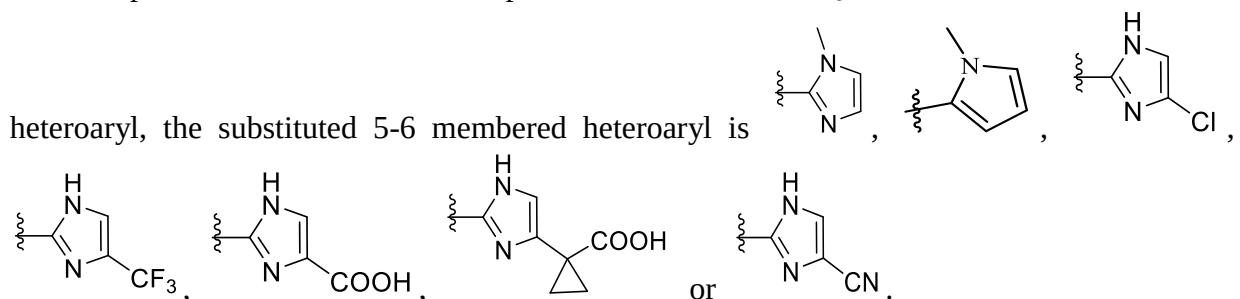
In a preferred embodiment of the present invention, when Q is a substituted C_3 - C_5



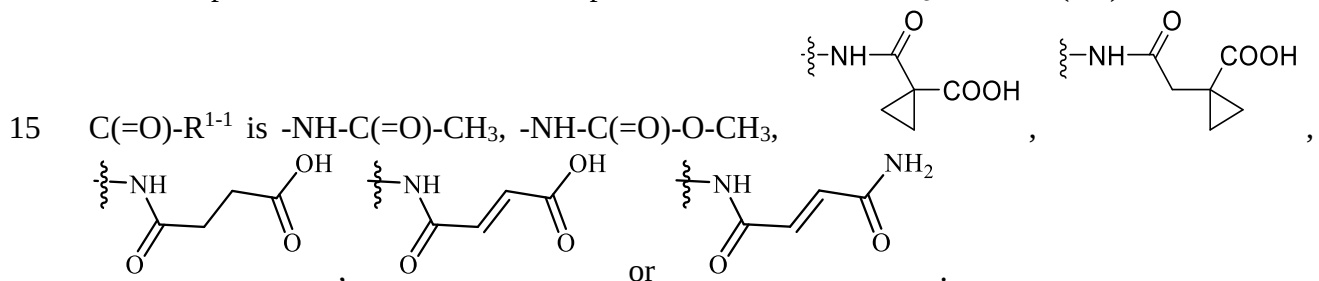
In a preferred embodiment of the present invention, when Q is a substituted C_3 - C_5



In a preferred embodiment of the present invention, when Q is a substituted 5-6 membered

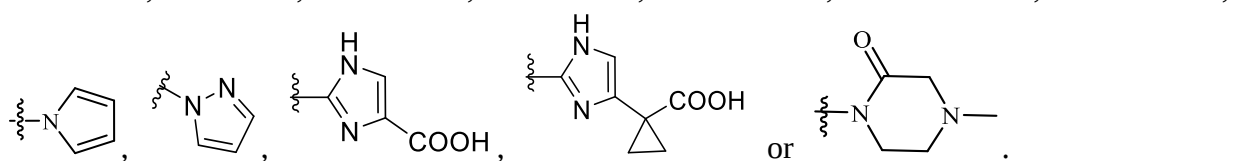
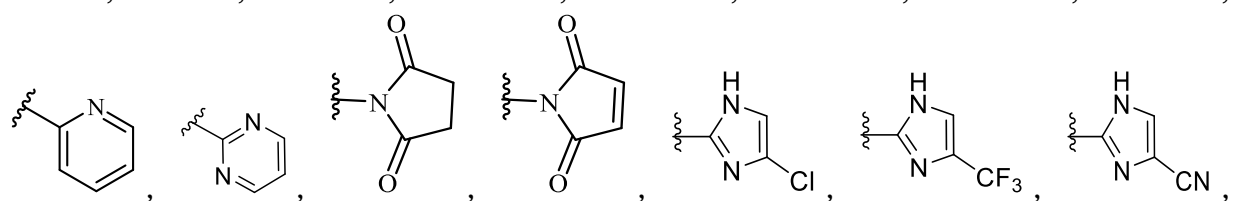
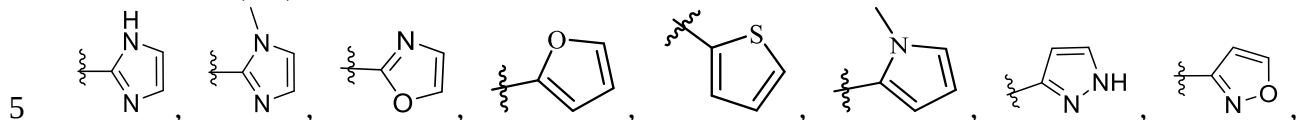
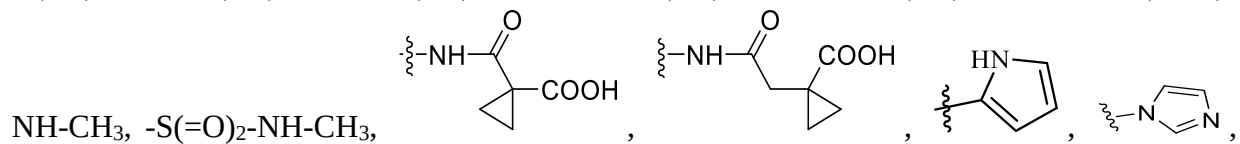
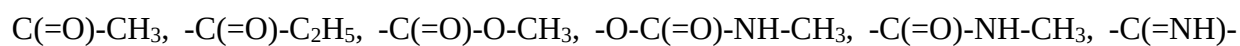
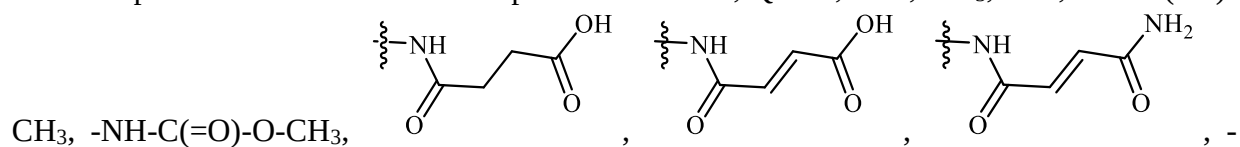


In a preferred embodiment of the present invention, when Q is $-NH-C(=O)-R^{1-1}$, the $-NH-$

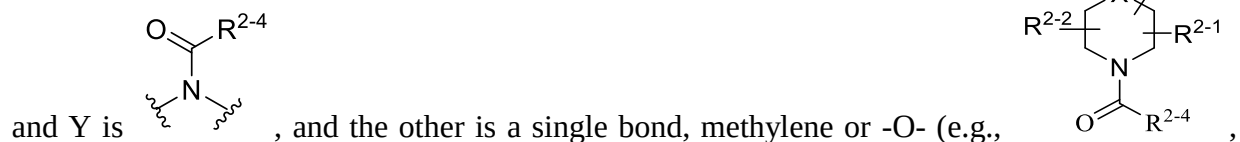
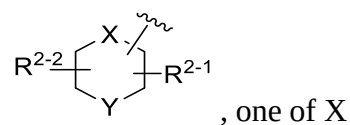


In a preferred embodiment of the present invention, when Q is $-S(=O)_2-N(R^{1-9}R^{1-10})$, the $-S(=O)_2-N(R^{1-9}R^{1-10})$ is $-S(=O)_2-NH-CH_3$.

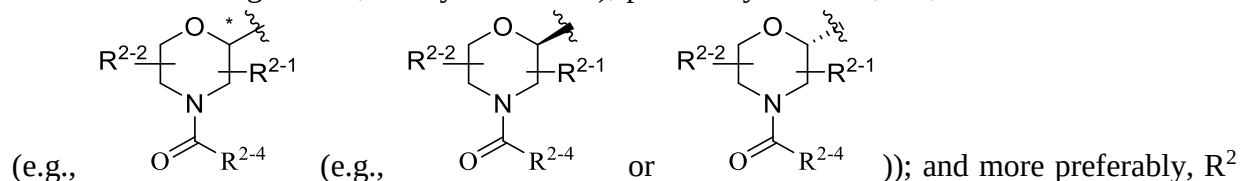
In a preferred embodiment of the present invention, Q is H, -OH, -CF₃, -CN, -NH-C(=O)-

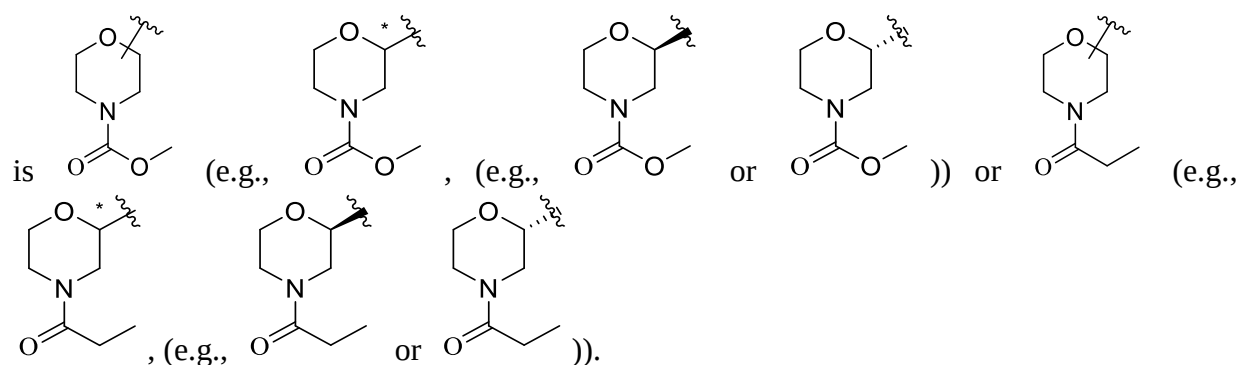


In a preferred embodiment of the present invention, when R² is

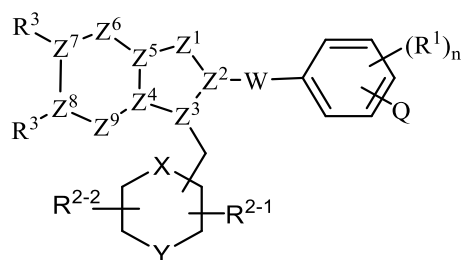


10 wherein X is a single bond, methylene or -O-; preferably X is -O-, i.e., R² is





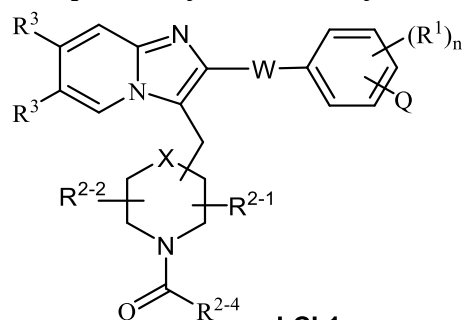
In a preferred embodiment of the present invention, the heterocyclic compound of formula I is of formula I-C' as shown below:



I-C'

, wherein Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl;

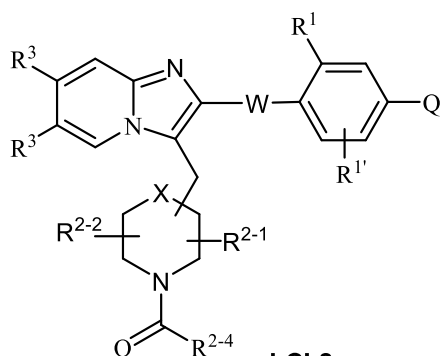
preferably, the heterocyclic compound of formula I is of formula I-C'-1 as shown below:



I-C'-1

, wherein Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl; and

more preferably, the heterocyclic compound of formula I is of formula I-C'-2 as shown



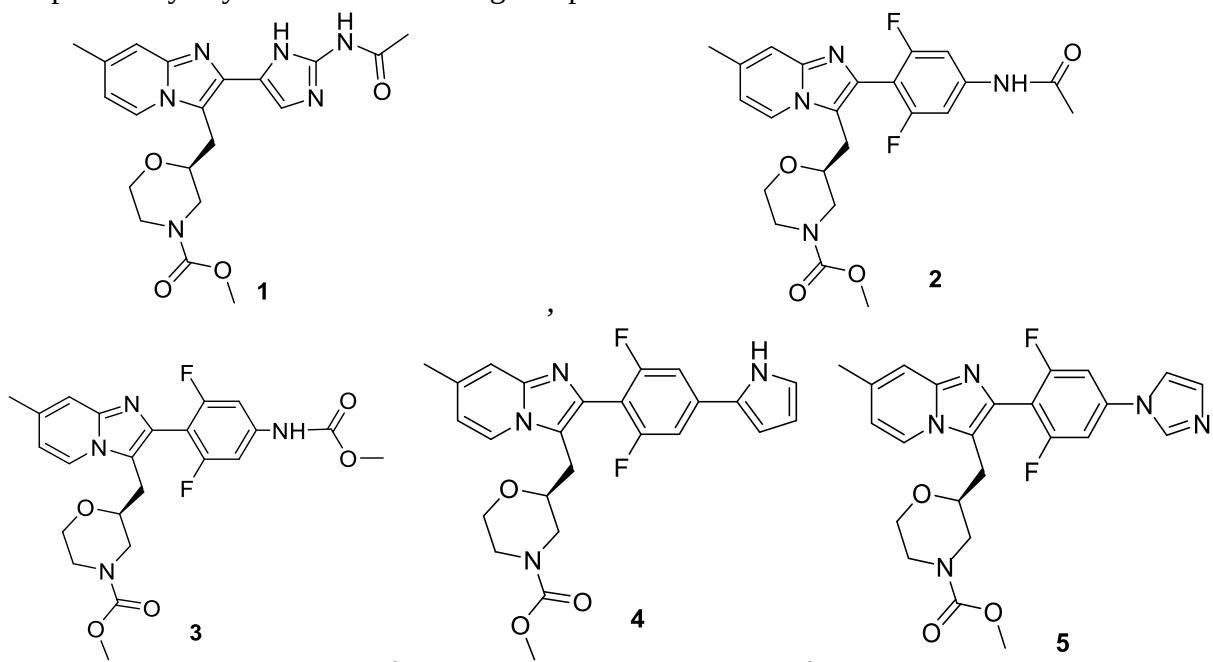
below:

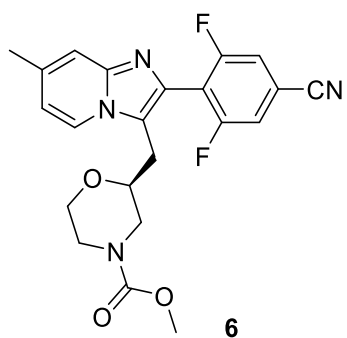
I-C'-2

, wherein $R^{1'}$ has the definition as described for R^1 ,

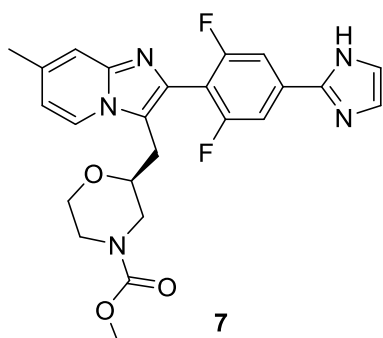
$R^{1'}$ is the same as or different from R^1 , and X is a single bond, methylene or -O-; Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-C(=O)- R^{1-1} , -C(=O)- R^{1-2} , -O-C(=O)-N(R^{1-3} R^{1-4}), -C(=NH)-N(R^{1-7} R^{1-8}), -S(=O)₂-N(R^{1-9} R^{1-10}), -S(=O)-N(R^{1-11} R^{1-12}), -O-C(=O)- R^{1-13} , an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl.

In a preferred embodiment of the present invention, the heterocyclic compound of formula I is preferably any one of the following compounds:

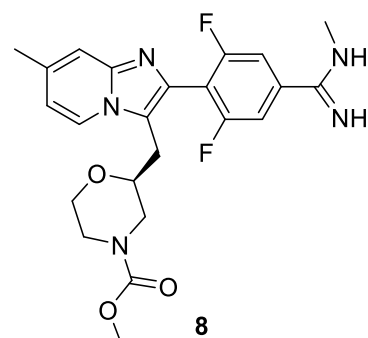




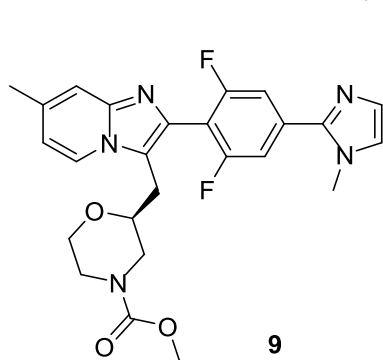
,



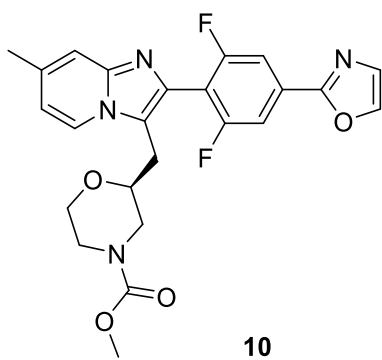
,



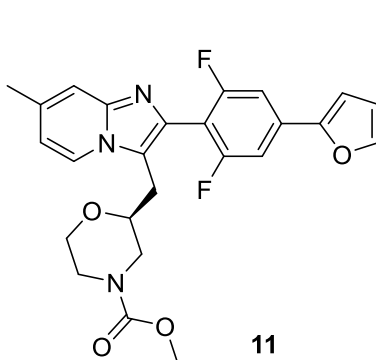
,



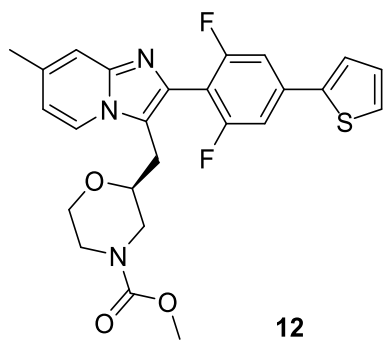
,



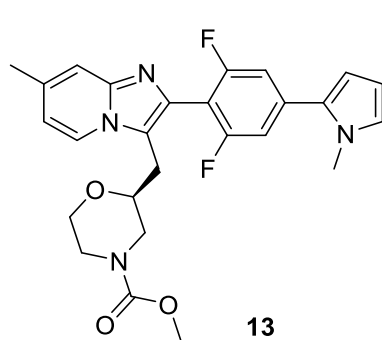
,



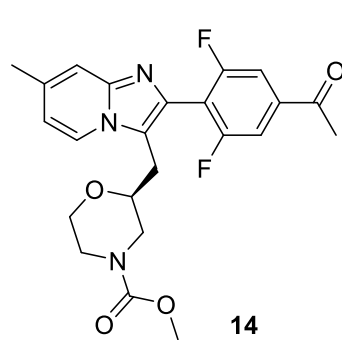
,



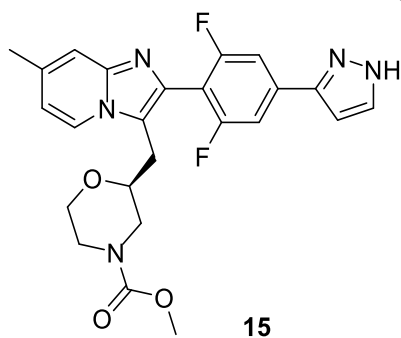
,



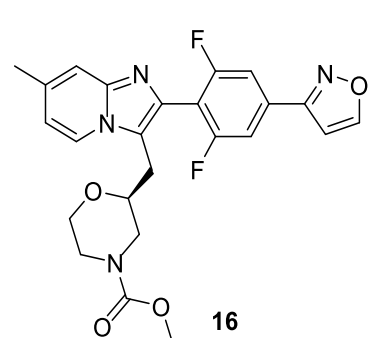
,



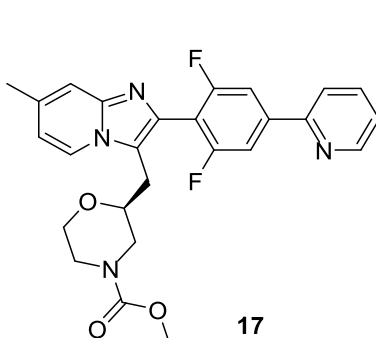
,



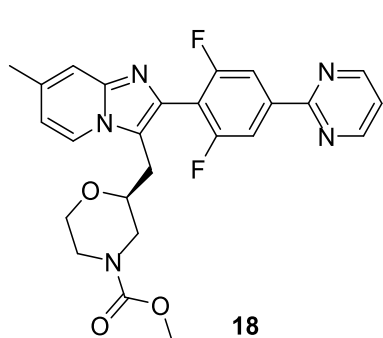
,



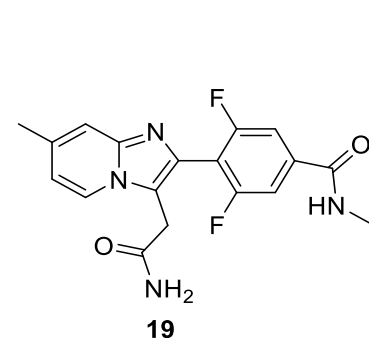
,



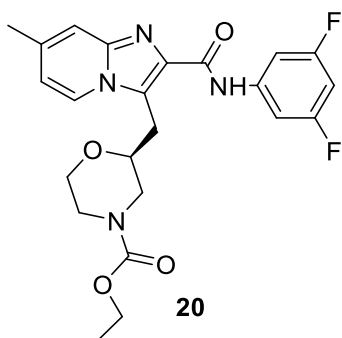
,



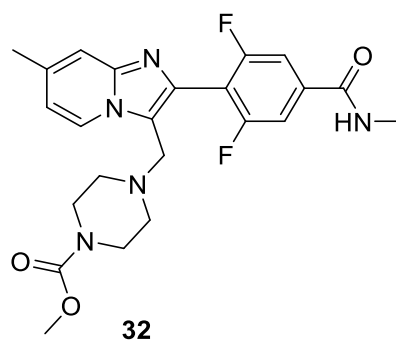
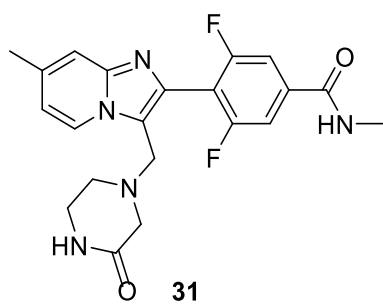
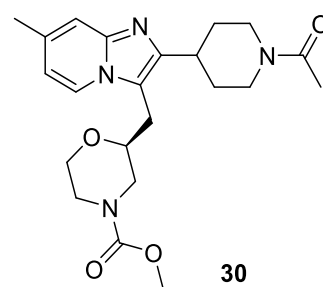
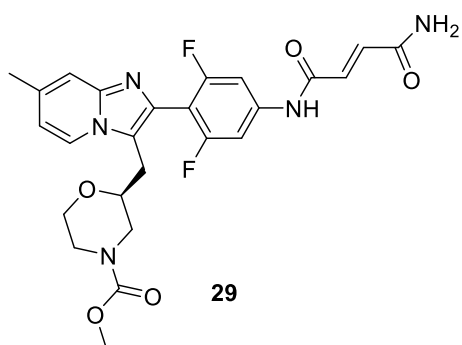
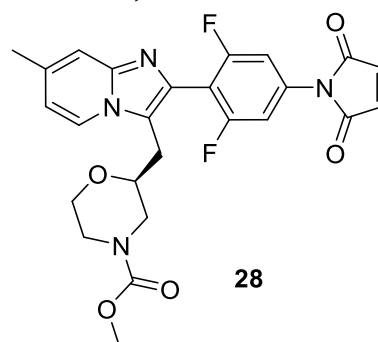
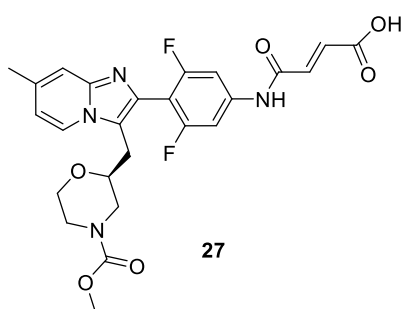
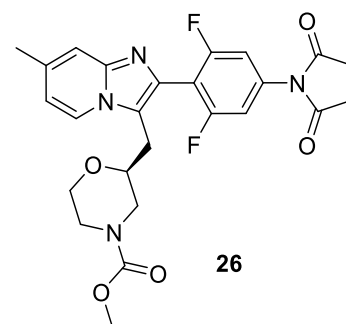
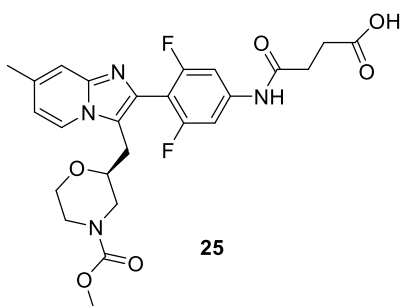
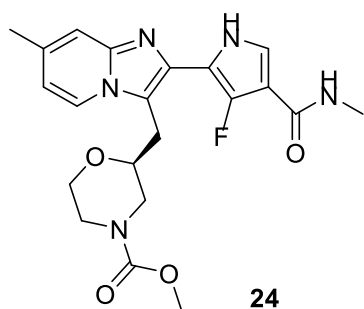
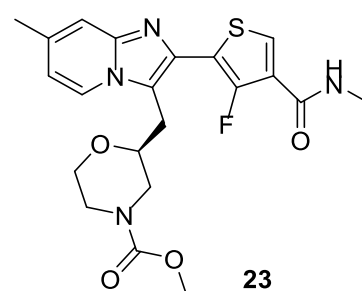
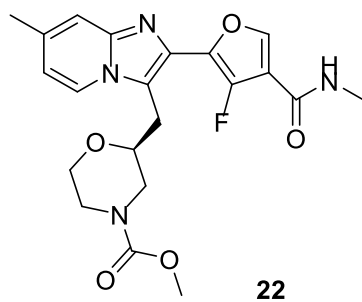
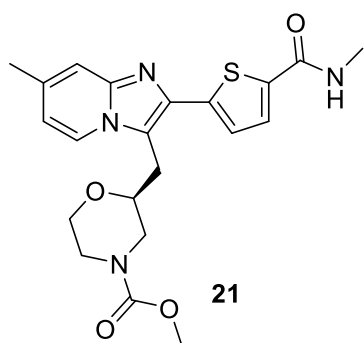
,

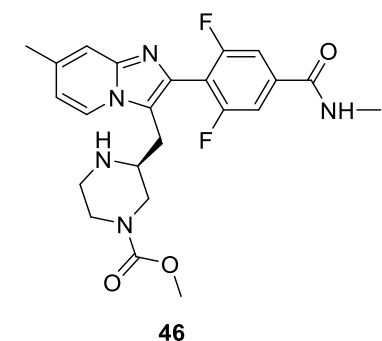
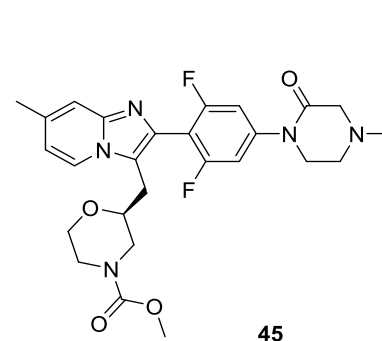
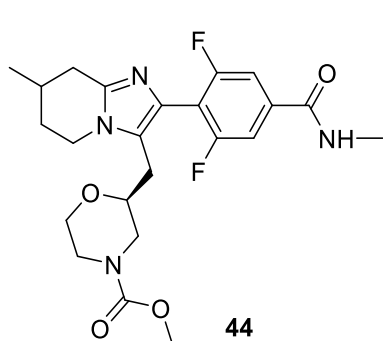
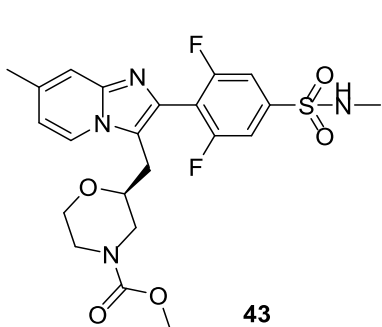
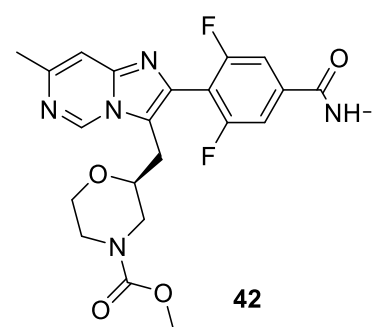
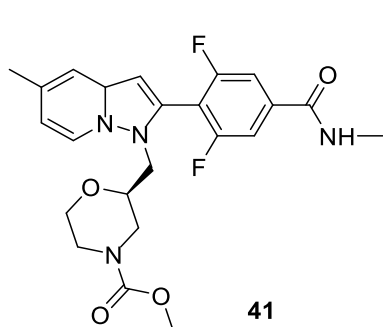
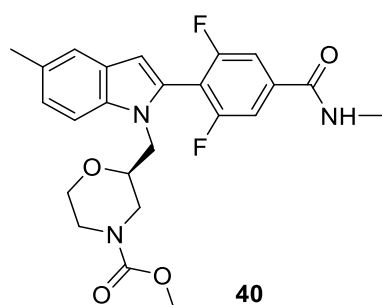
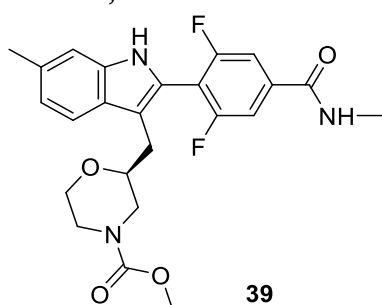
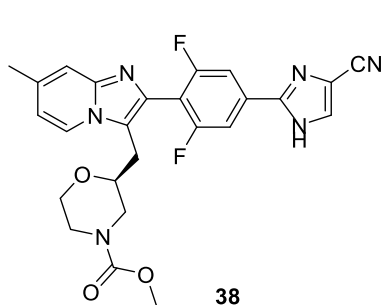
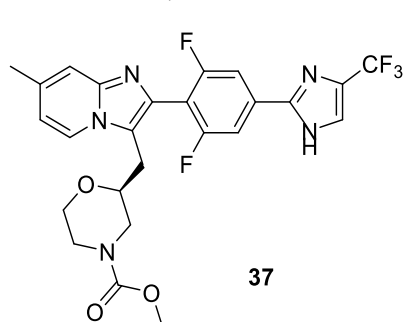
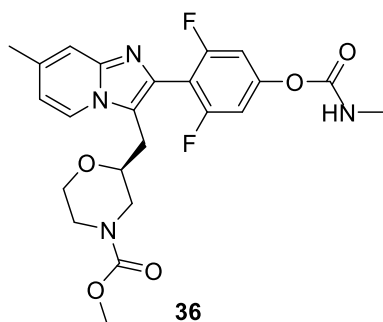
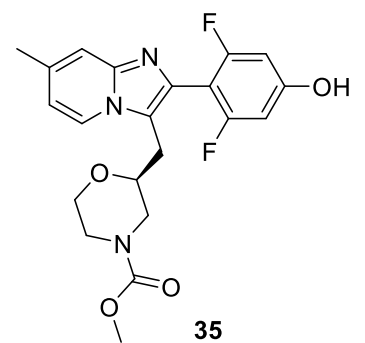
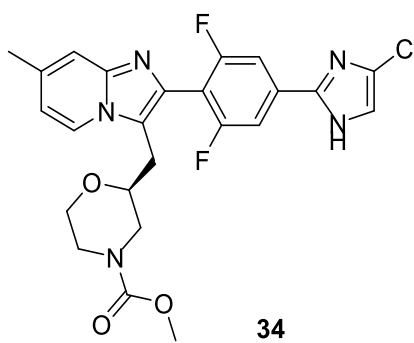
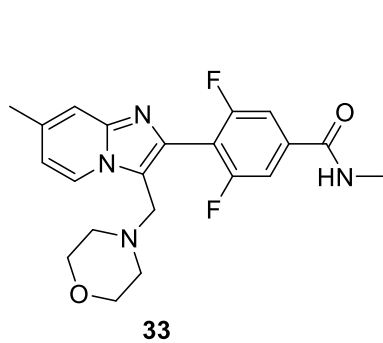


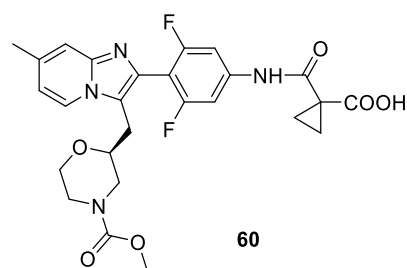
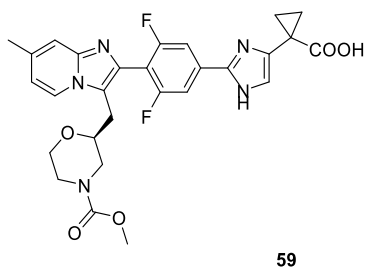
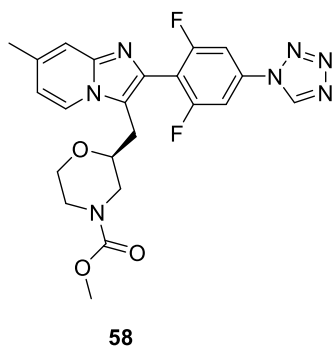
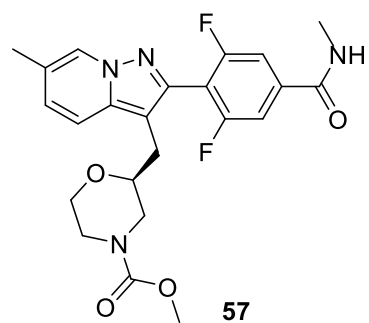
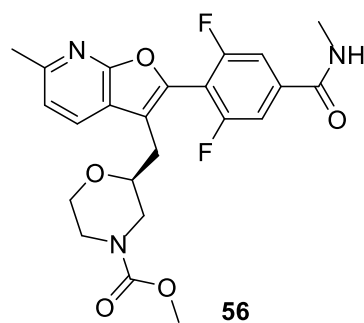
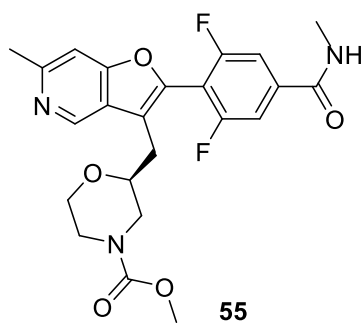
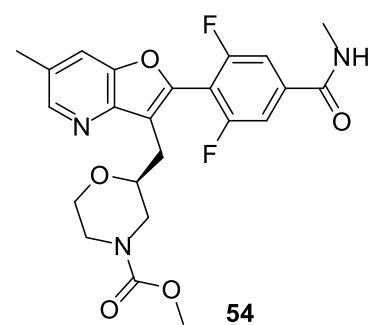
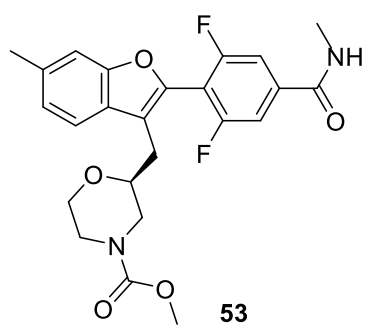
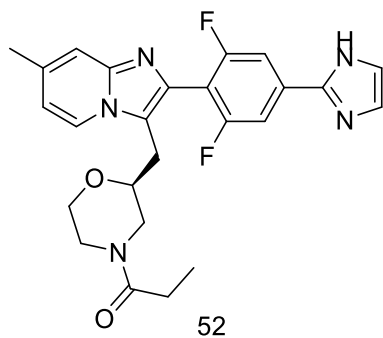
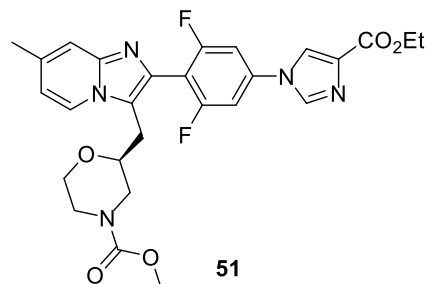
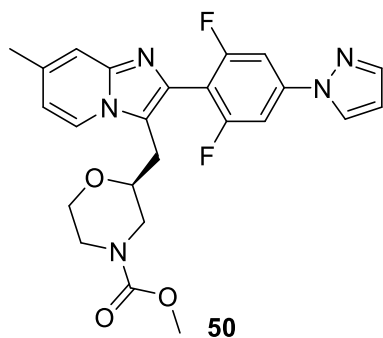
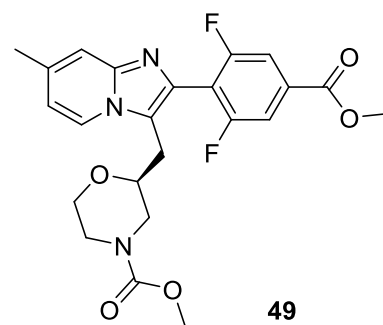
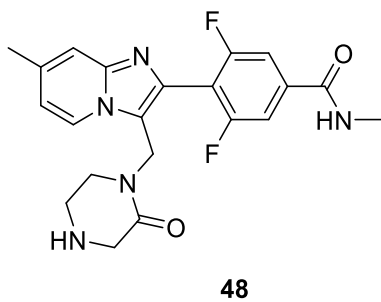
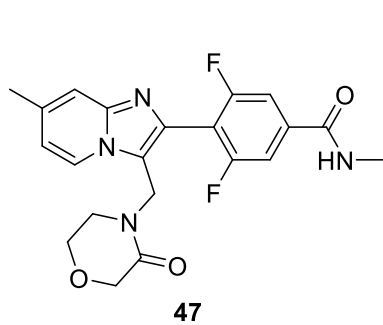
,

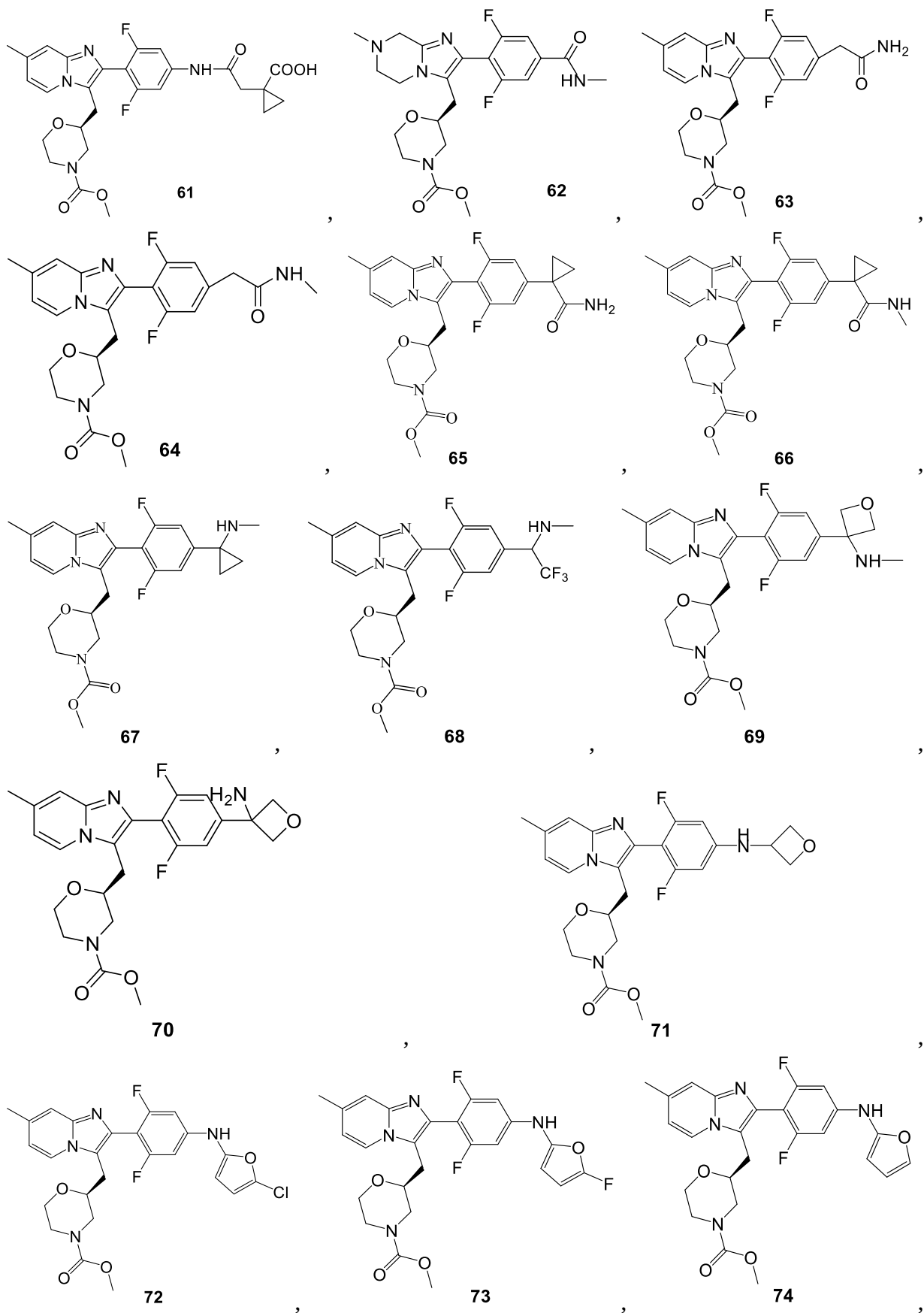


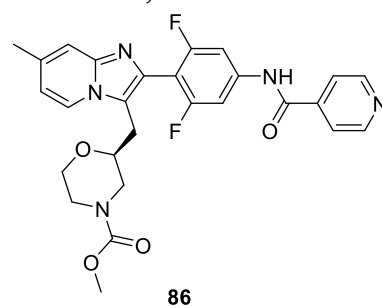
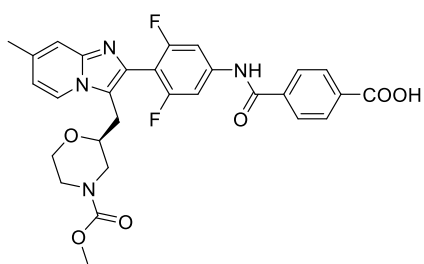
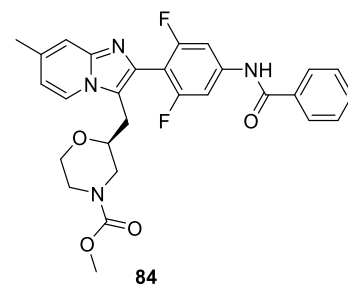
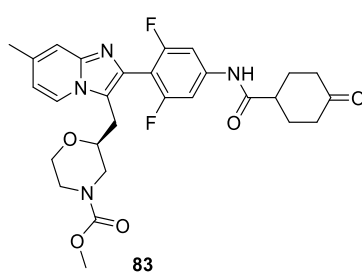
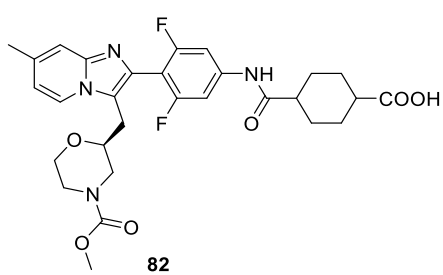
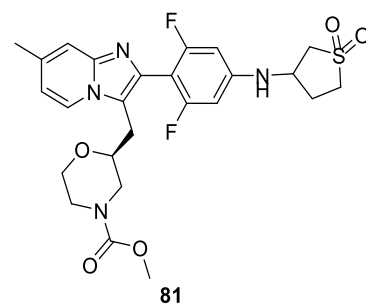
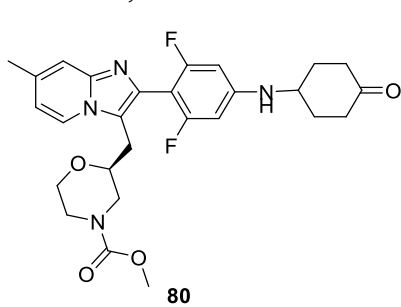
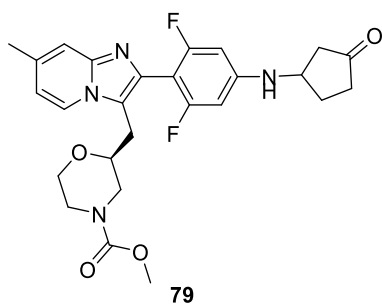
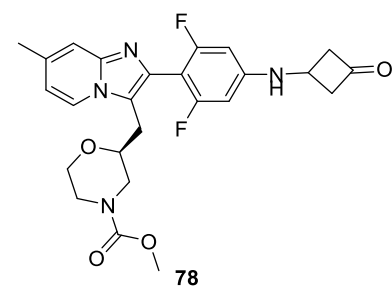
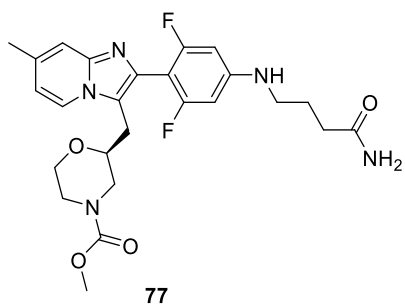
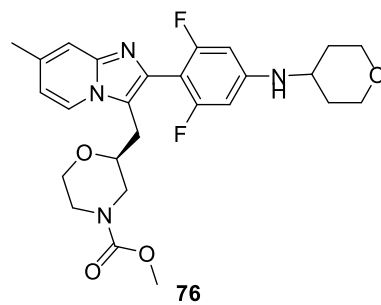
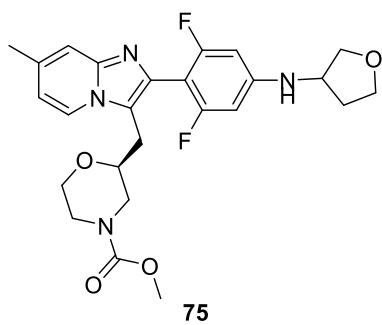
,

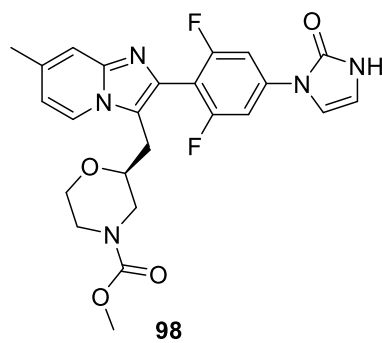
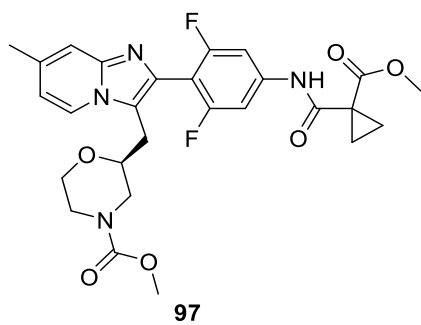
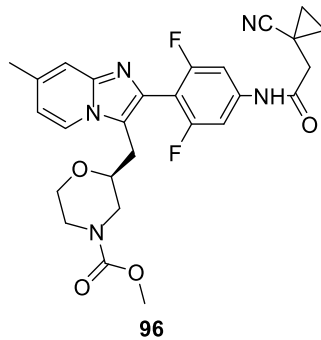
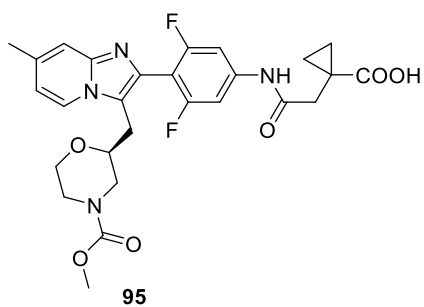
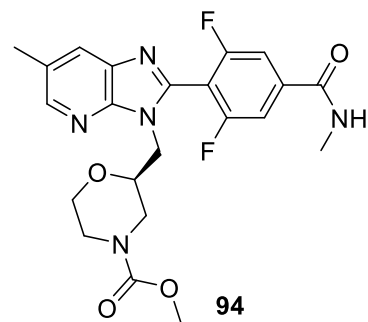
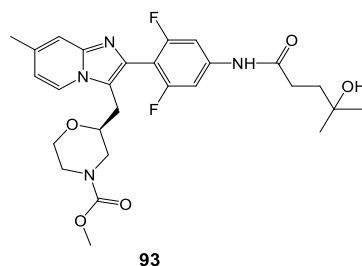
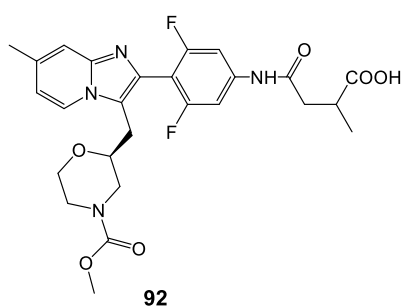
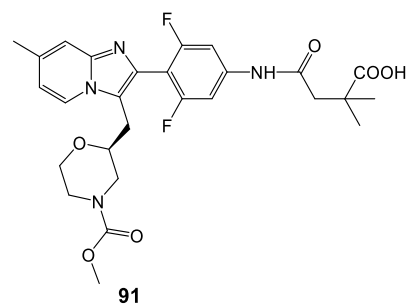
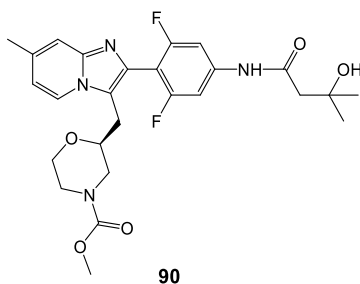
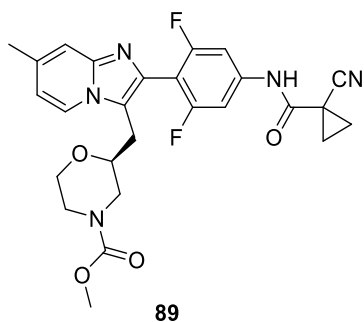
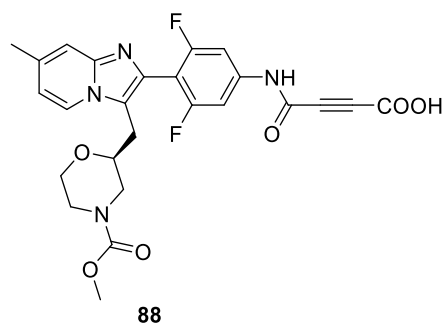
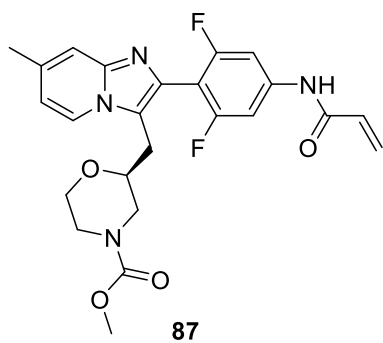


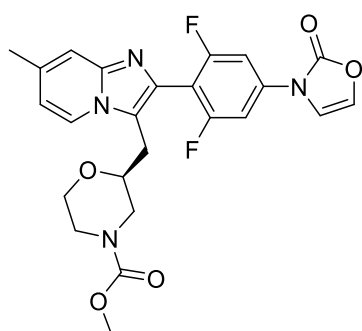




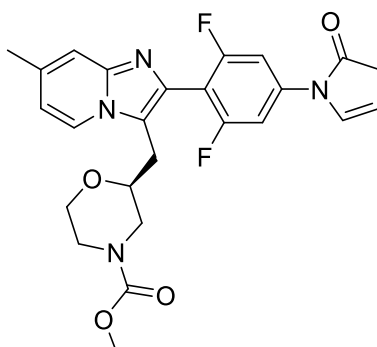




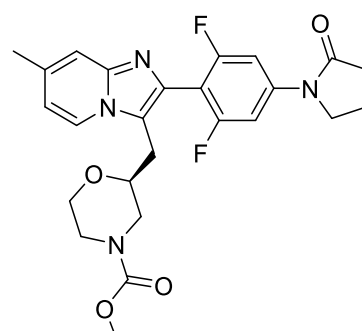




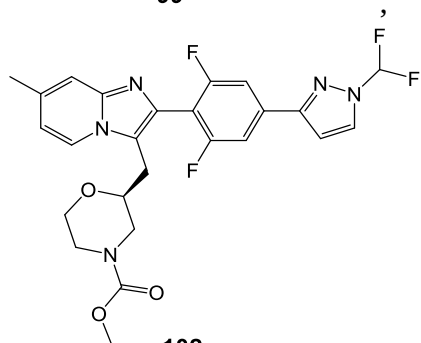
99



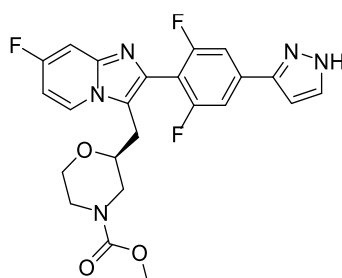
100



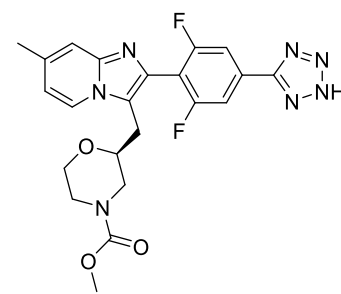
101



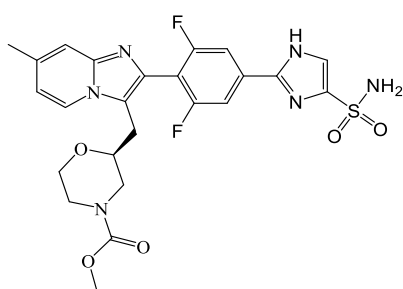
102



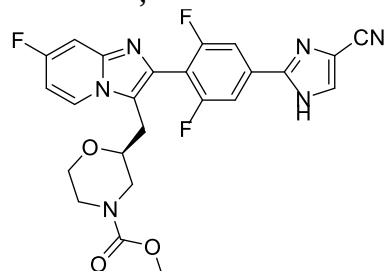
103



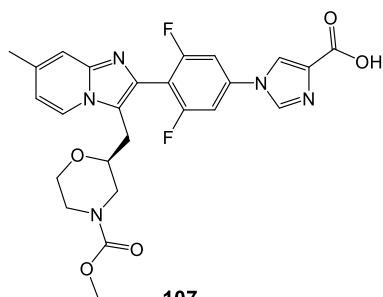
104



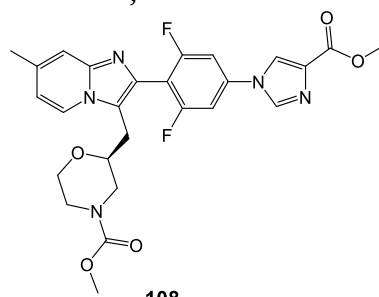
105



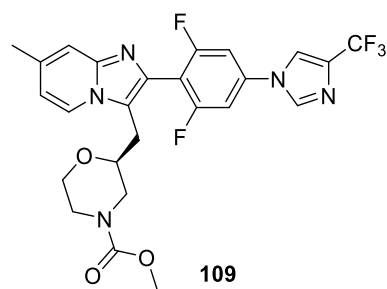
106



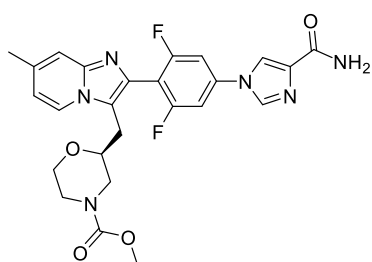
107



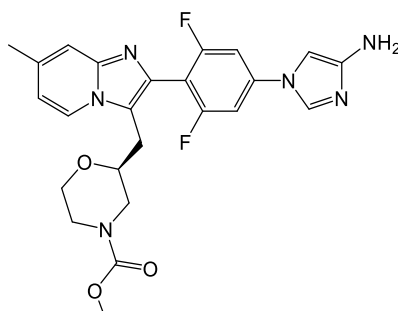
108



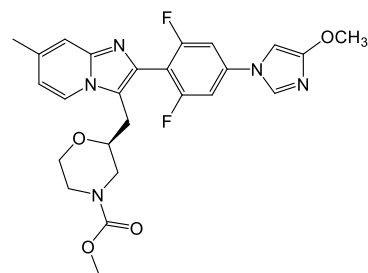
109



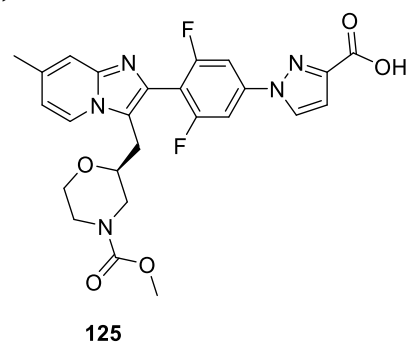
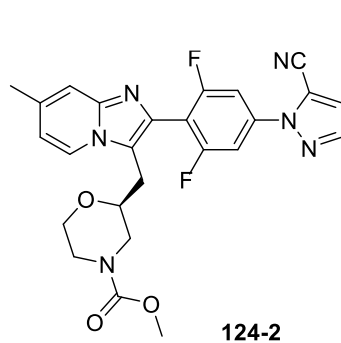
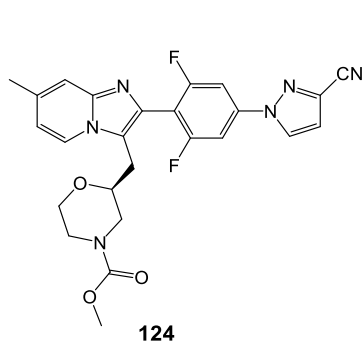
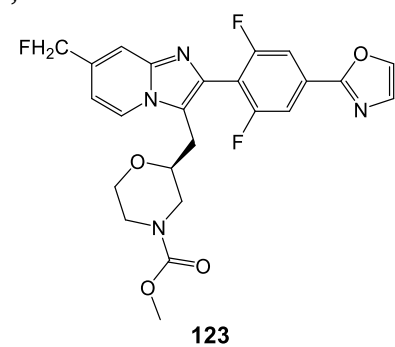
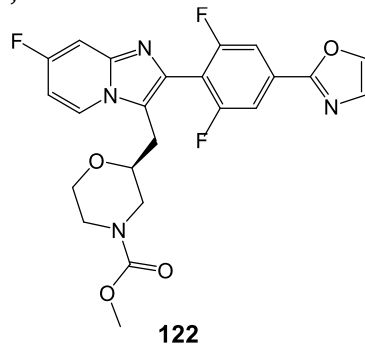
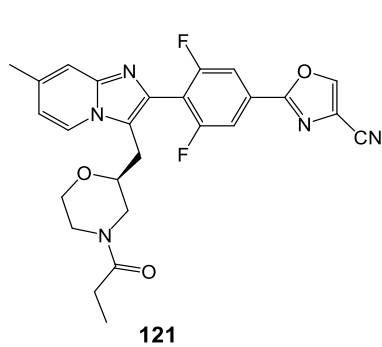
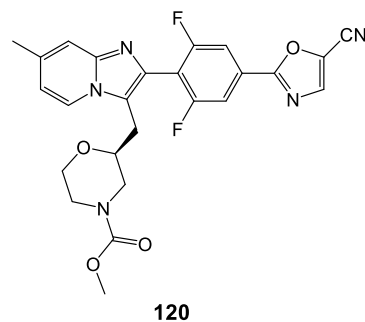
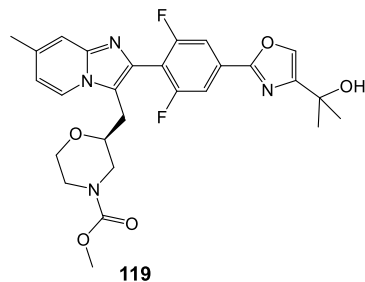
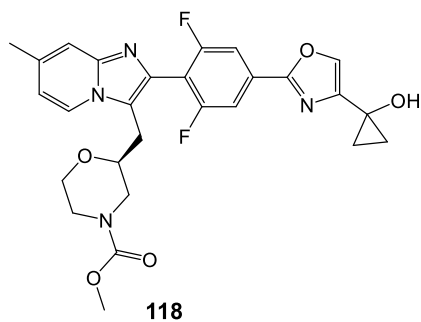
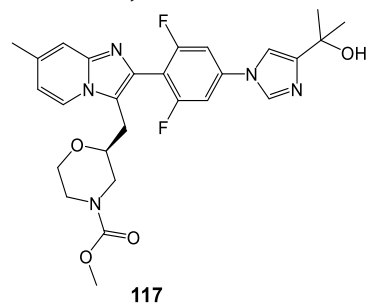
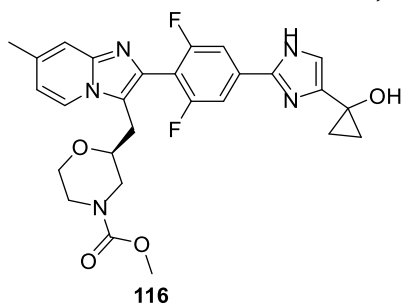
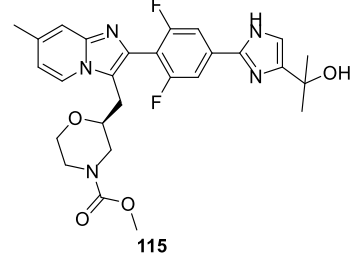
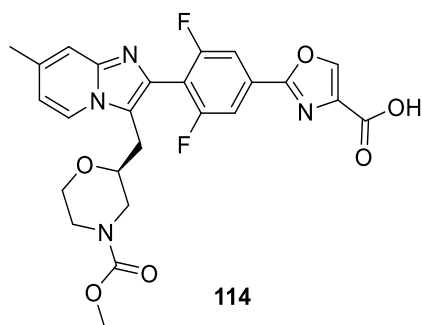
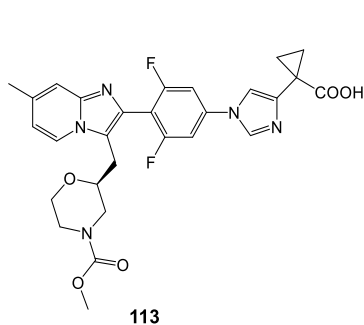
110

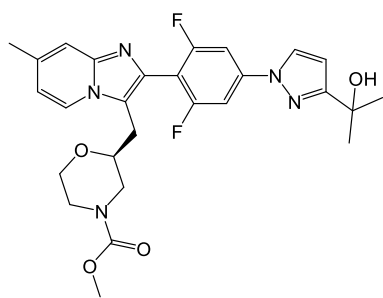


111

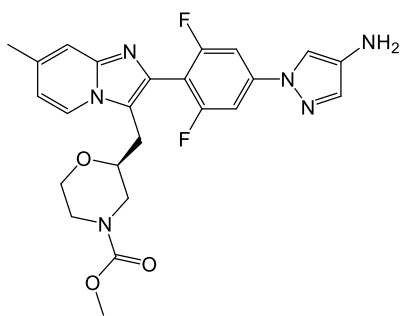


112

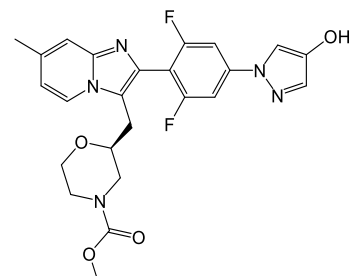




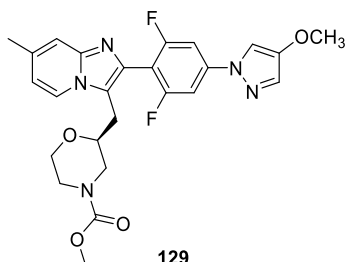
126



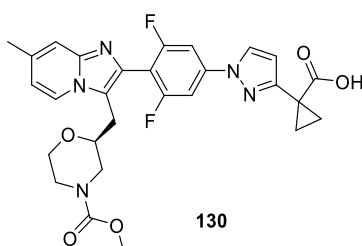
127



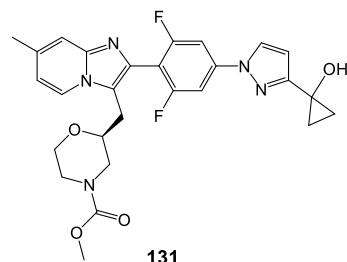
128



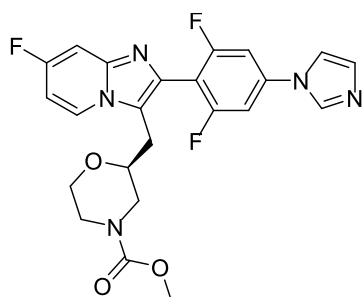
129



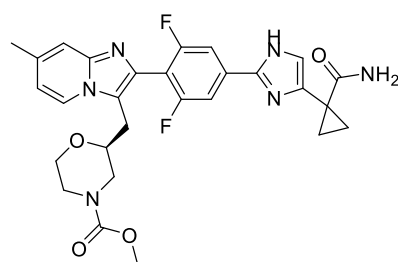
130



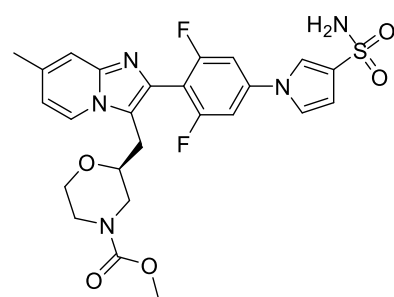
131



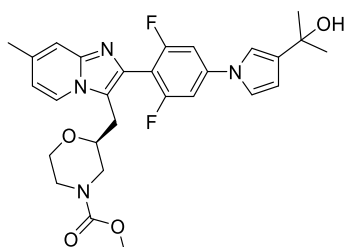
132



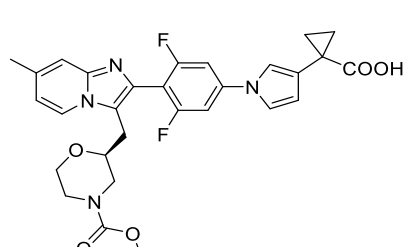
133



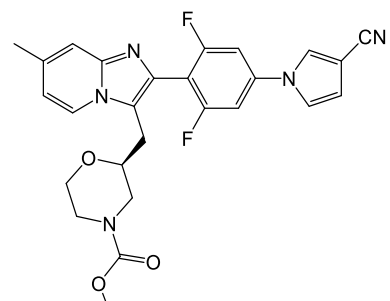
134



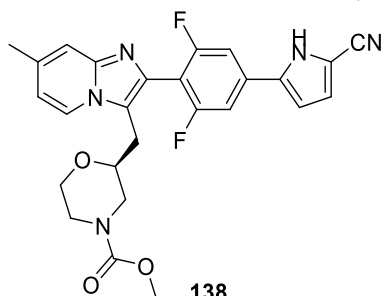
135



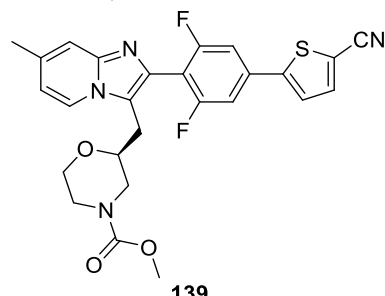
136



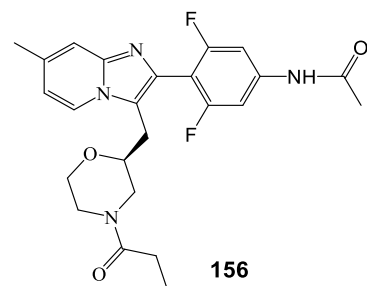
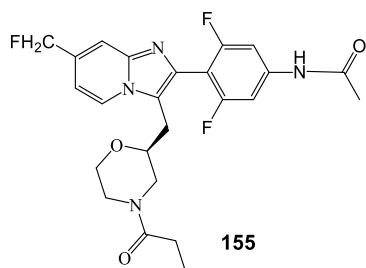
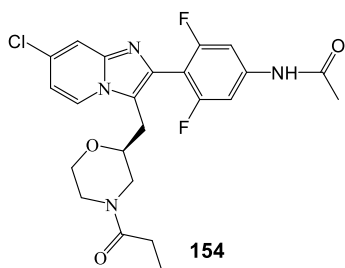
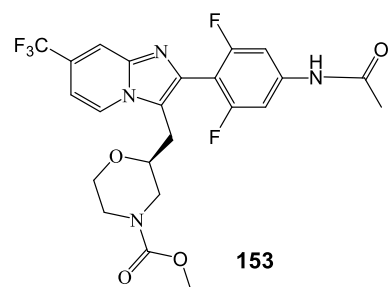
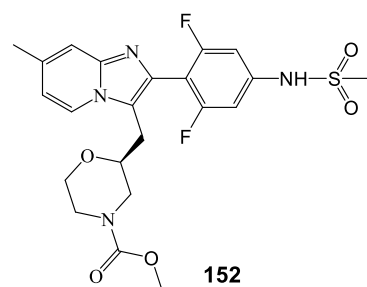
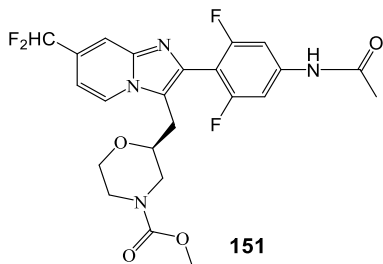
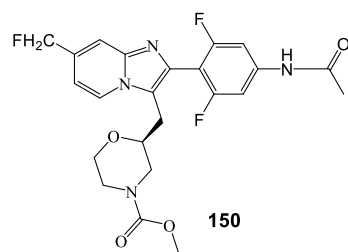
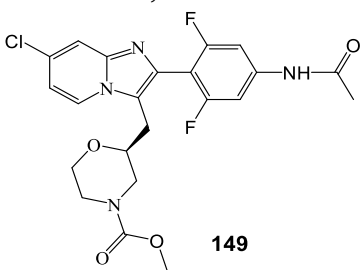
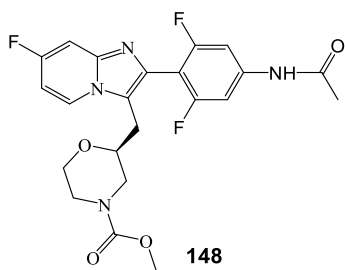
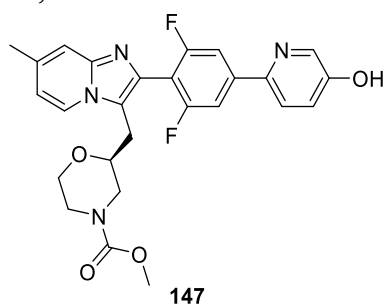
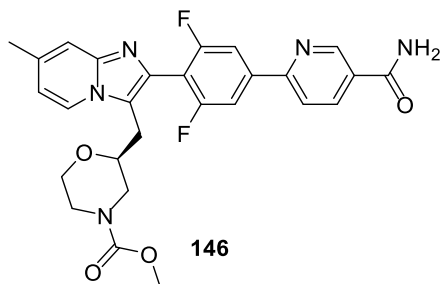
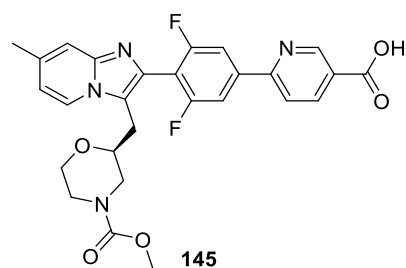
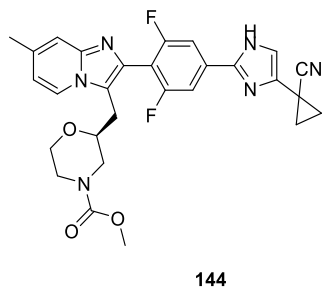
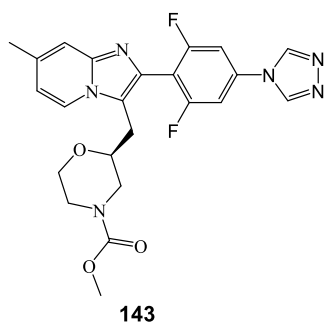
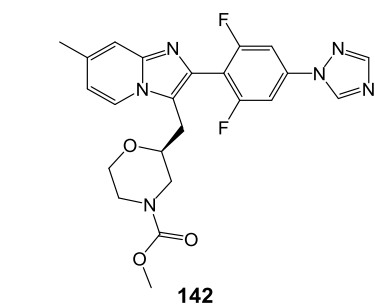
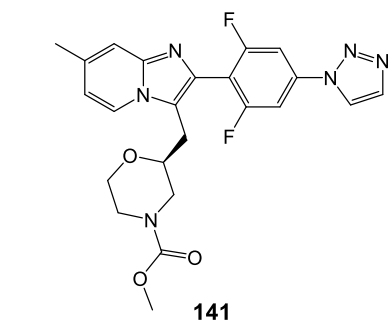
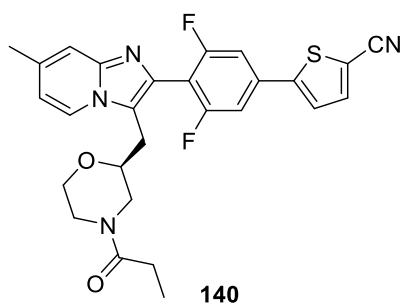
137

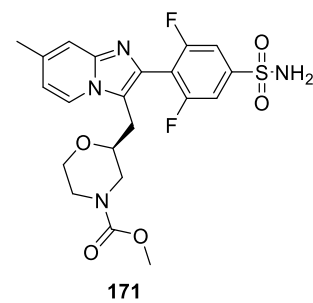
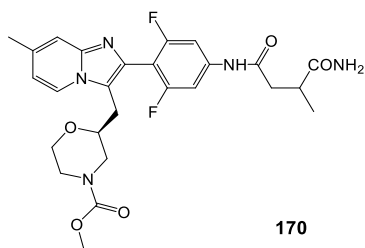
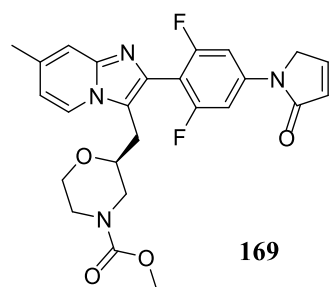
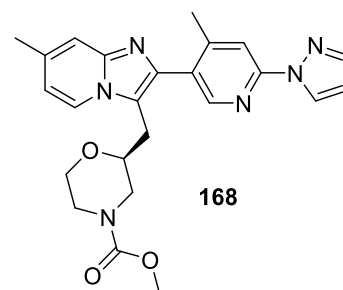
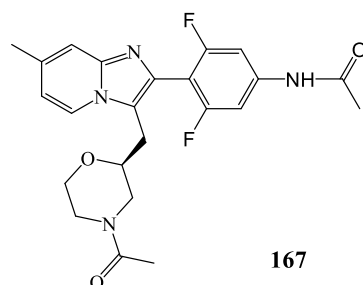
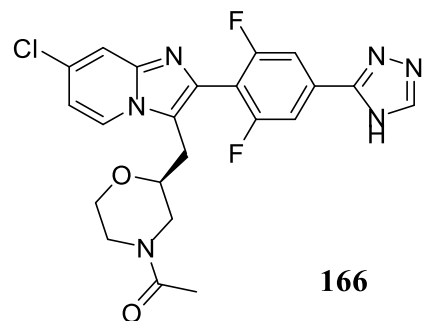
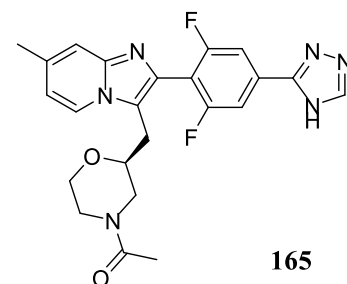
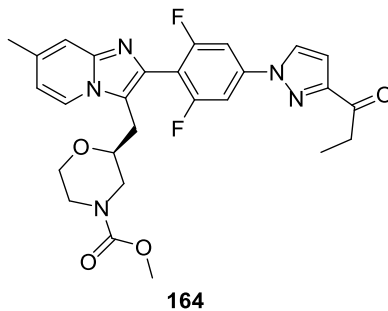
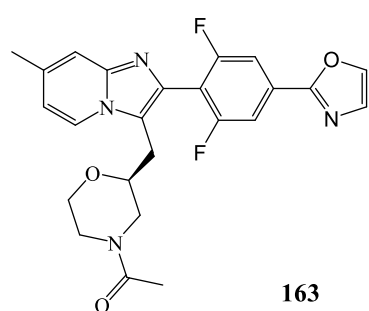
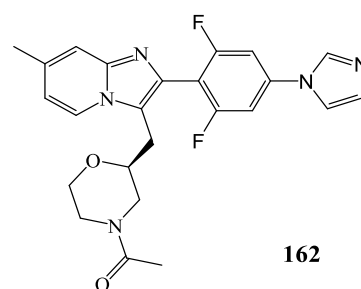
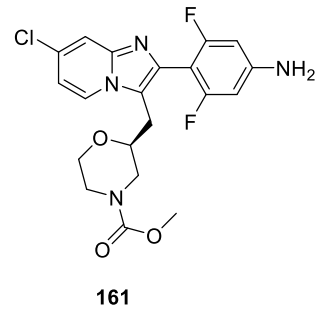
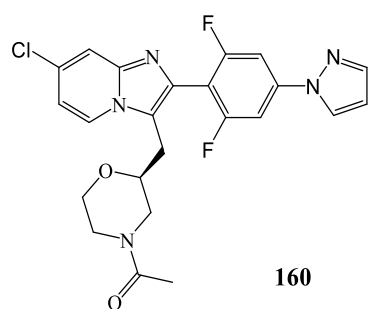
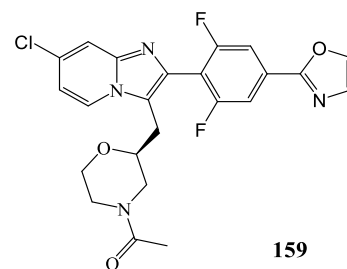
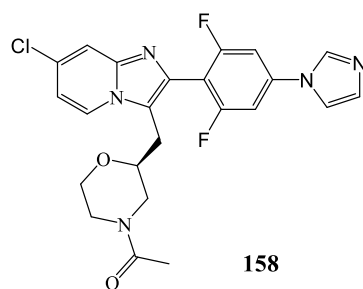
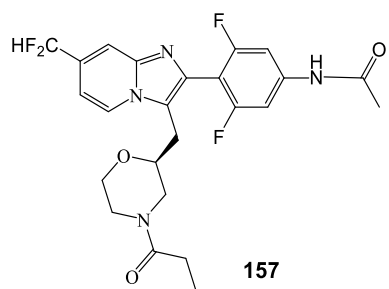


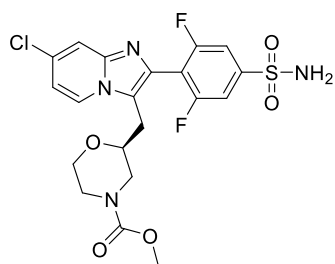
138



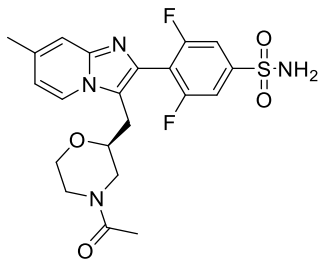
139



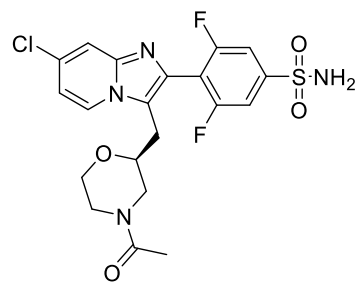




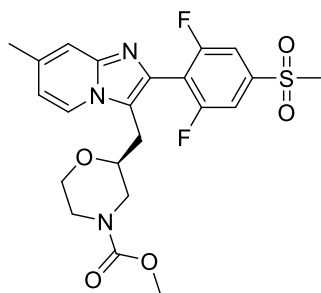
172



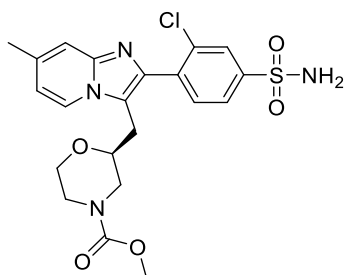
173



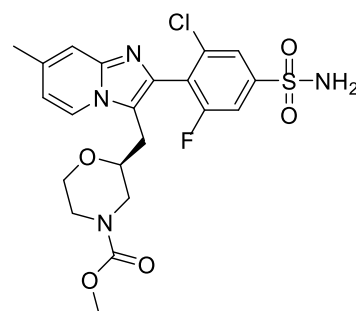
174



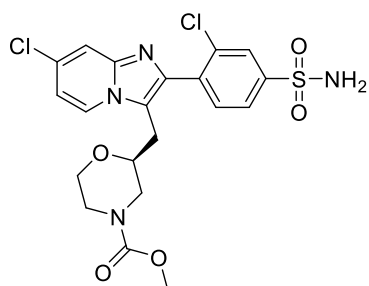
175



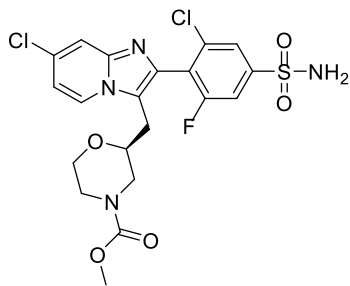
176



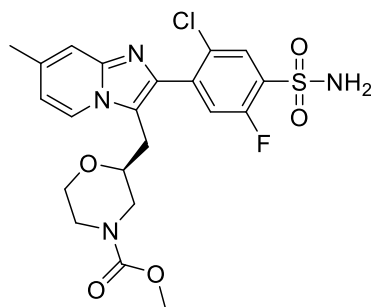
177



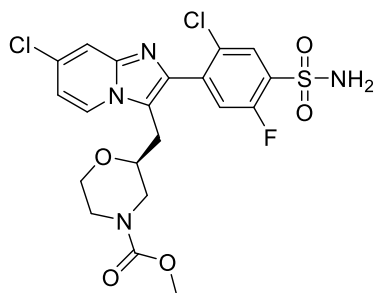
178



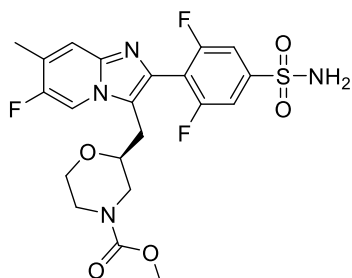
179



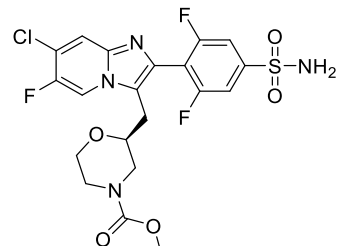
180



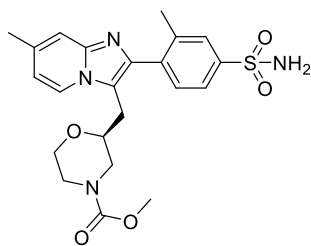
181



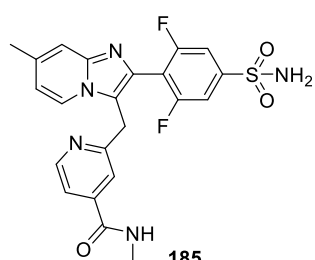
182



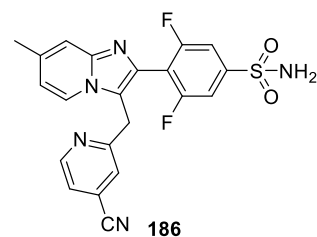
183



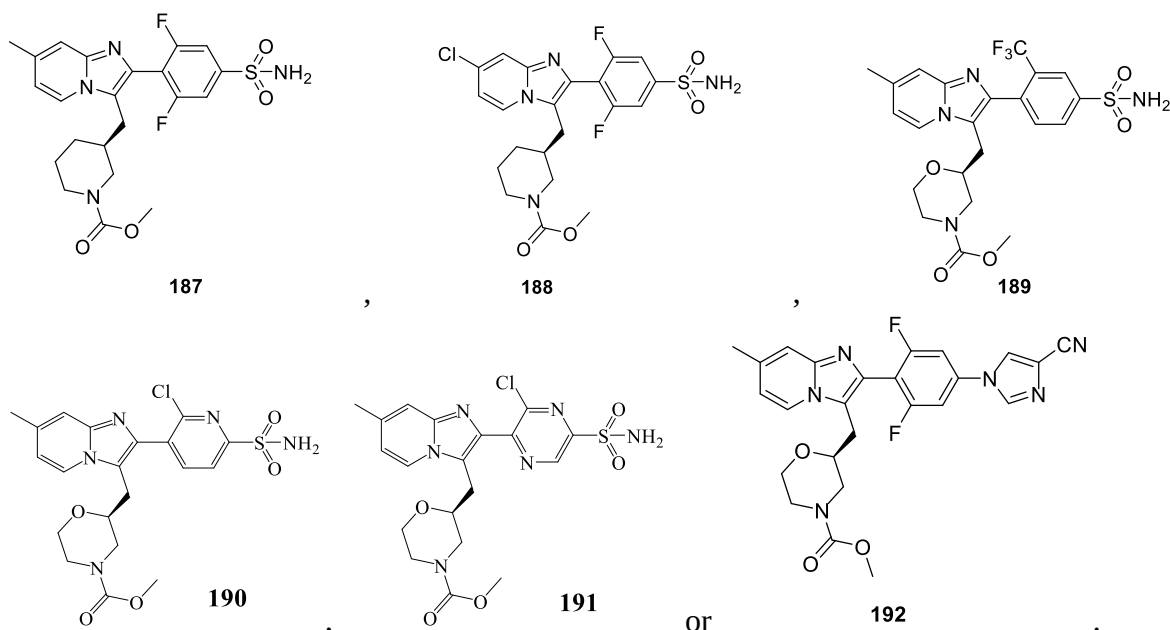
184



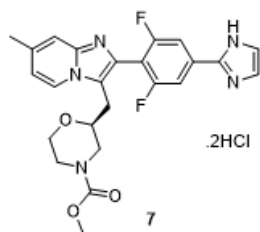
185



186



In a preferred embodiment of the present invention, the pharmaceutically acceptable salt of the heterocyclic compound of formula I is preferably the following compound:



Thus, throughout the present specification, the groups and substituents thereof in the heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof can be selected by those skilled in the art, to provide a stable heterocyclic compound of formula I or a stereoisomer, a geometric isomer, a tautomer, a nitrogen oxide, a hydrate, a solvate, a metabolite, an ester, a pharmaceutically acceptable salt or a prodrug thereof, including but not limited to the compounds described in the examples of the present invention.

The present invention further includes the isotopically labeled heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof disclosed herein, wherein one or more atoms are substituted with one or more atoms having a specified atomic mass or mass number. Examples of isotopes that can be incorporated into the compound disclosed herein include, but are not limited to, isotopes of hydrogen, carbon, nitrogen, oxygen, fluorine, sulfur, and chlorine (e.g., ^2H , ^3H , ^{13}C , ^{14}C , ^{15}N , ^{18}O , ^{17}O , ^{18}F , ^{35}S and ^{36}Cl). The isotopically labeled compound disclosed herein is useful in the assay of tissue

distribution of the compound, the prodrug and the metabolite thereof; and preferred isotopes for such assays include ^3H and ^{14}C . Furthermore, in some cases, substitution with heavier isotopes such as deuterium (^2H or D) can afford increased metabolic stability, which provides therapeutic advantages such as increased *in vivo* half-life or reduced dosage requirements.

The isotopically labeled compound disclosed herein can generally be prepared by substituting an isotopically labeled reagent for a non-isotopically labeled reagent according to the methods described herein.

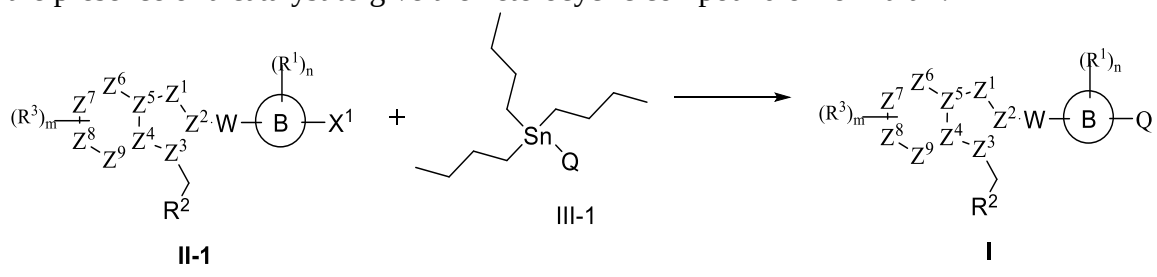
The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof disclosed herein can be synthesized by methods similar to those known in the field of chemistry, with the steps and conditions that can be referred to those of similar reactions in the art, in particular according to the description herein. Starting materials are generally from commercial sources (e.g., Aldrich), or can be readily prepared using methods well known to those skilled in the art (obtained by SciFinder and Reaxys online databases).

The other heterocyclic compounds of formula I can also be obtained by peripherally modifying the heterocyclic compound of formula I disclosed herein using the prepared compounds by conventional methods known in the art.

In general, the compound disclosed herein may be prepared by the methods described herein, wherein the substituents are defined as in formula I, unless otherwise specified. The following schemes and examples serve to further illustrate the context of the present invention.

The present invention provides a preparation method of the heterocyclic compound of formula I, which comprises the following schemes.

Scheme I comprises the following step: performing Stille coupling reaction as shown below on a compound of formula II-1 and an organic tin reagent of formula III-1 in an organic solvent in the presence of a catalyst to give the heterocyclic compound of formula I:



wherein X^1 is Br or Cl.

The methods and conditions for the Stille coupling reaction are conventional in the art for such reactions. For example:

The organic solvent is preferably a cyclic ether solvent (e.g., tetrahydrofuran and/or dioxane) and/or an aromatic hydrocarbon solvent (e.g., toluene). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

The catalyst is preferably Pd(PPh₃)₄. The molar ratio of the catalyst to the compound of formula II-1 may be 1:5-1:20 (e.g., 1:10).

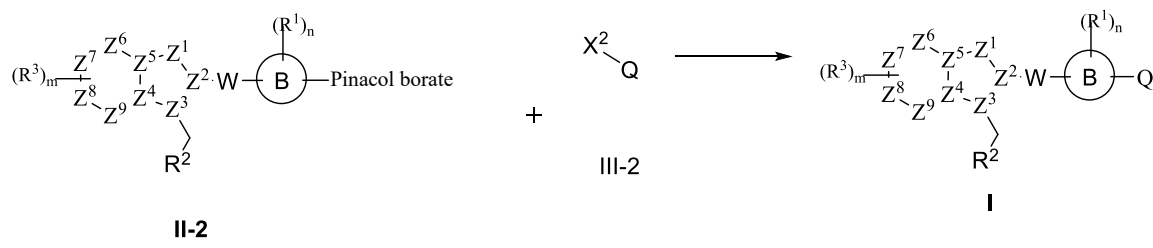
The molar ratio of the compound of formula II-1 to the organic tin reagent of formula III-1 may be a conventional molar ratio in the art, and the molar ratio of the compound of formula II-1 to the organic tin reagent of formula III-1 is preferably 1:3-1:4.

The temperature of the Stille coupling reaction may be 85-110 °C (e.g., 95-100 °C).

The progress of the reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-1 as the end point of the reaction.

In a preferred embodiment of the present invention, in Scheme I, Q is an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl.

Scheme II comprises the following step: performing Suzuki coupling reaction as shown below on a compound of formula II-2 and a compound of formula III-2 in a solvent in the presence of a catalyst and a basic reagent to give the heterocyclic compound of formula I:



wherein X² is Br or Cl.

The methods and conditions for the Suzuki coupling reaction are conventional in the art for such reactions. For example:

The solvent may be water and an organic solvent, and the organic solvent is preferably a cyclic ether solvent (e.g., tetrahydrofuran and/or dioxane). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

The catalyst is preferably Pd(PPh₃)₄. The molar ratio of the catalyst to the compound of formula II-2 may be 1:5-1:20 (e.g., 1:20).

The basic reagent may be Na₂CO₃, and the molar ratio of the basic reagent to the compound of formula II-2 may be 1:1-5:1 (e.g., 3:1).

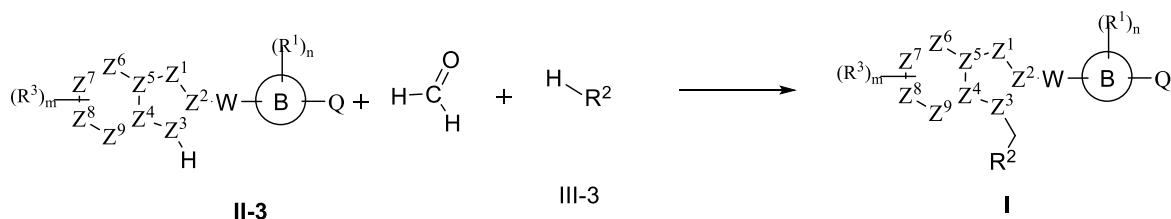
The molar ratio of the compound of formula II-2 to the compound of formula III-2 may be a conventional molar ratio in the art, and the molar ratio of the compound of formula II-2 to the compound of formula III-2 is preferably 1:1-1:1.2.

The temperature of the Suzuki coupling reaction may be 90-100 °C.

The progress of the reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-2 as the end point of the reaction.

In a preferred embodiment of the present invention, in Scheme II, Q is an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl.

Scheme III comprises the following step: performing the reaction as shown below on a compound of formula II-3, a compound of formula III-3 and formaldehyde in an organic solvent to give the heterocyclic compound of formula I:



The methods and conditions for the reaction are conventional in the art for such reactions. For example:

The organic solvent is preferably a carboxylic acid solvent (e.g., acetic acid). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

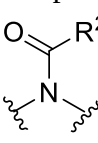
The formaldehyde may be aqueous formaldehyde solution, and preferably the aqueous formaldehyde solution at 37 wt.%; and the molar ratio of the formaldehyde to the compound of formula II-2 may be 1:1.

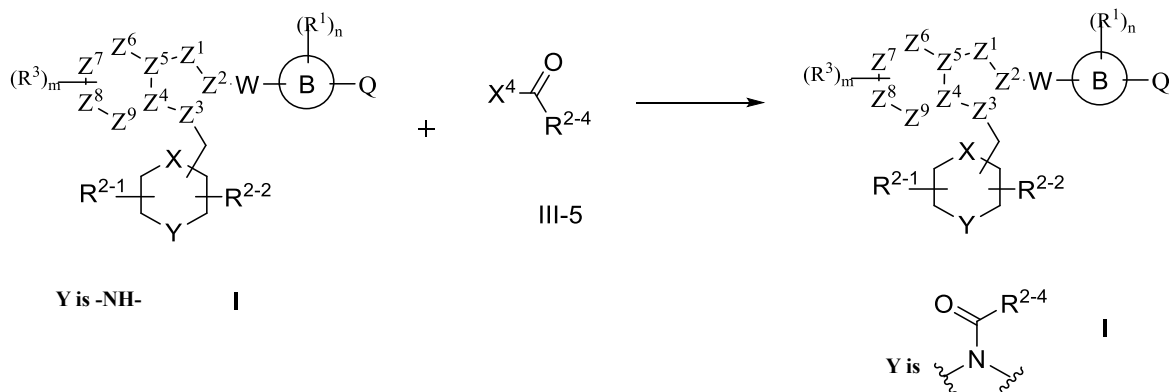
The molar ratio of the compound of formula II-2 to the compound of formula III-2 may be a conventional molar ratio in the art, and the molar ratio of the compound of formula II-2 to the compound of formula III-2 is preferably 1.2:1-1:1.2.

The temperature of the reaction may be 40-60 °C.

The progress of the reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-3 as the end point of the reaction.

formula III-5 in an organic solvent in the presence of a basic reagent to give the heterocyclic

compound of formula I (wherein Y is );



wherein X⁴ is halogen (e.g., Br or Cl).

5 The methods and conditions for the amidation reaction are conventional in the art for such reactions. For example:

The organic solvent is preferably a halogenated alkane solvent (e.g., dichloromethane). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

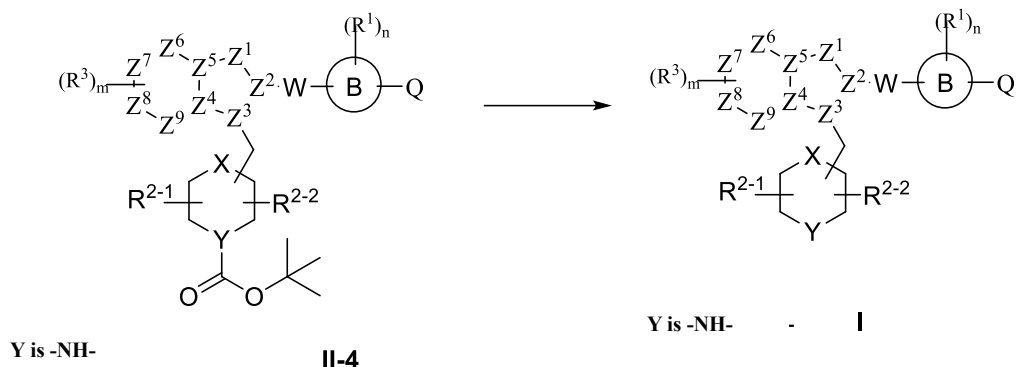
10 The basic reagent may be *N,N*-diisopropylethylamine, and the molar ratio of the basic reagent to the compound of formula II-4 may be 2:1-4:1 (e.g., 3:1).

The molar ratio of the compound of formula II-4 to the compound of formula III-5 may be a conventional molar ratio in the art, and the molar ratio of the compound of formula II-4 to the compound of formula III-5 is preferably 1.2:1-1:1.2.

The temperature of the amidation reaction may be -10-20 °C.

15 The progress of the amidation reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-4 as the end point of the reaction.

20 Scheme VI comprises the following step: performing deprotection reaction as shown below on a compound of formula II-4 and a compound of formula III-5 in an organic solvent in the presence of an acid to give the heterocyclic compound of formula I:



wherein Y is -NH-.

The methods and conditions for the deprotection reaction are conventional in the art for such reactions. For example:

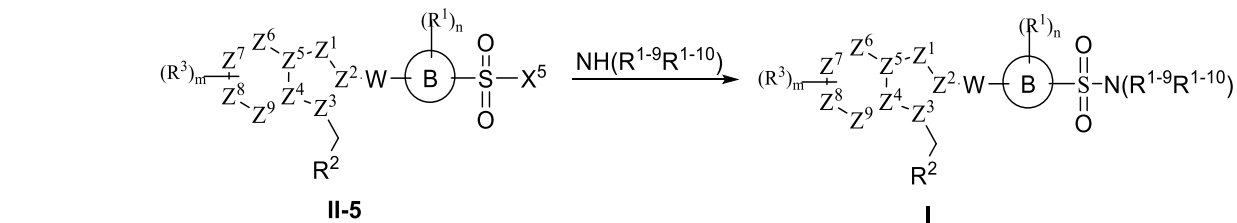
- 5 The organic solvent is preferably an alcohol solvent (e.g., ethanol). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

The acid may be hydrochloric acid, and the molar ratio of the acid to the compound of formula II-4 may be 1:1-4:1 (e.g., 1.5:1).

The temperature of the deprotection reaction may be -10-20 °C.

- 10 The progress of the amidation reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-4 as the end point of the reaction.

Scheme VII comprises the following step: performing substitution reaction as shown below on a compound of formula II-5 and NH(R¹⁻⁹R¹⁻¹⁰) in an organic solvent to give the heterocyclic compound of formula I (wherein Q is -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰)):



wherein X⁵ is Br or Cl.

The methods and conditions for the reaction are conventional in the art for such reactions. For example:

- 20 The organic solvent is preferably a nitrile solvent (e.g., acetonitrile). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

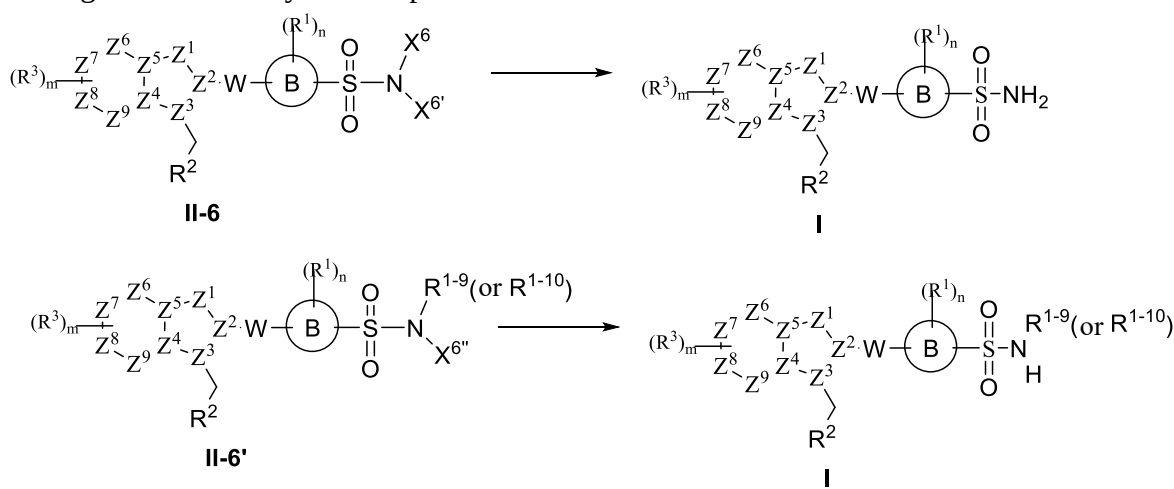
The molar ratio of the compound of formula II-5 to NH(R¹⁻⁹R¹⁻¹⁰) may be 1:1-1:5.

The NH(R¹⁻⁹R¹⁻¹⁰) may be in a form conventional in the art, for example in the form of aqueous ammonia when it is NH₃.

- 25 The temperature of the reaction may be -10-20 °C (e.g., 0 °C).

The progress of the reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-3 as the end point of the reaction.

Scheme VIII comprises the following step: performing deprotection reaction as shown below on a compound of formula II-6 or formula II-6' in an organic solvent in the presence of an acid to give the heterocyclic compound of formula I:



wherein X⁶, X^{6'} and X^{6''} are N protecting groups.

In a preferred embodiment, when X⁶, X^{6'} and X^{6''} are N protecting groups, the N protecting groups may be 4-methoxybenzyl (PMB).

The methods and conditions for the reaction are conventional in the art for such reactions. For example:

The organic solvent is preferably a halogenated alkane solvent (e.g., dichloromethane). The amount of the organic solvent is not particularly limited as long as the reaction proceeds.

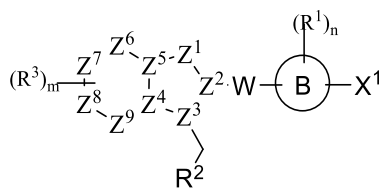
The acid may be trifluoromethanesulfonic acid, and the volume-to-mass ratio of the acid to the compound of formula II-6 or formula II-6' may be 1-5 mL/g (e.g., 3-4 mL/g).

The temperature of the reaction may be 0-100 °C (e.g., 30-50 °C).

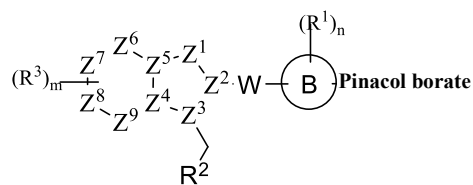
The progress of the reaction may be monitored by detection methods conventional in the art (e.g., TLC, GC, HNMR, or HPLC), and preferably by detecting the disappearance of the compound of formula II-6 or formula II-6' as the end point of the reaction.

In the present invention, after the reaction is completed, conventional work-up methods can be adopted for a treatment. In the present invention, after work-up, if the heterocyclic compound crude product of formula I is obtained, conventional means such as preparative HPLC, preparative TLC or recrystallization can be adopted for separation and purification.

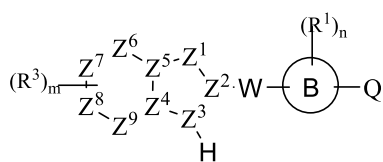
The present invention also provides a compound of formula II-1, II-2, II-3, II-4, II-5, II-6 or II-6':



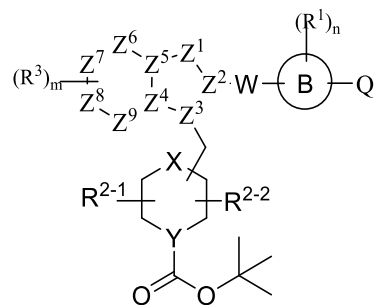
II-1



II-2

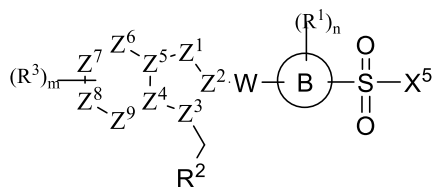


II-3

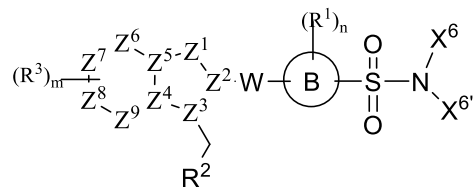


Y is -NH-

II-4

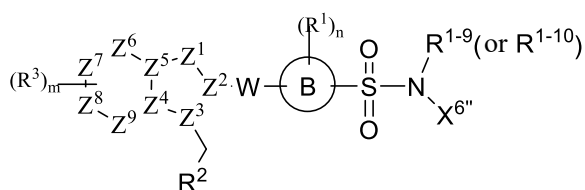


II-5



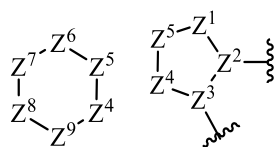
II-6

or



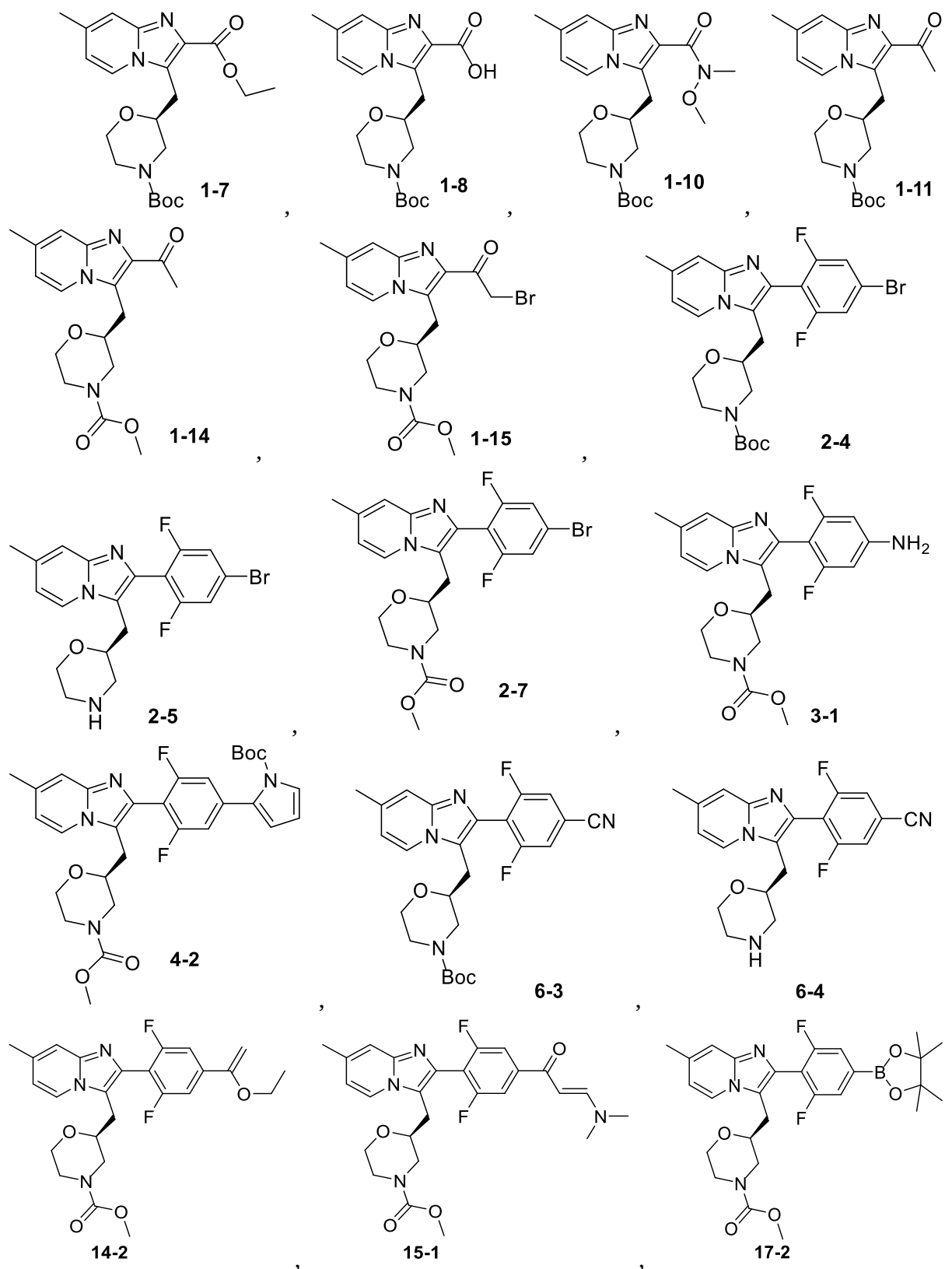
II-6'

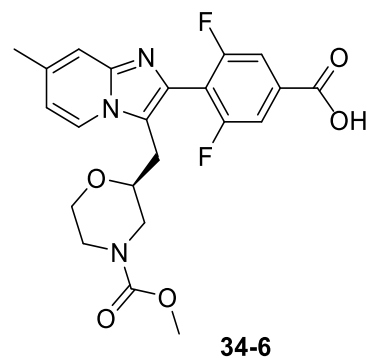
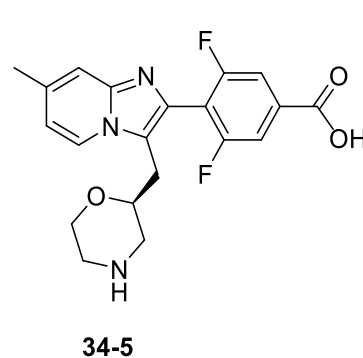
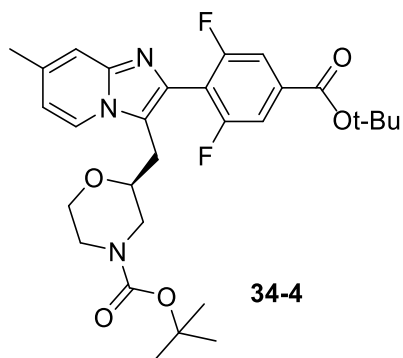
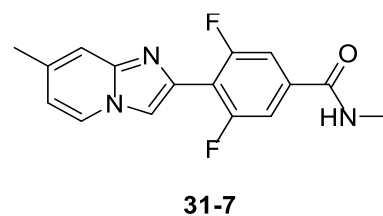
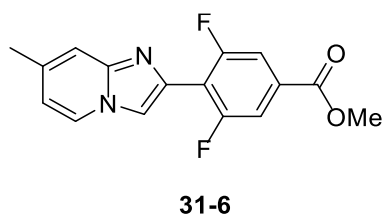
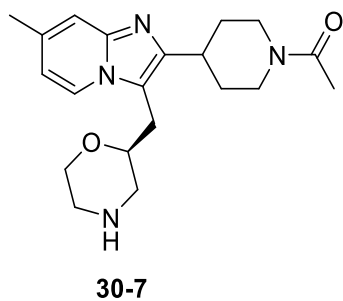
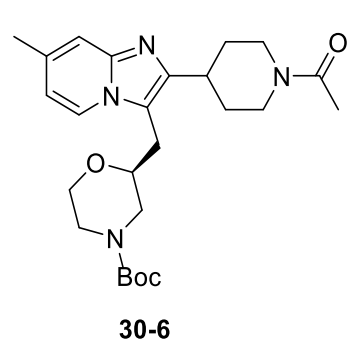
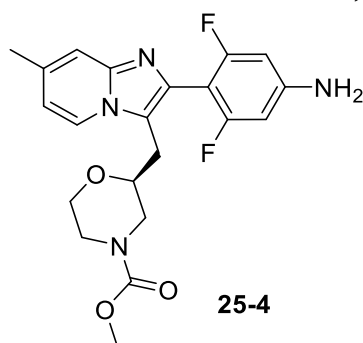
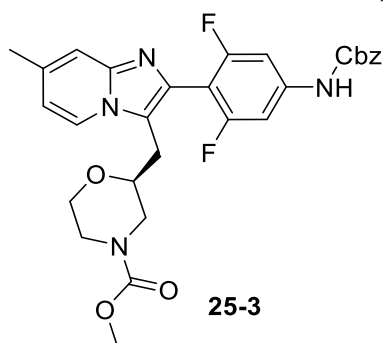
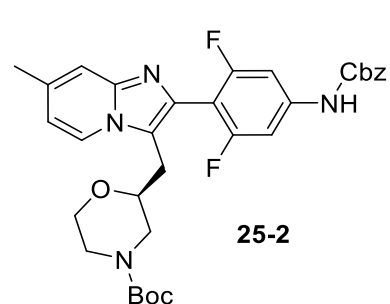
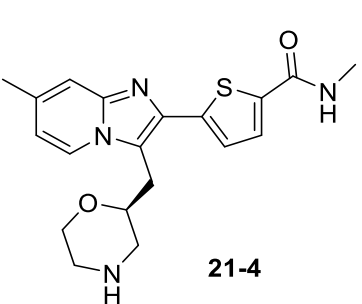
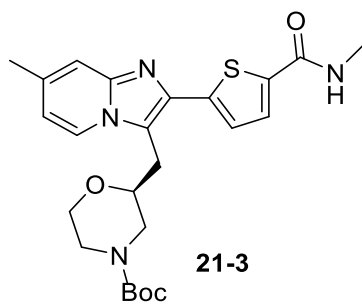
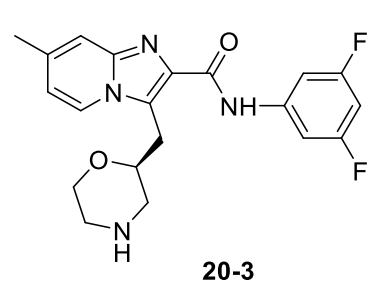
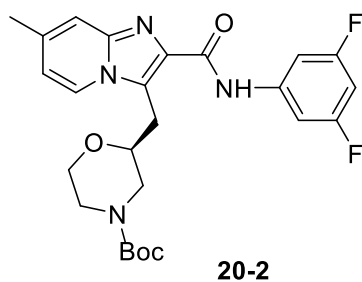
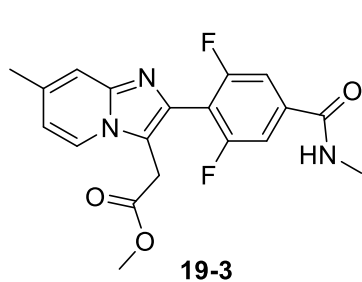
;

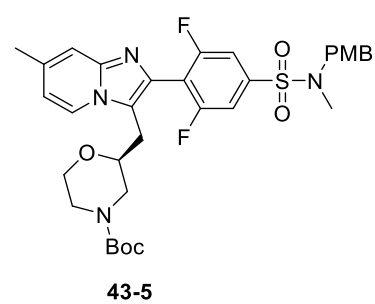
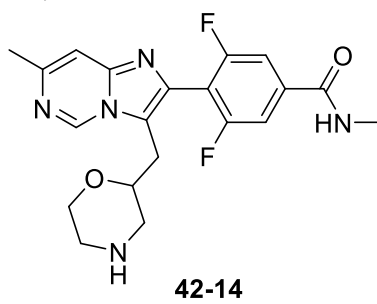
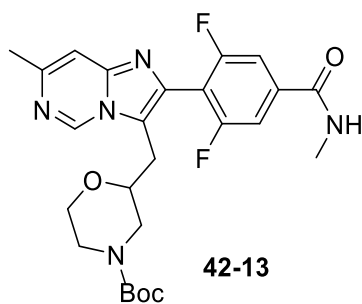
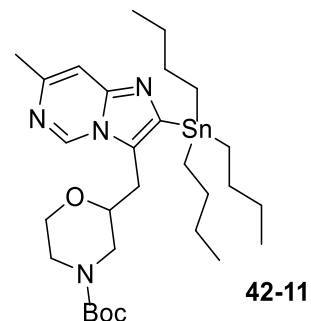
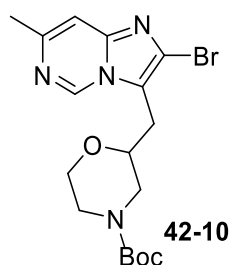
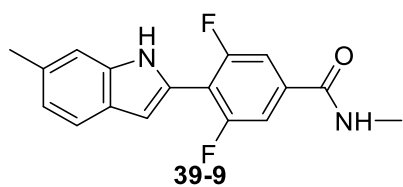
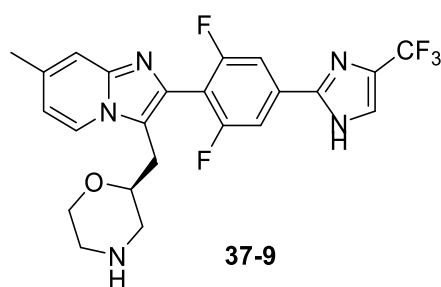
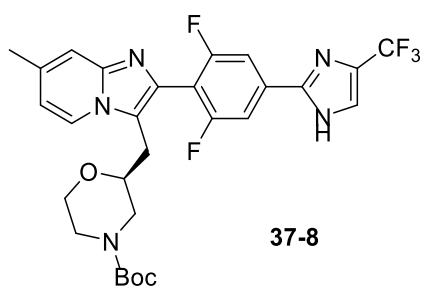
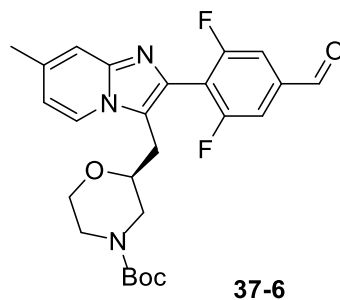
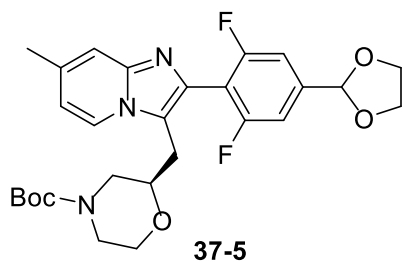
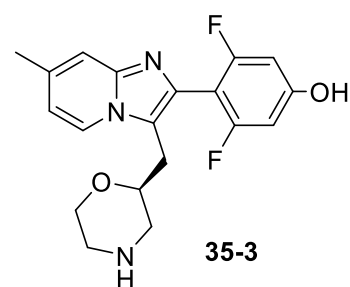
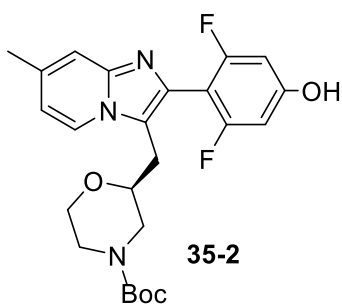
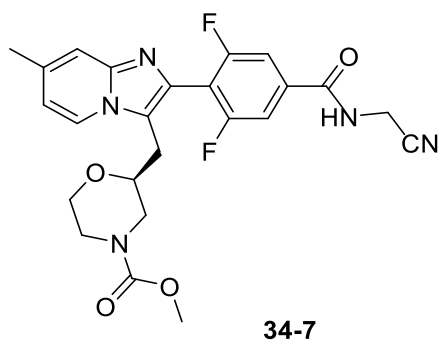


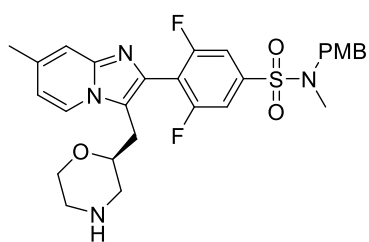
5 wherein the , , R^3 , R^2 , R^1 , W , X , Y , R^{2-1} , R^{2-2} , n , m , X^5 , X^6 , $X^{6'}$, $X^{6''}$ and X^1 have the definitions as described above.

The present invention also provides compounds as shown below:

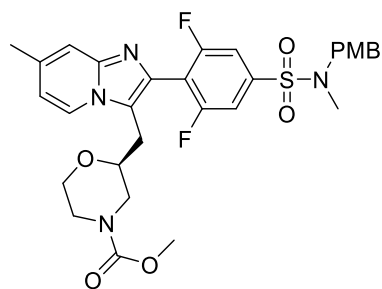




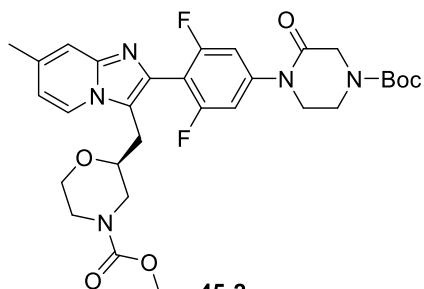




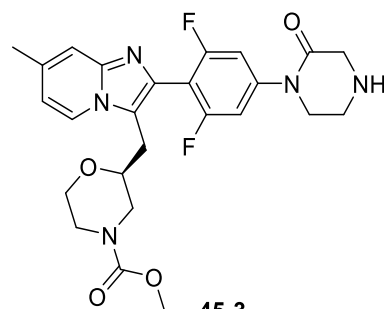
43-6



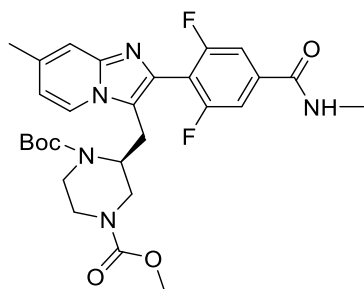
43-7



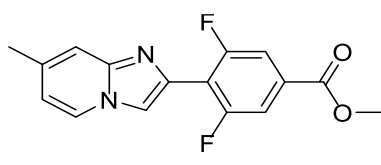
45-2



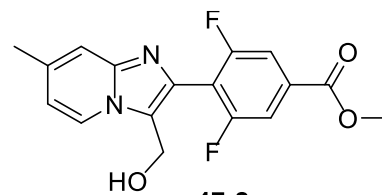
45-3



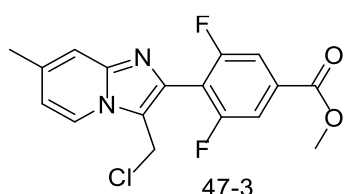
46-6



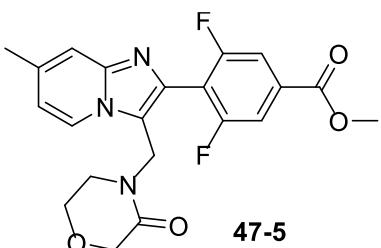
47-1



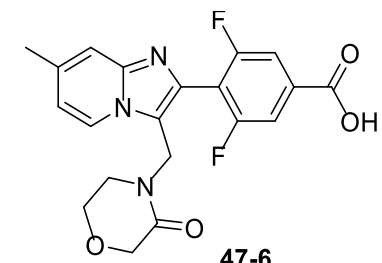
47-2



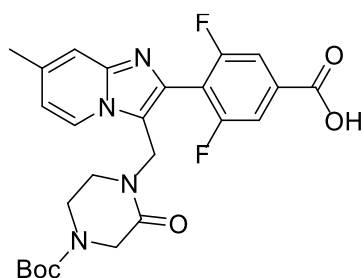
47-3



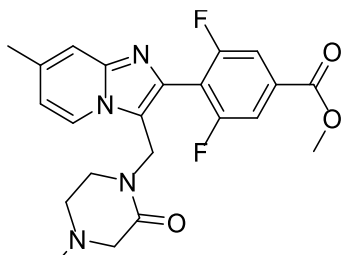
47-5



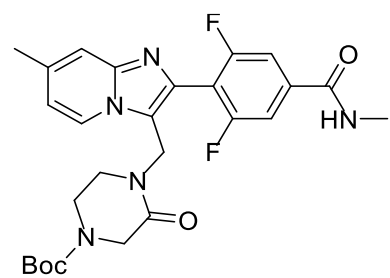
47-6



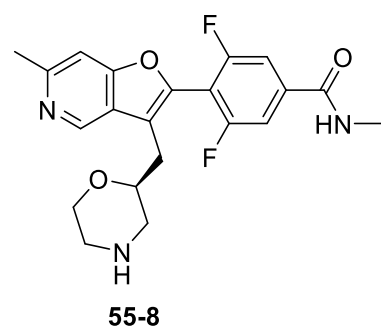
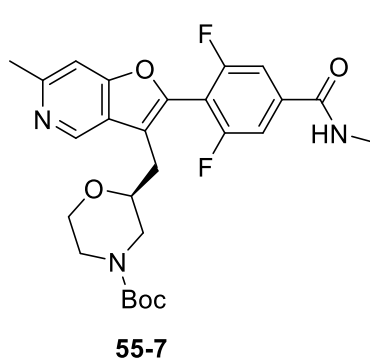
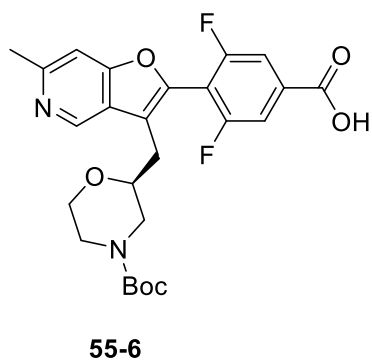
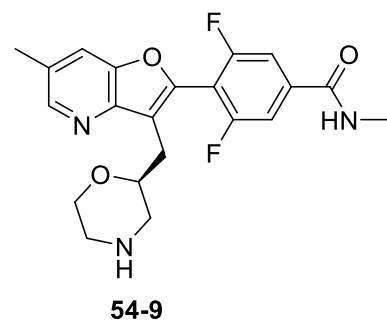
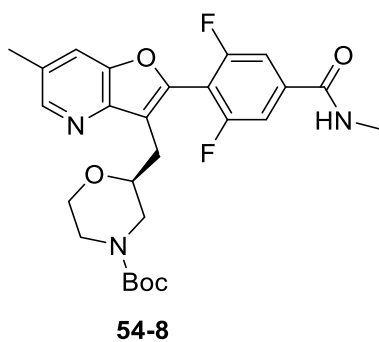
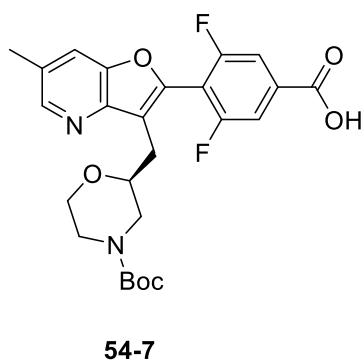
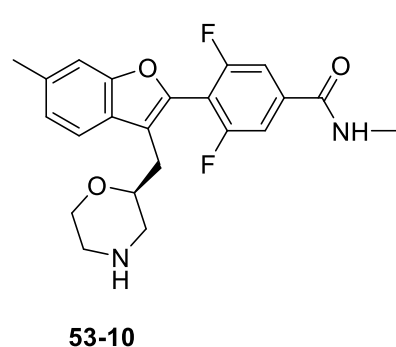
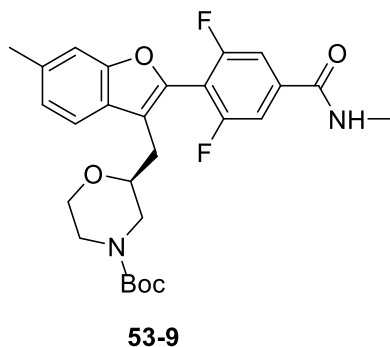
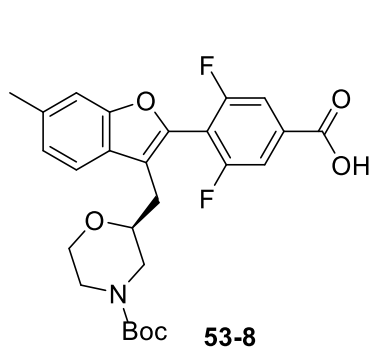
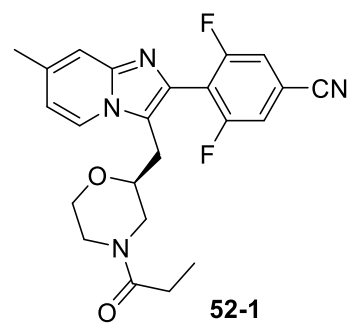
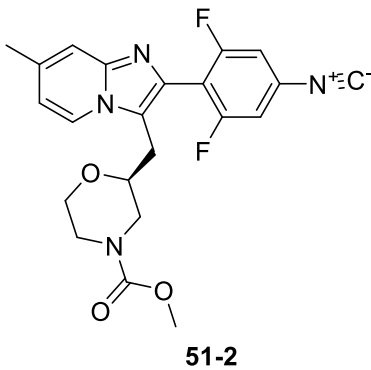
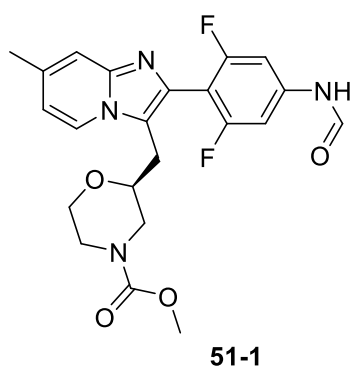
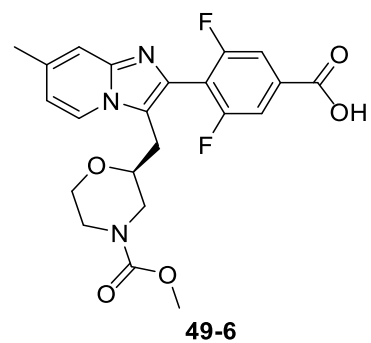
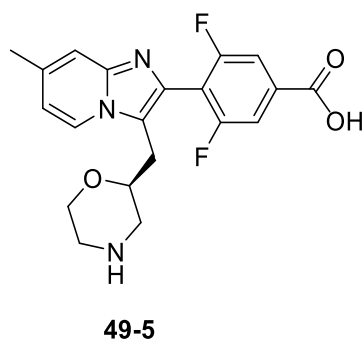
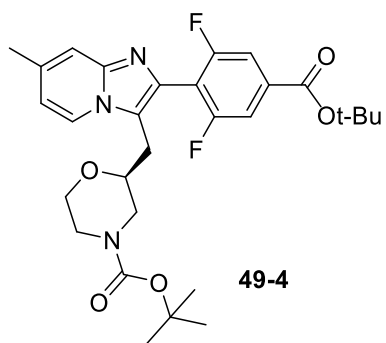
48-1

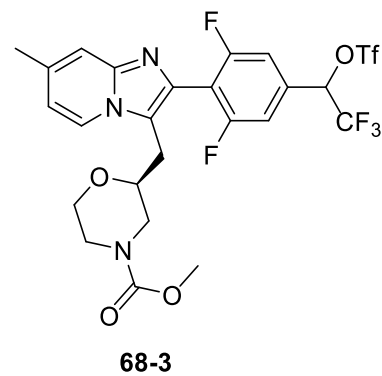
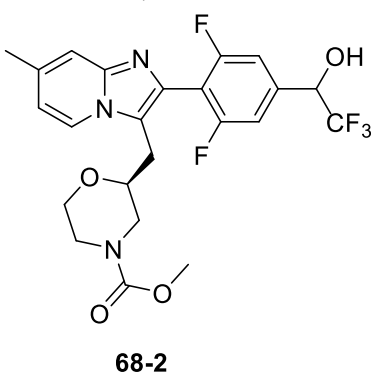
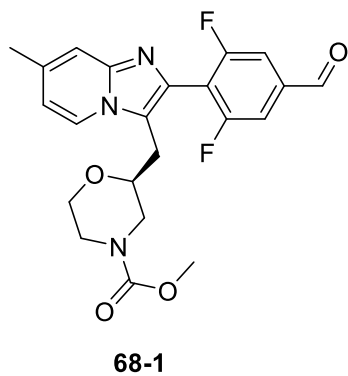
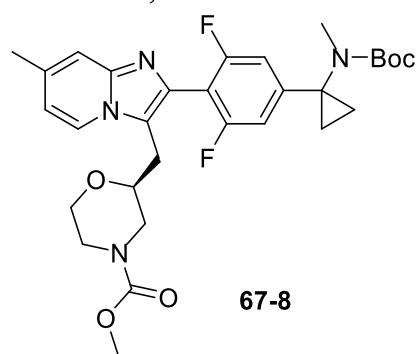
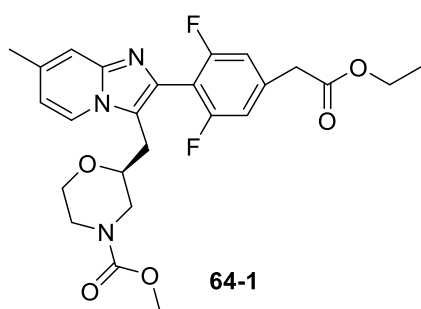
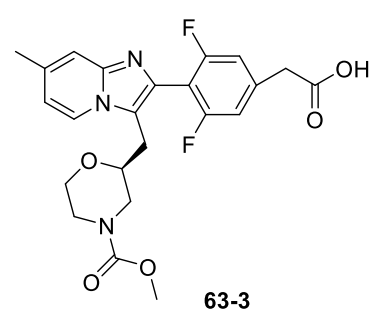
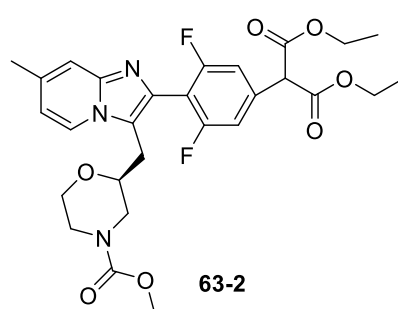
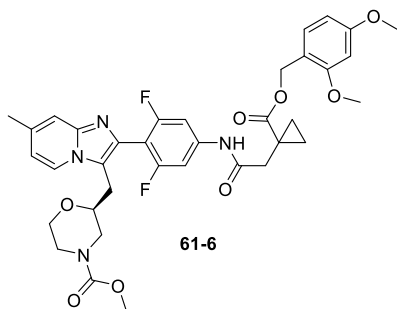
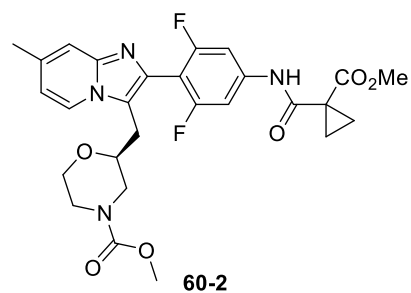
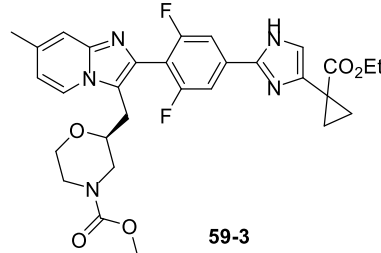
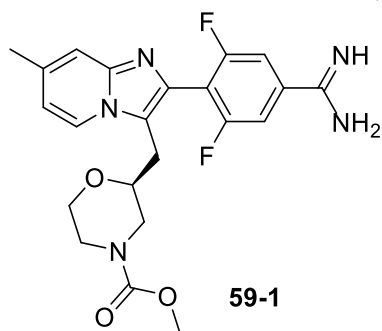
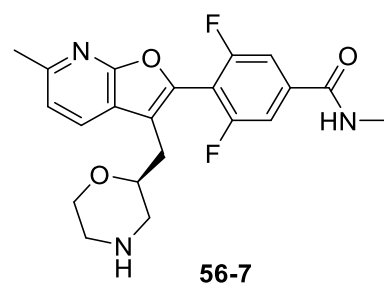
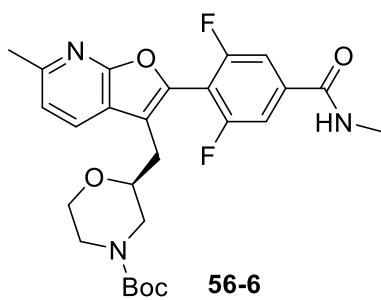
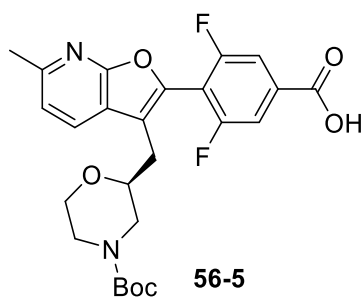


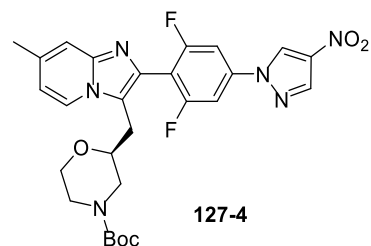
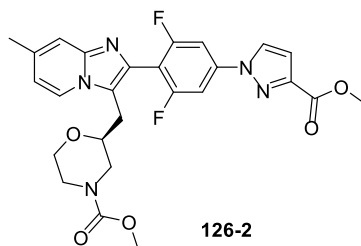
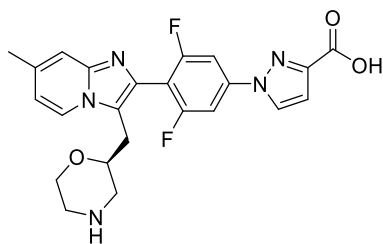
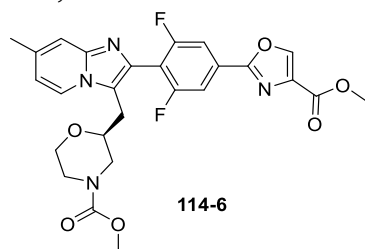
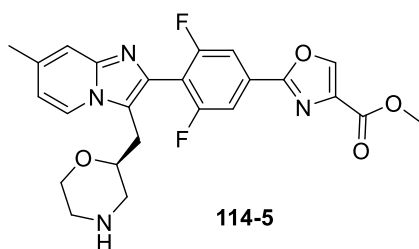
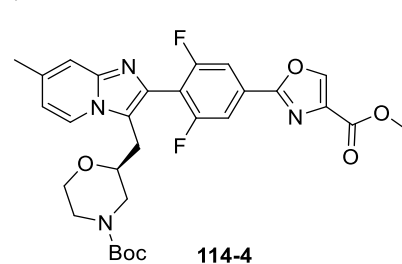
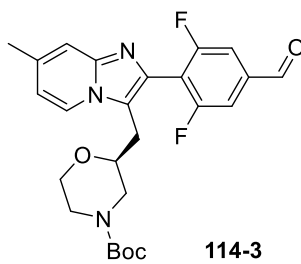
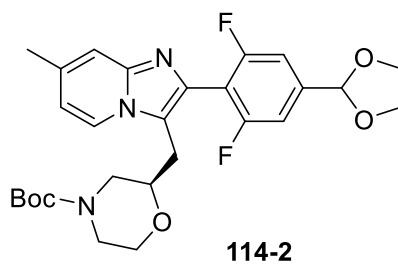
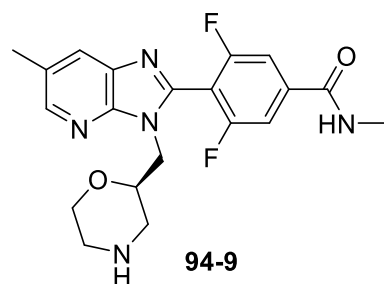
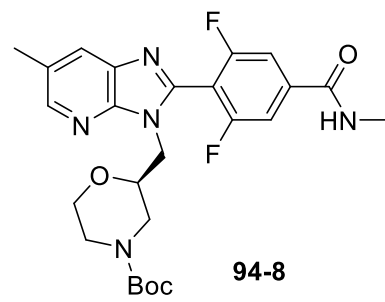
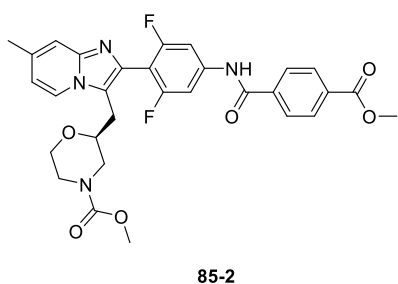
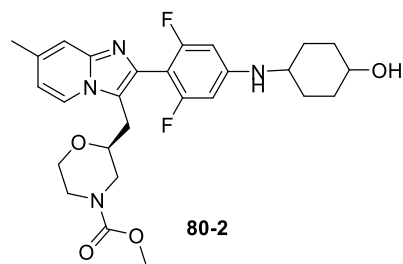
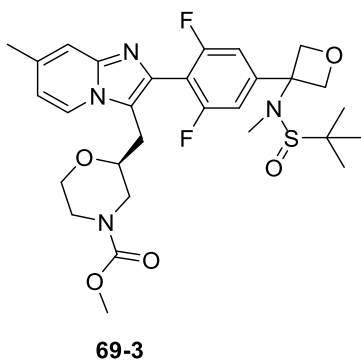
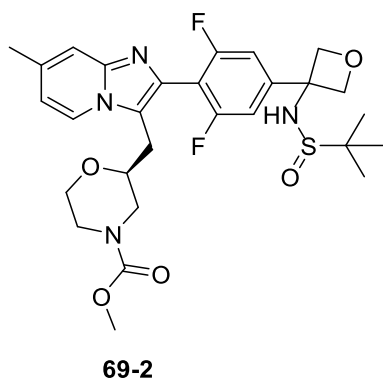
48-2

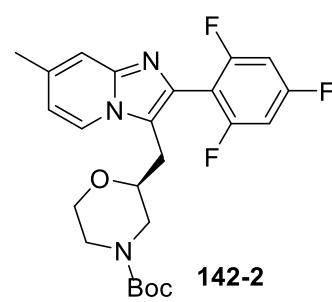
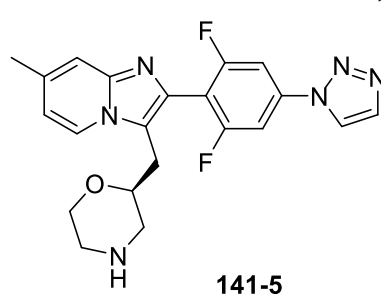
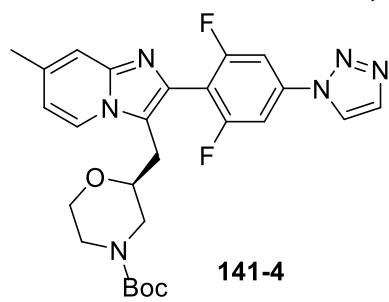
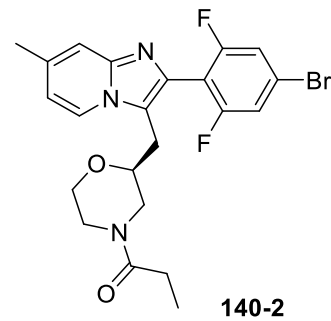
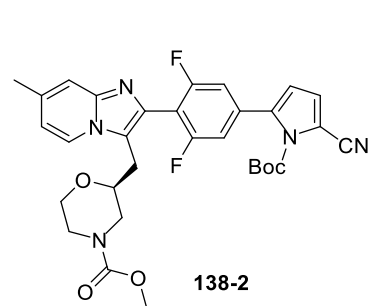
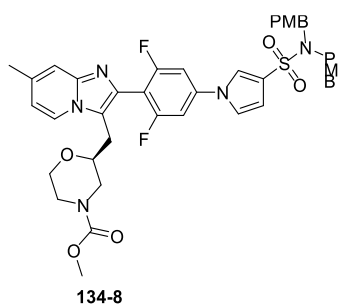
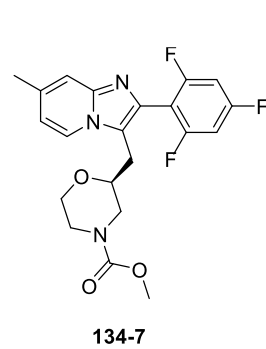
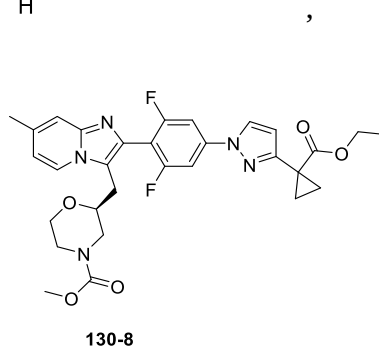
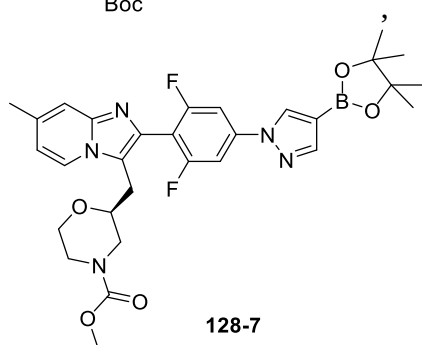
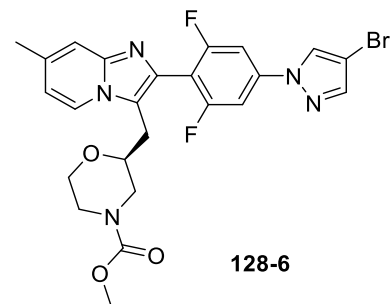
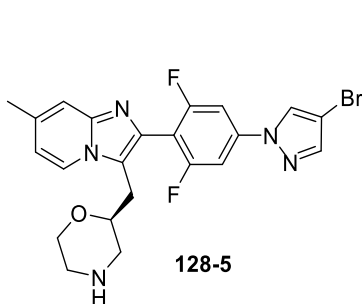
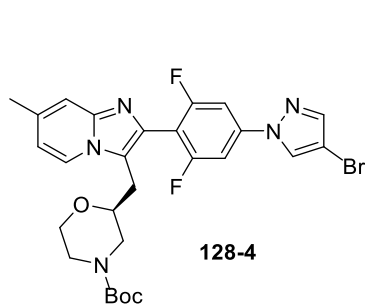
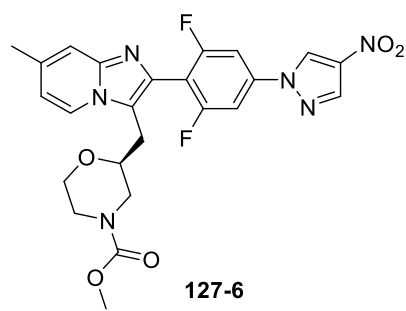
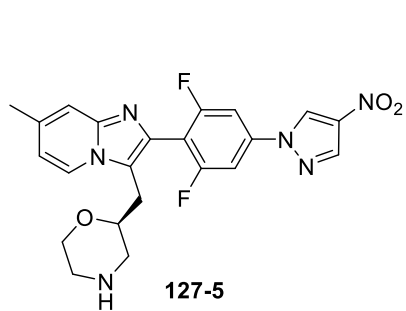


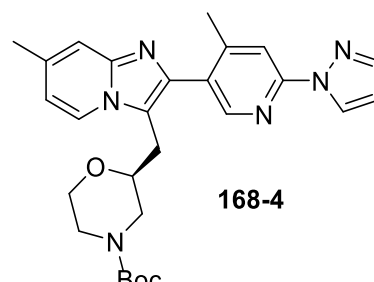
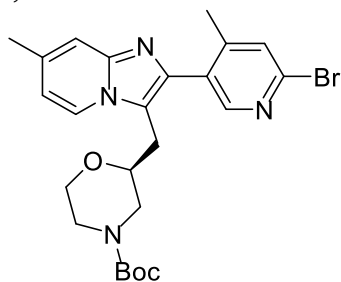
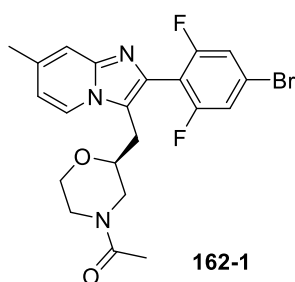
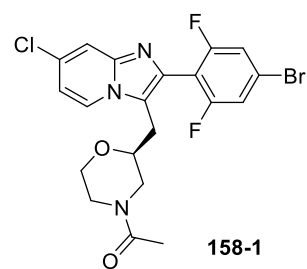
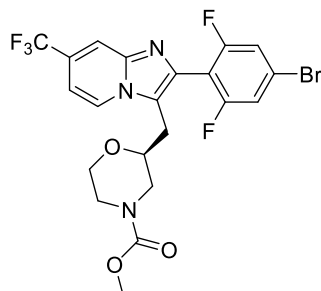
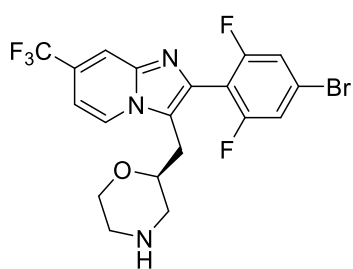
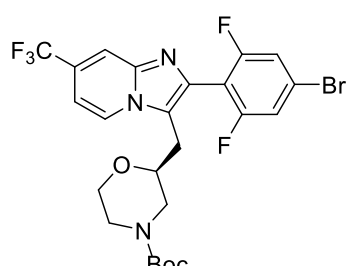
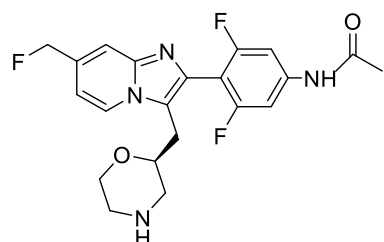
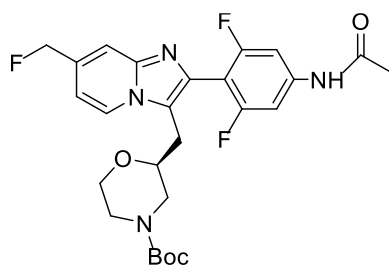
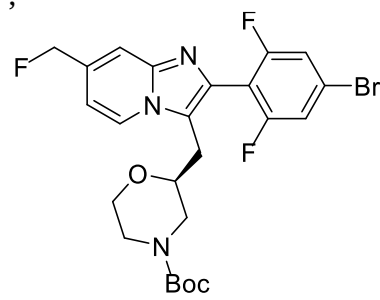
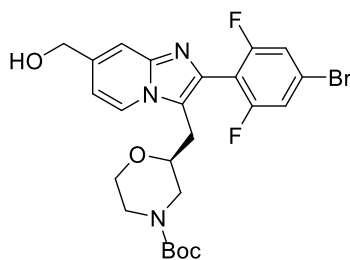
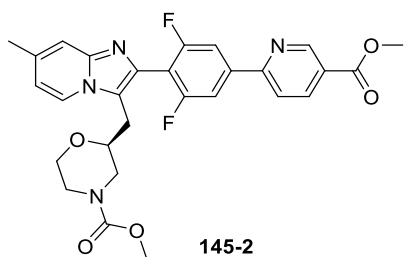
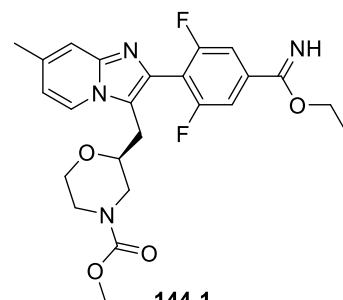
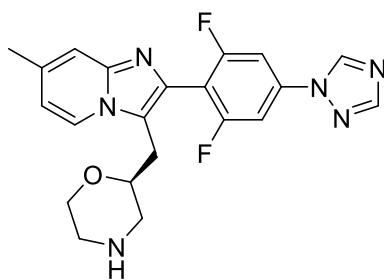
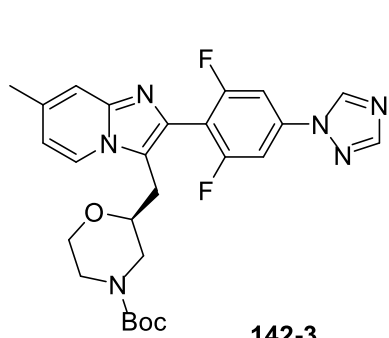
48-3

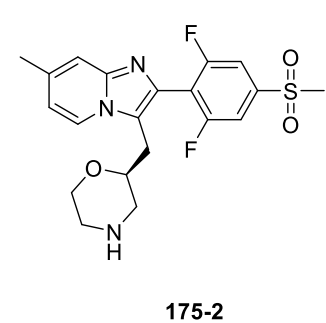
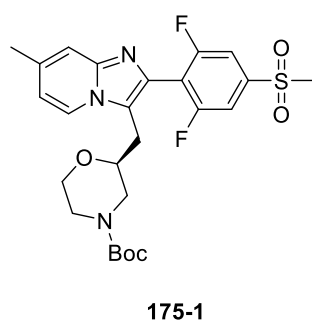
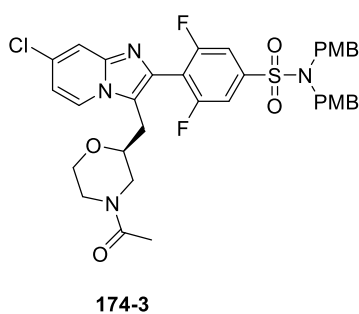
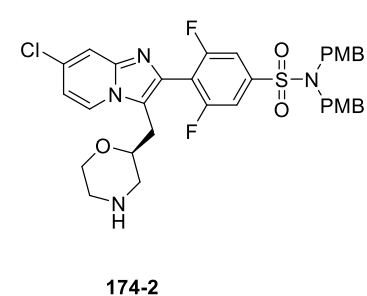
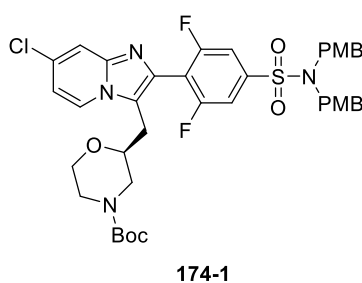
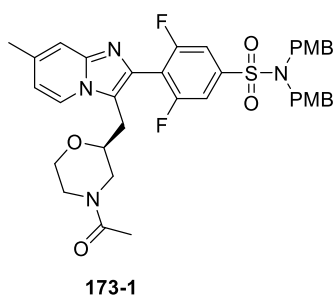
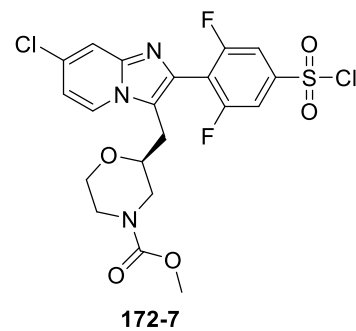
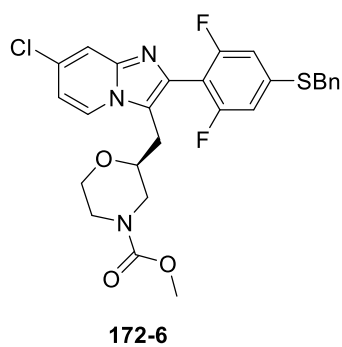
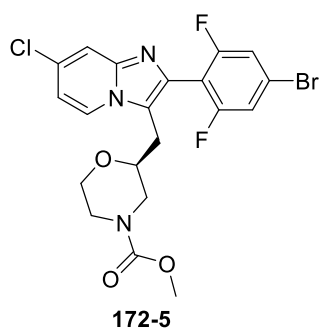
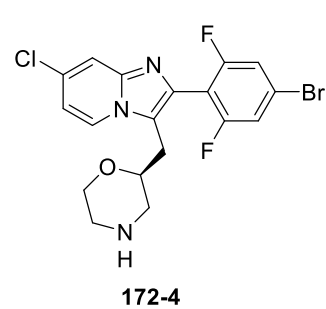
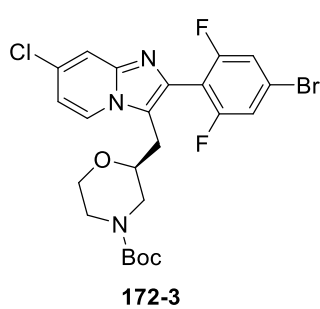
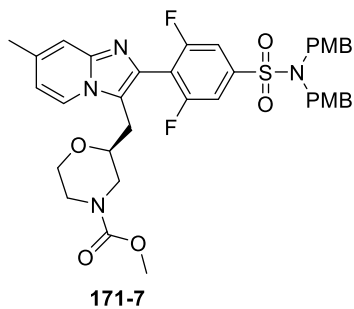
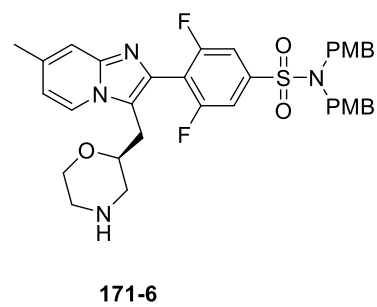
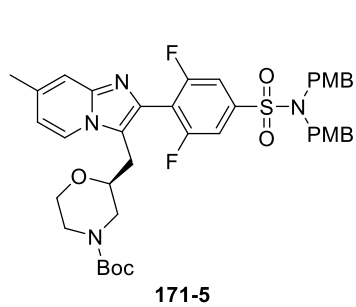
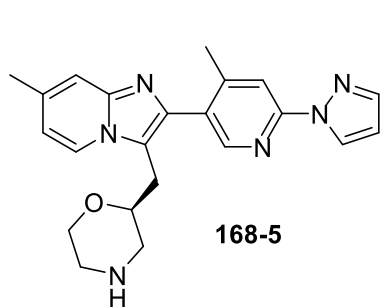


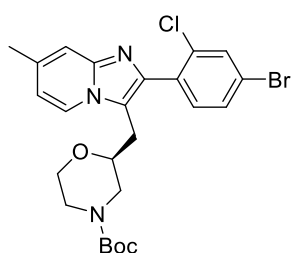




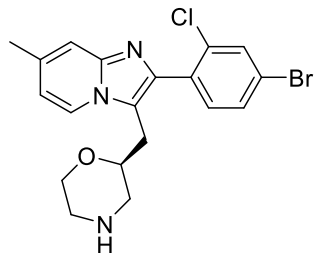




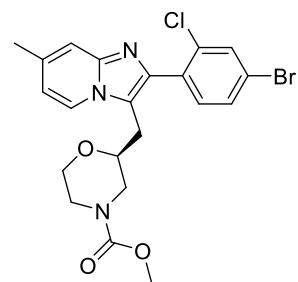




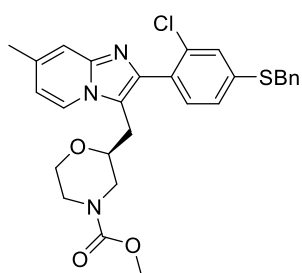
176-2



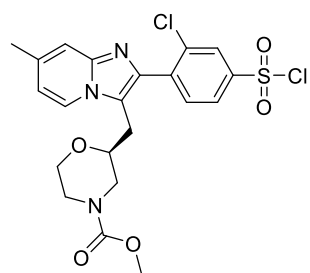
176-3



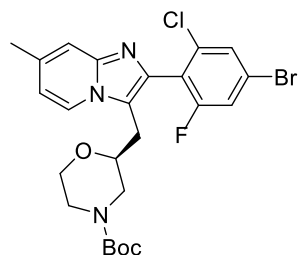
176-4



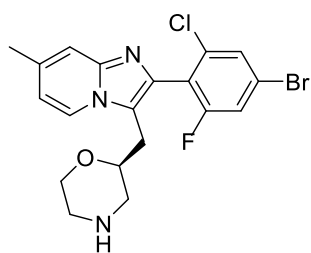
176-5



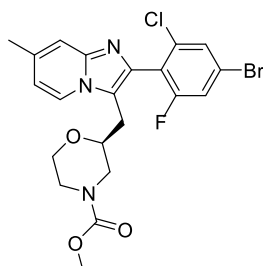
176-6



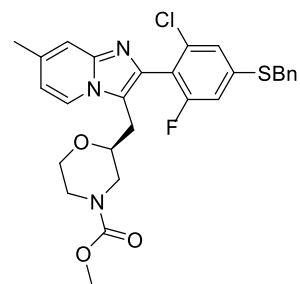
177-3



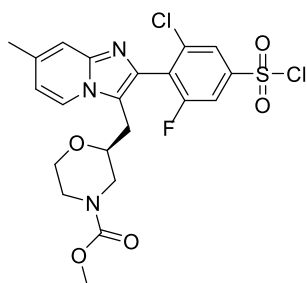
177-4



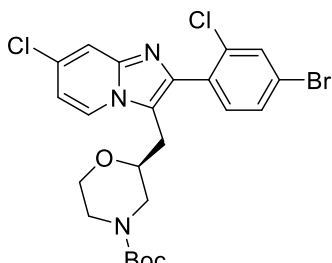
177-5



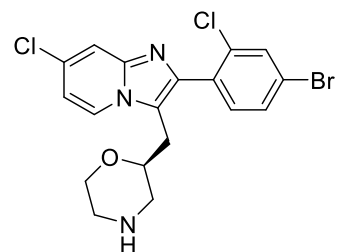
177-7



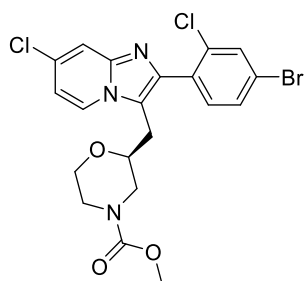
177-8



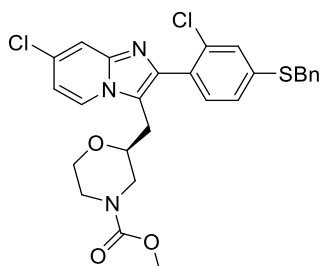
178-2



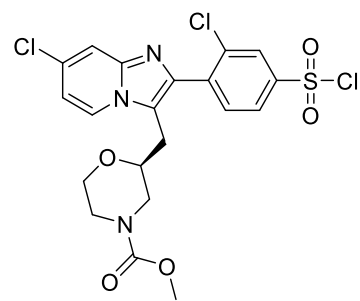
178-3



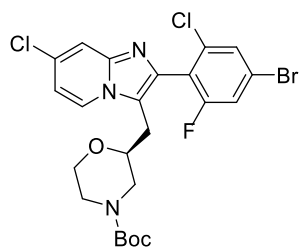
178-4



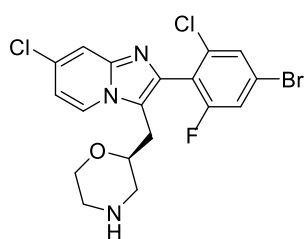
178-5



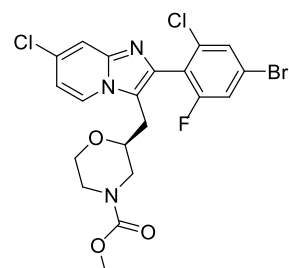
178-6



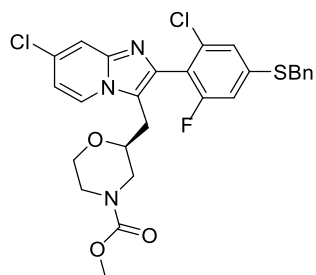
179-2



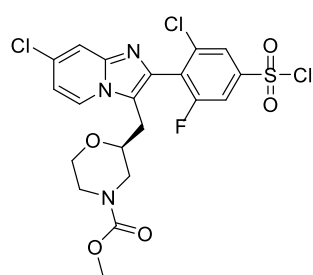
179-3



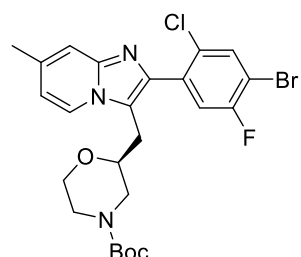
179-4



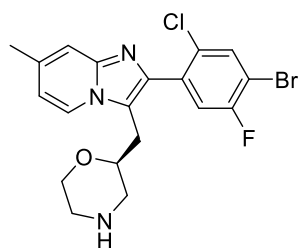
179-5



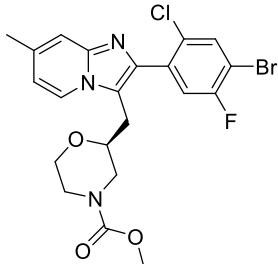
179-6



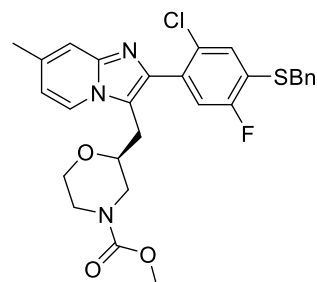
180-2



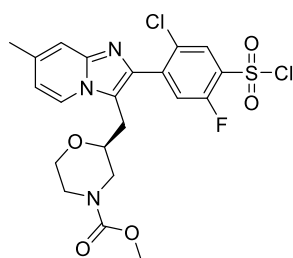
180-3



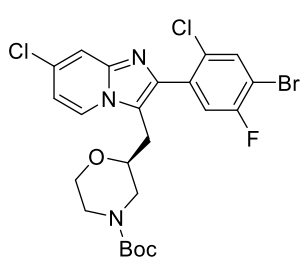
180-4



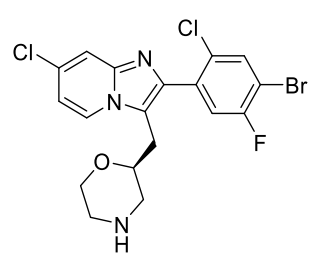
180-5



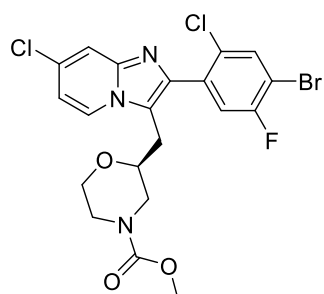
180-6



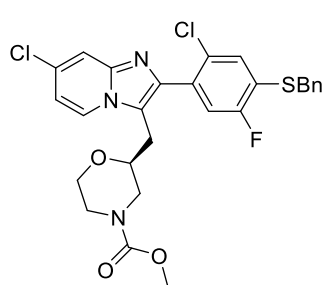
181-1



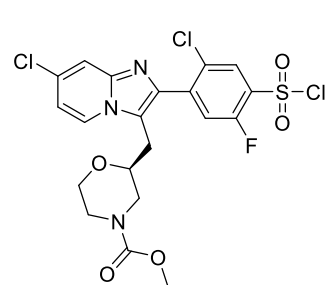
181-2



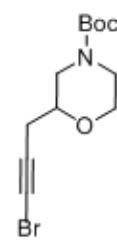
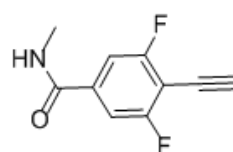
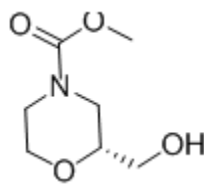
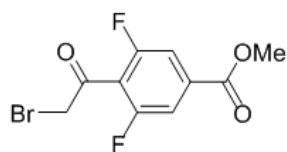
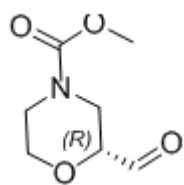
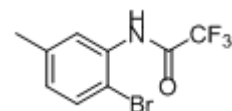
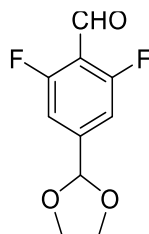
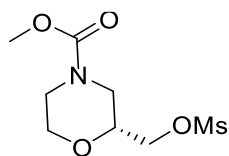
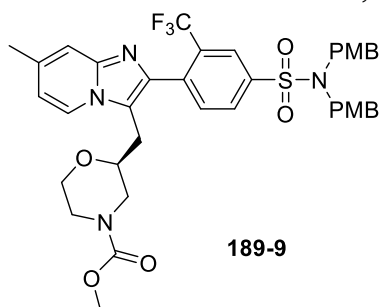
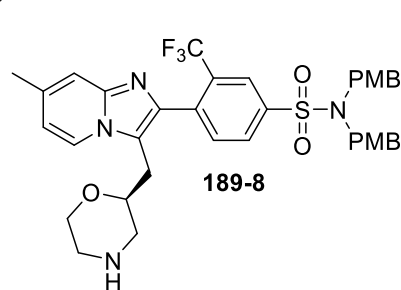
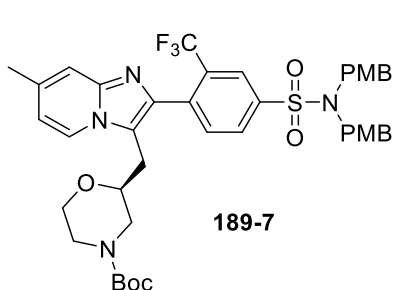
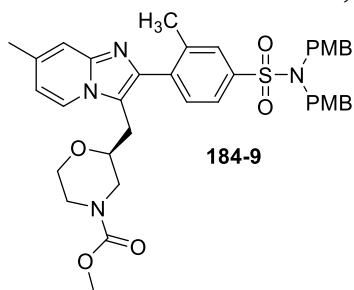
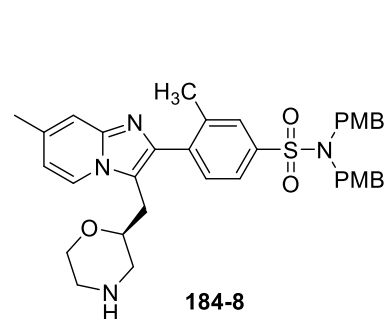
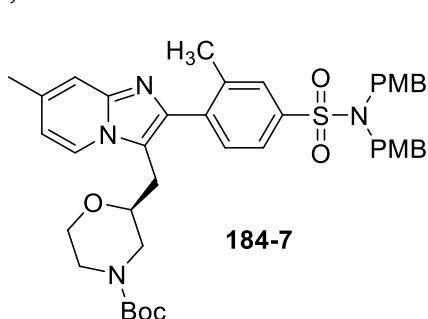
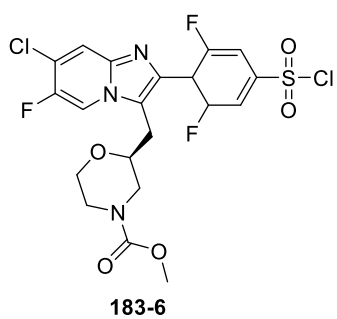
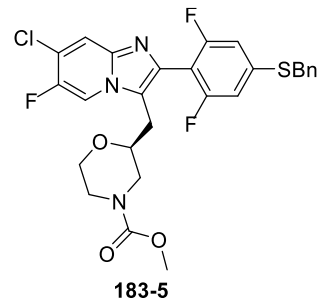
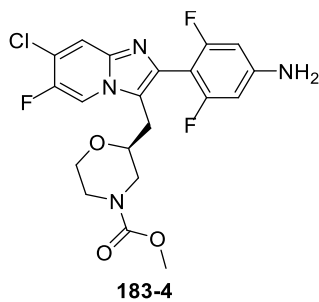
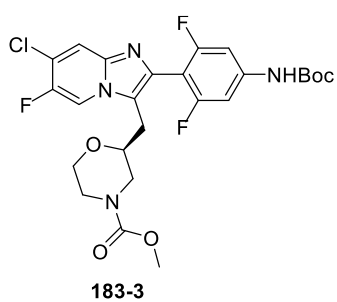
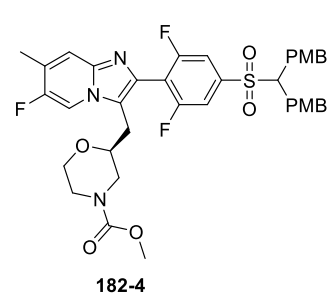
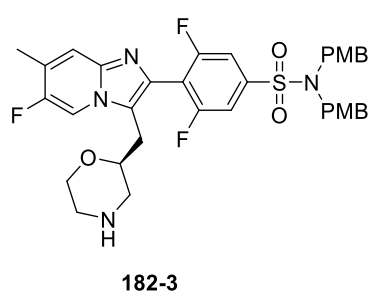
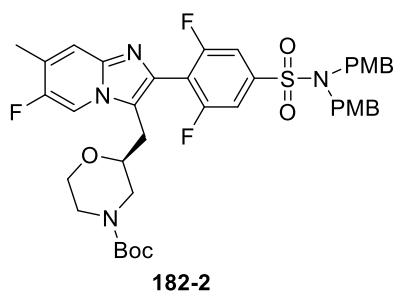
181-3



181-4



181-5



5

The present invention also provides a pharmaceutical composition comprising the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof and at least one pharmaceutically acceptable excipient. The pharmaceutical composition may also comprise one or more additional active ingredients. For example, such pharmaceutical composition may comprise one or more additional heterocyclic compounds of formula I or pharmaceutically acceptable salts, hydrates, solvates, metabolites, stereoisomers, tautomers or prodrugs thereof. Alternatively or additionally, the pharmaceutical composition may, for example, comprise one or more active ingredients other than a heterocyclic compound of formula I or a pharmaceutically acceptable salt, a hydrate, a solvate, a metabolite, a stereoisomer, a tautomer or a prodrug thereof.

The pharmaceutical composition comprise an therapeutically effective amount of the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof.

The effective amount of the compound, the pharmaceutical composition or the drug disclosed herein can be readily determined by routine experiments, and the most effective and convenient administration route and the most appropriate preparation can also be determined by routine experiments.

The pharmaceutically acceptable excipients may be those widely used in the field of pharmaceutical production. The excipients are primarily used to provide a safe, stable and functional pharmaceutical composition and may also provide methods for dissolving the active ingredients at a desired rate or for promoting effective absorption of the active ingredients after administration of the composition to a subject. The pharmaceutically acceptable excipients may be inert fillers or provide a function such as stabilizing the overall pH of the composition or preventing degradation of the active ingredients of the composition. The pharmaceutically acceptable excipients may include one or more of the following: binders, suspending agents, emulsifiers, diluents, fillers, granulating agents, gluing agents, disintegrants, lubricants, anti-adherents, glidants, wetting agents, gelling agents, absorption delaying agents, dissolution inhibitors, reinforcing agents, adsorbents, buffering agents, chelating agents, preservatives, colorants, flavoring agents and sweeteners.

Substances that may serve as pharmaceutically acceptable excipients include, but are not limited to, ion exchangers; aluminum; aluminum stearate; lecithin; serum proteins such as human serum protein; buffer substances such as phosphate; glycine; sorbic acid; potassium sorbate; partial glyceride mixture of saturated vegetable fatty acid; water; salts or electrolytes such as protamine sulfate, disodium hydrogen phosphate, potassium hydrogen phosphate, sodium

chloride and zinc salt; colloidal silica; magnesium trisilicate; polyvinylpyrrolidone; polyacrylate; waxes; polyethylene-polyoxypropylene-blocking polymer; wool fat; sugars such as lactose, glucose and sucrose; starches such as corn starch and potato starch; cellulose and its derivatives such as sodium carboxymethyl cellulose, ethyl cellulose and cellulose acetate; gum powder; malt; gelatin; talc powder; adjuvants such as cocoa butter and suppository waxes; oils such as peanut oil, cottonseed oil, safflower oil, sesame oil, olive oil, corn oil and soybean oil; glycols such as propylene glycol and polyethylene glycol; esters such as ethyl oleate and ethyl laurate; agar; buffers such as magnesium hydroxide and aluminum hydroxide; alginic acid; pyrogen-free water; isotonic salts; Ringer's solution; ethanol, phosphate buffered solution, and other non-toxic suitable lubricants such as sodium lauryl sulfate and magnesium stearate, coloring agents, releasing agents, coating agents, sweeteners, flavoring agents and perfumes, preservatives and antioxidants.

The pharmaceutical composition disclosed herein may be prepared in accordance with the disclosure by any method known to those skilled in the art, for example, conventional mixing, dissolving, granulating, emulsifying, levigating, encapsulating, embedding or lyophilizing processes.

The compound disclosed herein may be provided in pharmaceutical dosage forms including immediate release, controlled release, sustained release or targeted drug release systems. For example, common dosage forms include solutions and suspensions, (micro)emulsions, ointments, gels and patches, liposomes, tablets, dragees, soft or hard shell capsules, suppositories, ovules, implants, amorphous or crystalline powders, aerosols and lyophilized preparations. Depending on the administration route, special devices may be required to administer or deliver the drugs, such as syringes and needles, inhalers, pumps, injection pens, applicators or special flasks. Pharmaceutical dosage forms often comprises drugs, excipients, and containers/closure systems. One or more excipients (also known as inactive ingredients) may be added to the compound disclosed herein to improve or facilitate the manufacture, stability, administration and safety of drugs, and may provide a means to obtain a desired drug release profile. Thus, the types of the excipients to be added to a drug may depend on various factors, such as physical and chemical properties of the drug, administration route and preparation methods. Those pharmaceutical excipients in the art, including those listed in various pharmacopoeias, may be used. (See *U.S. Pharmacopeia* (USP), *Japanese Pharmacopoeia* (JP), *European Pharmacopoeia* (EP) and *British Pharmacopoeia* (BP); publications from the Center for Drug Evaluation and Research (CEDR) of U.S. Food and Drug Administration

(www.fda.gov), for example, *Inactive Ingredient Guide*, 1996; and *Handbook of Pharmaceutical Additives*, 2002, edited by Ash & Ash, Synapse Information Resources, Inc., Endicott NY; etc.)

Pharmaceutical dosage forms of the compound disclosed herein may be prepared by any method well known in the art, for example, by conventional mixing, sieving, dissolving, melting, granulating, dragee-making, tableting, suspending, extruding, spray-drying, grinding, emulsifying, (nano/micro)encapsulating, embedding or lyophilizing processes. As described above, the composition disclosed herein may include one or more physiologically acceptable inactive ingredients that facilitate the processing of active molecules into preparations for pharmaceutical use.

Appropriate preparations will be determined according to the desired administration route. For example, for intravenous injection, the composition may be prepared in aqueous solution and, if necessary, physiologically compatible buffers, including, for example, phosphate, histidine or citrate for adjusting the pH of the preparation, and tonicity agents such as sodium chloride or dextrose. For transmucosal or nasal administration, a semi-solid, liquid preparation or patch, possibly containing a penetration enhancer, may be preferred; and the penetration enhancer is generally known in the art. For oral administration, the compound may be prepared into liquid or solid dosage forms and used as immediate release or controlled/sustained release preparations. Suitable dosage forms for oral ingestion by an individual include tablets, pills, dragees, hard and soft shell capsules, liquids, gels, syrups, ointments, suspensions and emulsions. The compound may also be prepared in rectal compositions such as suppositories or retention enemas, e.g., containing conventional suppository matrices such as cocoa butter or other glycerides.

Oral solid dosage forms can be obtained using excipients including fillers, disintegrants, binders (dry and wet), dissolution retardants, lubricants, glidants, anti-adherents, cationic exchange resins, wetting agents, antioxidants, preservatives, colorants and flavoring agents. These excipients may be of synthetic or natural sources. Examples of such excipients include cellulose derivatives, citric acid, dicalcium phosphate, gelatin, magnesium carbonate, magnesium/sodium lauryl sulfate, mannitol, polyethylene glycol, polyvinylpyrrolidone, silicates, silica, sodium benzoate, sorbitol, starches, stearic acid or salts thereof, sugars (i.e., dextrose, sucrose, lactose, etc.), talc, tragacanth mucilage, vegetable oils (hydrogenated) and waxes. Ethanol and water may be used as adjuvants for granulation. In some cases, it is desirable to coat tablets with, for example, a taste-masking film, a gastric acid-resistant film or a sustained release film. Natural and synthetic polymers are often used to coat tablets in combination with colorants, sugars, and organic solvents or water to produce dragees. When capsules are superior

to tablets, the pharmaceutical powders, suspensions or solutions may be delivered in the form of compatible hard or soft shell capsules.

In some embodiments, the compound disclosed herein may be administered topically, e.g. by means of a skin patch, a semi-solid or liquid preparation such as gels, (micro)emulsions, ointments, solutions, (nano/micro)suspensions or foams. Skin and underlying tissue penetration of drugs may be modulated by: for example, using penetration enhancers; using appropriate selection and combination of lipophilic, hydrophilic and amphiphilic excipients, including water, organic solvents, waxes, oils, synthetic and natural polymers, surfactants, and emulsifiers; by adjusting the pH value; and using complexing agents. Other techniques, such as iontophoresis, may also be used to modulate skin penetration of the compound disclosed herein. Transdermal or topical administration will be preferred, for example, in situations where topical administration with minimal systemic exposure is desired.

For inhalation administration or nasal administration, the compound used herein is conveniently administered in the form of solutions, suspensions, emulsions or semi-solid aerosols from pressurised packs or nebulisers, usually with the aid of propellants, such as halocarbons derived from methane and ethane, carbon dioxide or any other suitable gas. For topical aerosols, hydrocarbons such as butane, isobutene and pentane are suitable. In the case of a pressurized aerosol, the appropriate dosage unit may be determined by providing a valve delivery measurement. Capsules and cartridges having, for example, gelatin may be prepared for use in an inhaler or insufflator. These generally contain powder mixtures of compounds with suitable powder matrices such as lactose or starch.

Compositions prepared for parenteral administration by injection are generally sterile and may be presented in unit dosage form, for example, in ampoules, syringes, injection pens, or multiple-dose containers, the latter typically containing preservatives. The compositions may take such forms as suspensions, solutions, or emulsions in oily or aqueous carriers, and may contain preparations such as buffers, tonicity agents, viscosity enhancers, surfactants, suspending and dispersing agents, antioxidants, biocompatible polymers, chelating agents, and preservatives. Depending on the injection site, the carrier may contain water, synthetic or vegetable oils, and/or organic co-solvents. In some cases, e.g., for lyophilized products or concentrates, parenteral preparations will be reconstituted or diluted prior to administration. Depot preparations that provide controlled or sustained release of the compound disclosed herein may include injectable suspensions of nano/microparticles or nano/micro or non-micronized crystals. Other matrices well known in the art, polymers such as poly(lactic acid), poly(glycolic acid) or copolymers thereof, may be used as the controlled/sustained release matrices. Other

depot administration systems may be presented in the form of implants and pumps requiring incisions.

Suitable carriers for the compound disclosed herein for intravenous injection are well known in the art and include water-based solutions containing a base (e.g., sodium hydroxide) for forming an ionic compound, sucrose or sodium chloride as a tonicity agent, and buffers, for example, containing phosphate or histidine. A co-solvent such as polyethylene glycol may be added. These water-based systems can effectively dissolve the compound disclosed herein and produce low toxicity after systemic administration. The proportion of components of the solution system can be varied considerably without destroying the solubility and toxicity characteristics. In addition, the properties of the components may be varied. For example, low toxicity surfactants such as polysorbate or poloxamer may be used, polyethylene glycol or other co-solvents may also be used, biocompatible polymers such as polyvinylbenzidine may be added, and other sugars and polyols may be used in place of dextrose.

The therapeutically effective dosage may first be estimated using various methods well known in the art. Initial dosage for animal studies may be based on established effective concentrations in cell culture assays. Dosage ranges suitable for human individuals may be determined, for example, using data obtained from animal studies and cell culture assays. In certain embodiments, the compound disclosed herein may be prepared as medicaments for oral administration.

An effective or therapeutically effective amount or dosage of a medicament (e.g., the compound disclosed herein) refers to the amount of the medicament or compound that causes improvement of symptoms or prolongation of survival in an individual. Toxicity and therapeutic efficacy of the molecule can be determined by standard pharmaceutical procedures in cell cultures or experimental animals, e.g., by measuring LD₅₀ (the dosage lethal to 50% of the population) and ED₅₀ (the dosage therapeutically effective in 50% of the population). The dose ratio of toxic to therapeutic effects is the therapeutic index and can be expressed as LD₅₀/ED₅₀. Medicaments that exhibit high therapeutic indexes are preferred.

An effective or therapeutically effective amount is the amount of the compound or pharmaceutical composition that will elicit the biological or medical response of a tissue, system, animal or human that is being sought by the researcher, veterinarian, physician or other clinicians. The dosage is preferably within a range of circulating concentrations that include ED₅₀ with minimal or no toxicity. The dosage may vary within this range depending on the dosage form employed and/or the administration route utilized. The correct preparation,

administration route, dosage and time interval between administrations should be selected based on methods known in the art while taking the specificity of the individual condition into account.

The dosage and time interval may be adjusted individually to provide plasma levels of the active moiety sufficient to achieve the desired effect; i.e., the minimum effective concentration (MEC). The MEC for each compound will vary, but can be estimated, for example, from *in vitro* data and animal experiments. The dosage necessary to obtain MEC will depend on the individual characteristics and administration route. In the case of topical administration or selective uptake, the effective local concentration of drugs may not be related to the plasma concentration.

The amount of the medicament or composition administered may depend on various factors, including the gender, age and weight of a subject being treated, the severity of affliction, the administration route and the judgment of the prescribing physician.

When required, the composition disclosed herein may be presented in a package or with a dispensing device containing one or more unit dosage forms containing active ingredients. For example, the package or device may comprise a metal or plastic foil (such as a foaming package) or glass and a rubber stopper, such as in a vial. The package or dispensing device may be accompanied by instructions for administration. The composition comprising the compound disclosed herein formulated in a compatible pharmaceutical carrier may also be prepared, placed in an appropriate container, and labeled for treatment of a specified condition.

The present invention also provides use of the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof or the pharmaceutical composition in the manufacture of a P2X3 inhibitor.

The present invention also provides use of the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof or the pharmaceutical composition in the manufacture of a medicament. The medicament is used for preventing, dealing with, treating or ameliorating diseases in an animal mediated at least in part by P2X3 or associated with P2X3 activity; alternatively, the medicament is used for treating pains, urinary tract diseases or respiratory diseases.

In some embodiments, the medicament may be used for treating urinary tract diseases in an animal (e.g., a human) including, but not limited to: urinary incontinence, overactive bladder, dysuria, or cystitis.

In some embodiments, the medicament may be used for treating respiratory diseases in an animal (e.g., a human) including, but not limited to: respiratory disorders including idiopathic

pulmonary fibrosis, chronic obstructive pulmonary disease, asthma, bronchospasm, or chronic cough.

In some embodiments, the medicament may be used for treating pains in an animal (e.g., a human) including, but not limited to: inflammatory pain, surgical pain, visceral pain, dental pain, premenstrual pain, central pain, pain caused by burns, migraine, cluster headache, or chronic pain.

The present invention also provides use of the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof or the pharmaceutical composition in treating and preventing diseases.

The present invention also provides use of the heterocyclic compound of formula I or the pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof or the pharmaceutical composition in preventing, dealing with, treating or ameliorating diseases in an animal (e.g., a human) mediated at least in part by P2X3 or associated with P2X3 activity. The diseases include, but are not limited to: respiratory diseases, cough, chronic cough, idiopathic pulmonary fibrosis, chronic lung obstruction, asthma, pains, urinary incontinence, autoimmune diseases, overactive bladder, dysuria, inflammation, senile dementia, Parkinson's disease, sleep disorders, epilepsy, psychiatric diseases, arthritis, neurodegeneration, traumatic brain injury, myocardial infarction, rheumatoid arthritis, brain stroke, thrombosis, atherosclerosis, colon syndrome, inflammatory bowel diseases, digestive tract diseases, gastrointestinal dysfunction, respiratory failure, sexual dysfunction, cardiovascular system diseases, heart failure, hypertension, urinary incontinence, cystitis, arthritis, endometriosis, hematopathy, musculoskeletal and connective tissue development disorders, or systemic disorder diseases.

In some embodiments, the diseases include pains including, but not limited to: inflammatory pain, surgical pain, visceral pain, dental pain, premenstrual pain, central pain, pain caused by burns, migraine, or cluster headache.

In some embodiments, the diseases include urinary tract diseases.

In some embodiments, the diseases include respiratory diseases including, but not limited to: respiratory disorders including idiopathic pulmonary fibrosis, chronic obstructive pulmonary disease, asthma, bronchospasm, or chronic cough.

The present invention further provides a method for treating or preventing diseases, which comprises administering an effective dosage of the heterocyclic compound of formula I or the

pharmaceutically acceptable salt, the hydrate, the solvate, the metabolite, the stereoisomer, the tautomer or the prodrug thereof or the pharmaceutical composition to a patient.

Definitions and General Terms

Unless otherwise stated, all technical and scientific terms used herein have the same meaning as commonly understood by those skilled in the art to which the present invention belongs. All patents and publications referred to herein are incorporated herein by reference in their entirety.

Unless otherwise stated, the following definitions used herein should be applied. For the purposes of the present invention, the chemical elements are in accordance with the Periodic Table of the Elements (CAS version), and the "Handbook of Chemistry and Physics", 75th edition, 1994. In addition, general principles of organic chemistry can be found in the descriptions of "Organic Chemistry", Thomas Sorrell, University Science Books, Sausalito:1999, and "March's Advanced Organic Chemistry" by Michael B.Smith and Jerry March, John Wiley & Sons, New York: 2007, which are incorporated herein by reference in their entirety.

The term "include", "includes" or "including" is open-ended, i.e., including what is meant by the present invention, but not excluding other aspects.

The term "stereoisomer" refers to compounds having the same chemical structure, but different spatial arrangements of the atoms or groups. Stereoisomers include enantiomers, diastereomers, conformers (rotamers), geometric isomers (cis/trans), atropisomers, and the like.

The term "enantiomer" refers to two isomers of a compound that do not overlap but are mirror images of each other.

The term "diastereoisomer" refers to stereoisomers having two or more chiral centers and whose molecules are not mirror images of each other. Diastereomers have different physical properties, such as melting points, boiling points, spectral properties, and reactivities. Mixtures of diastereomers can be separated by high resolution analytical procedures such as electrophoresis and chromatography (e.g., HPLC).

The stereochemical definitions and rules used herein generally follow S.P. Parker, Ed., McGraw-Hill Dictionary of Chemical Terms (1984) McGraw-Hill Book Company, New York; and Eliel, E. and Wilen, S., "Stereochemistry of Organic Compounds", John Wiley & Sons, Inc., New York, 1994.

Any asymmetric atom (e.g., carbon, etc.) of the compound disclosed herein can exist in a racemate or enantiomer enriched form, e.g., the (R)-, (S)- or (R,S)-configuration. In certain embodiments, each asymmetric atom has at least 0% enantiomeric excess, at least 60%

enantiomeric excess, at least 70% enantiomeric excess, at least 80% enantiomeric excess, at least 90% enantiomeric excess, at least 95% enantiomeric excess, or at least 99% enantiomeric excess in the (R)- or (S)-configuration.

Any resulting mixture of stereoisomers may be separated into pure or substantially pure geometric isomers, enantiomers and diastereomers depending on the differences in the physicochemical properties of the components, for example, by chromatography and/or fractional crystallization.

The term "tautomer" or "tautomeric form" refers to structural isomers having different energies that are interconvertible by a low energy barrier. If a tautomer is possible (e.g., in solution), the chemical equilibrium of the tautomer can be reached. For example, a proton tautomer, also known as a prototropic tautomer, includes the interconversion by proton transfer, such as keto-enol isomerization and imine-enamine isomerization. A valence tautomer includes the interconversion by recombination of some bonding electrons. A specific example of keto-enol tautomerization is the interconversion of the pentan-2,4-dione and 4-hydroxypent-3-en-2-one tautomers. Another example of tautomerization is phenol-keto tautomerization. A specific example of phenol-keto tautomerization is the interconversion of pyridin-4-ol and pyridin-4(1*H*)-one tautomers. Unless otherwise indicated, all tautomeric forms of the compound disclosed herein are within the scope of the present invention.

In general, the term "substituted" means that one or more hydrogen atoms in a given structure are substituted with a particular substituent. Further, when the group is substituted with one or more substituents described above, the substituents are independent of each other, that is, the one or more substituents may be different from each other or the same. Unless otherwise indicated, the substitution of a substituent may occur at various substitutable positions of the substituted group. When more than one position in a given structure can be substituted with one or more substituents selected from particular groups, the substitution of the substituents may occur at various positions, identically or differently.

As will be understood by those skilled in the art, " $\overset{\text{z}}{\text{---}}$ " used in the structural formula describing groups herein means that the corresponding groups are connected with other fragments and groups in the compound through this site, according to conventions used in the art.

In each part of this specification, substituents for the disclosed compound are disclosed according to group types or ranges. It is specifically noted that each separate sub-combination of the various members of these group types and ranges is encompassed by the present invention. For example, the term "C₁-C₆ alkyl" or "C₁₋₆ alkyl" may be linear or branched, and specifically

refers to the independently disclosed methyl, ethyl, C₃ alkyl, C₄ alkyl, C₅ alkyl, and C₆ alkyl; "C₁₋₄ alkyl" specifically refers to the independently disclosed methyl, ethyl, C₃ alkyl (i.e., propyl, including *n*-propyl and isopropyl), and C₄ alkyl (i.e., butyl, including *n*-butyl, isobutyl, *sec*-butyl, and *tert*-butyl).

5 In each part of the present invention, connecting substituents are described. When a connecting group is clearly required to the structure, the Markush variables listed for that group should be considered as connecting groups. For example, if a connecting group is required to the structure and the "alkyl" or "aryl" is listed for the Markush group definition of the variable, it is to be understood that the "alkyl" or "aryl" represents a connected alkylene group or arylene group, respectively.

10 The term "alkyl" used herein refers to linear or branched saturated monovalent hydrocarbon radical containing 1-20 carbon atoms. Examples of alkyl groups include, but are not limited to, methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl, *tert*-butyl, *n*-pentyl, 2-pentyl, 3-pentyl, 2-methyl-2-butyl, 3-methyl-2-butyl, 3-methyl-1-butyl, 2-methyl-1-butyl, *n*-hexyl, 2-hexyl, 3-hexyl, 2-methyl-2-pentyl, 3-methyl-2-pentyl, 4-methyl-2-pentyl, 3-methyl-3-pentyl, 2-methyl-3-pentyl, 2,3-dimethyl-2-butyl, 3,3-dimethyl-2-butyl, *n*-heptyl, *n*-octyl, and the like.

In some specific structures, when an alkyl group is explicitly indicated as a connecting group, the alkyl group represents a connected alkylene group, e.g., C₁-C₆ alkyl in the group "halogenated C₁-C₆ alkyl" should be considered as C₁-C₆ alkylene.

20 The term "alkylene" refers to a saturated divalent hydrocarbon radical resulting from the removal of two hydrogen atoms from a linear or branched saturated hydrocarbon radical. Examples of alkylene groups include methylene (-CH₂-), ethylidene (including -CH₂CH₂- or -CH(CH₃)-), isopropylene (including -CH(CH₃)CH₂- or -C(CH₃)₂-), and the like.

25 The term "alkenyl" refers to a linear or branched monovalent hydrocarbon radical containing 2-12 carbon atoms, wherein there is at least one unsaturated site, i.e., a carbon-carbon sp² double bond, which includes the positioning of "cis" and "trans", or the positioning of "E" and "Z". Examples of alkenyl groups include, but are not limited to, ethenyl (-CH=CH₂), allyl (-CH₂CH=CH₂), and the like.

30 The term "alkoxy" or "alkyl-O-" means that the alkyl group is connected with the rest of the molecule through an oxygen atom, wherein the alkyl group are described as herein. Examples of alkoxy groups include, but are not limited to, methoxy, ethoxy, 1-propoxy, 2-propoxy, 1-butoxy and the like.

The term "haloalkyl", "haloalkoxy" or "haloalkyl-O-" means that the alkyl or alkoxy group is substituted with one or more halogen atoms, examples of which include, but are not limited to, trifluoromethyl, trifluoromethoxy and the like.

The term "cycloalkyl" refers to a monovalent or polyvalent saturated monocyclic or bicyclic system containing 3-12 ring carbon atoms. Examples of cycloalkyl groups include, but are not limited to, cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl, cyclooctyl, and the like; wherein the C₃-C₆ cycloalkyl includes cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl.

The term "cycloalkenyl" refers to an unsaturated monocyclic carbocycloalkenyl group containing 3-6 ring carbon atoms ("C₃-C₆ cycloalkenyl"). Examples of cycloalkenyl groups include, but are not limited to, cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl or cyclohexadienyl, and the like.

The term "heterocycloalkyl" refers to a saturated monocyclic or bicyclic system containing 3-10 ring atoms, which can include fused, bridged, or spiro ring systems (e.g., bicyclic systems ("bicyclic heterocycloalkyl")); wherein the monocyclic or bicyclic system contains 3-9 ring carbon atoms and at least one ring heteroatom selected from nitrogen, sulfur, and oxygen ("C₃-C₉ heterocycloalkyl"). The bicyclic heterocycloalkyl system may include one or more heteroatoms in one or two rings, and is saturated. Unless otherwise stated, heterocycloalkyl may be carbon- or nitrogen-based, and the -CH₂-group may be optionally substituted with -C(=O)-. The sulfur atom of the ring may be optionally oxidized to an S-oxide. The nitrogen atom of the ring may be optionally oxidized to an N-oxide. In some embodiments, heterocycloalkyl is C₃-C₅ heterocycloalkyl, meaning that heterocycloalkyl contains 3-5 ring carbon atoms and at least one ring heteroatom selected from O, S and N. Examples of heterocycloalkyl include, but are not limited to: oxiranyl, thietanyl, pyrrolidinyl, pyrazolinyl, pyrazolidinyl, imidazolinyl, imidazolidinyl, oxazolidinyl, tetrahydrofuranyl, piperidinyl, morpholinyl, tetrahydropyrimidinyl, oxazinanyl, thiomorpholinyl, piperazinyl, and the like. Examples of the substitution of the -CH₂-group with -C(=O)- in the heterocycloalkyl include, but are not limited to, 2-oxopyrrolidinyl, 2-piperidonyl, 3-morpholinonyl, 3-thiomorpholinonyl, and oxotetrahydropyrimidinyl, and the like, or isomers and stereoisomers thereof. In some embodiments, exemplary C₃-C₉ heterocycloalkyl groups include, but are not limited to, the C₃-C₅ heterocycloalkyl groups described above in conjunction with azepanyl, oxepanyl, thiepanyl, diazepanyl, azocanyl, oxecanyl, thiocanyl, quinuclidinyl, octahydroindolyl, octahydroisoindolyl, decahydroquinolyl, decahydroisoquinolyl, or isomers and stereoisomers thereof.

The term "heterocycloalkenyl" refers to a monocyclic or bicyclic system containing partially unsaturated alkenyl groups, which contains 3-9 ring carbon atoms and at least one ring

atom selected from nitrogen, sulfur, and oxygen atoms ("C₃-C₉ heterocycloalkenyl"); wherein the heterocycloalkenyl is non-aromatic and does not contain any aromatic ring. Unless otherwise stated, heterocycloalkenyl may be carbon- or nitrogen-based, and the -CH₂-group may be optionally substituted with -C(=O)-. The sulfur atom of the ring may be optionally oxidized to an *S*-oxide. The nitrogen atom of the ring may be optionally oxidized to an *N*-oxide. In some embodiments, heterocycloalkenyl is preferably C₃-C₅ heterocycloalkenyl; examples of C₃-C₅ heterocycloalkenyl include, but are not limited to, dihydrofuranyl, dihydrothienyl, dihydropyrrolyl, dioxolyl, dihydroimidazolyl, dihydropyrazolyl, dihydrothiazolyl, dihydroisothiazolyl, dihydrooxadiazolyl, dihydrothiadiazolyl, dihydrotriazolyl, dihydrotetrazolyl, tetrahydropyridinyl, 3,4-dihydro-2*H*-pyran, pyranyl, thiopyranyl, dihydropyridinyl, dihydropyrazinyl, dihydropyrimidinyl, oxazinyl, and dihydrotetrazolyl, or isomers and stereoisomers thereof. In some embodiments, exemplary C₃-C₉ heterocycloalkenyl groups include, but are not limited to, the C₃-C₅ heterocycloalkenyl groups described above in conjunction with octahydroisoquinolinyl, 3*H*-indolyl, dihydroisoquinolinyl, dihydroquinolyl or 4*H*-quinolizinyl, or isomers and stereoisomers thereof.

The term "halogen" or "halo" refers to fluorine (F), chlorine (Cl), bromine (Br), or iodine (I).

The term "aryl" refers to monocyclic, bicyclic and tricyclic carbon ring systems containing 6-14 ring atoms, or 6-10 ring atoms. Examples of aryl groups may include phenyl, naphthyl and anthryl. Unless otherwise stated, the group "C₆-C₁₀ aryl" refers to an aryl group containing 6-10 ring carbon atoms.

The term "heteroaryl" refers to monocyclic, bicyclic and tricyclic systems containing 5-6 ring atoms, or 5-10 ring atoms, or 5-12 ring atoms, wherein at least one ring contains one or more ring heteroatoms selected from nitrogen, oxygen, and sulfur. Unless otherwise stated, the heteroaryl group may be connected to the rest of the molecule (e.g., the host structure in the formula) via any reasonable site (which may be C in CH, or N in NH). Examples of heteroaryl groups include, but are not limited to, furanyl, imidazolyl, isoxazolyl, oxazolyl, pyrrolyl, pyrazolyl, pyridinyl, pyrimidinyl, pyridazinyl, pyrazinyl, thienyl, thiazolyl, and the like; and include, but are not limited to, the following bicyclic rings: benzimidazolyl, benzofuranyl, benzothienyl, indolyl, oxindolyl, imidazopyridinyl, pyrazolopyridinyl, pyrazolopyrimidinyl, quinolinyl, isoquinolinyl, quinazolinyl, and the like.

In addition, it should be noted that, unless otherwise explicitly indicated, the description of "... is independently" used herein is to be understood broadly and means that each individual group described is independent from the others and may be independently the same or different

specific groups. In more detail, the description of "... is independently" can mean that the specific options expressed by the same symbols in different groups do not affect each other; it can also mean that the specific options expressed by the same symbols in the same group do not affect each other.

5 The term "pharmaceutically acceptable" refers to molecular entities and compositions that are physiologically tolerable and do not typically produce an allergic or similar untoward reaction, such as gastrointestinal distress and dizziness, when administered to a human.

10 The term "carrier" refers to a diluent, adjuvant, excipient, or matrix with which the compound is administered. Such pharmaceutical carriers can be sterile liquid, such as water and oil, including those derived from petroleum, animals, plants or synthesis, such as peanut oil, soybean oil, mineral oil and sesame oil. Water or aqueous saline solutions and aqueous dextrose and glycerol solutions are preferably used as carriers, particularly injectable solutions. Suitable pharmaceutical carriers are described in "Remington's Pharmaceutical Sciences" by E.W. Martin.

15 The term "prodrug" used herein represents a compound that is converted *in vivo* to a compound of formula I. Such conversion is effected by hydrolysis of the prodrug in the blood or by enzymatic conversion of the prodrug into the parent structure in the blood or tissue. The prodrugs disclosed herein can be esters, and in the prior art, the esters that can be used as prodrugs include phenyl esters, aliphatic (C₁₋₂₄) esters, acyloxymethyl esters, carbonates, carbamates and
20 amino acid esters. For example, a compound disclosed herein containing hydroxyl can be acylated to give a prodrug. Other prodrugs include phosphate esters, and those phosphate esters are obtained by phosphorylating via the hydroxyl on the parent structure. For a complete discussion of prodrugs, reference can be made to the following: T. Higuchi and V. Stella, Prodrugs as Novel Delivery Systems, Vol.14 of the *A.C.S. Symposium Series*; Edward B. Roche, ed., *Bioreversible Carriers in Drug Design*, American Pharmaceutical Association and Pergamon Press, 1987, J. Rautio et al.; Prodrugs: Design and Clinical Applications, *Nature Review Drug Discovery*, 2008, 7, 255-270; and S. J. Hecker et al., Prodrugs of Phosphates and Phosphonates, *Journal of Medicinal Chemistry*, 2008, 51, 2328-2345.

25 The term "metabolite" used herein refers to a product obtained by the metabolism of a particular compound or salt thereof *in vivo*. Metabolites of a compound can be identified by
30 techniques well known in the art, and their activities can be characterized by assays as described herein. Such products may be obtained by the oxidation, reduction, hydrolysis, amidation, deamidation, esterification, defatting, enzymatic cleavage, and the like of the administered compound. Accordingly, the present invention includes metabolites of the compound, including

metabolites produced by contacting the compound disclosed herein with a mammal for a sufficient period of time.

The term "pharmaceutically acceptable salt" used herein refers to both organic and inorganic salts of the compound disclosed herein. Pharmaceutically acceptable salts are well known in the art. For example, S. M. Berge et al. describe pharmaceutically acceptable salts in detail in J. Pharmaceutical Sciences, 1977, 66: 1-19. Pharmaceutically acceptable salts formed by non-toxic acids include, but are not limited to, inorganic acid salts such as hydrochloride, hydrobromide, phosphate, sulfate, perchlorate; organic acid salts such as acetate, oxalate, maleate, tartrate, citrate, succinate, malonate; or salts obtained by other methods described in the literature, such as ion exchange. Other pharmaceutically acceptable salts include adipate, alginate, ascorbate, aspartate, benzenesulfonate, benzoate, bisulfate, borate, butyrate, camphorate, camphorsulfonate, cyclopentylpropionate, digluconate, dodecylsulfate, ethanesulfonate, formate, fumarate, glucoheptonate, glycerophosphate, gluconate, hemisulfate, heptanoate, hexanoate, hydriodate, 2-hydroxy-ethanesulfonate, lactobionate, lactate, laurate, lauryl sulfate, malate, methanesulfonate, 2-naphthalenesulfonate, nicotinate, nitrate, oleate, palmitate, pamoate, pectinate, persulfate, 3-phenylpropionate, picrate, pivalate, propionate, stearate, thiocyanate, *p*-toluenesulfonate, undecanoate, valerate, and the like. Salts formed by suitable bases include salts of alkali metal, alkaline earth metal, ammonium and $N^+(C_{1-4} \text{ alkyl})_4$. The present invention also contemplates quaternary ammonium salts formed by any compound containing an N group. Water-soluble or oil-soluble or dispersible products can be obtained by quaternization. Alkali metal or alkaline earth metal that can form salts include sodium, lithium, potassium, calcium, magnesium, and the like. Pharmaceutically acceptable salts further include suitable and non-toxic ammonium, quaternary ammonium salts and amine cations formed by counterions, such as halides, hydroxides, carboxylates, sulfates, phosphates, nitrates, C_{1-8} sulfonates and aromatic sulfonates.

"Solvate" disclosed herein refers to an association compound of one or more solvent molecules with the compound disclosed herein. Solvents that form the solvate include, but are not limited to, water, isopropanol, ethanol, methanol, dimethyl sulfoxide, ethyl acetate, acetic acid and aminoethanol. The term "hydrate" refers to an association compound of solvent molecules with water.

"Ester" disclosed herein refers to an ester that is hydrolyzed *in vivo* and formed by a compound containing hydroxyl or carboxyl. Such esters are, for example, pharmaceutically acceptable esters that are hydrolyzed in human or animal to produce parent alcohol or acid. The compound of formula I disclosed herein contains carboxyl, which can form an ester that is

hydrolyzed *in vivo* with appropriate groups including, but not limited to, alkyl, arylalkyl and the like.

"Nitrogen oxide" disclosed herein refers to an *N*-oxide formed by oxidizing one or more nitrogen atoms when the compound contains several amine functional groups. Specific examples of *N*-oxides are *N*-oxides of tertiary amines or *N*-oxides of nitrogen-containing heterocyclic nitrogen atoms. The corresponding amines can be treated with an oxidizing agent such as hydrogen peroxide or peracid (e.g., peroxyacetic acid) to form *N*-oxides (see Advanced Organic Chemistry, Wiley Interscience, 4th edition, Jerry March, pages). In particular, *N*-oxides may be prepared by the method of L. W. Deady (Syn. Comm. 1977, 7, 509-514) in which an amine compound is reacted with *m*-chloroperoxybenzoic acid (MCPBA), for example, in an inert solvent such as dichloromethane.

In some embodiments, the term "treat", "treating" or "treatment" used herein refers to ameliorating diseases or disorders (i.e., slowing or arresting or reducing the progression of the diseases or at least one clinical symptom thereof). In other embodiments, "treat", "treating" or "treatment" refers to mitigating or improving at least one physical parameter, including physical parameters that may not be perceived by a patient. In other embodiments, "treat", "treating" or "treatment" refers to modulating a disease or disorder, either physically (e.g., stabilizing a perceptible symptom) or physiologically (e.g., stabilizing a physical parameter), or both. In other embodiments, "treat", "treating" or "treatment" refers to preventing or delaying the onset, occurrence, or deterioration of a disease or disorder.

Unless otherwise indicated, abbreviations for any of protecting groups, amino acids and other compounds used herein are provided based on their commonly used and accepted abbreviations, or by referring to IUPAC-IUB Commission on Biochemical Nomenclature (see Biochem. 1972, 11:942-944).

The biological activity of the compound disclosed herein can be assessed by using any conventionally known method. Appropriate detection methods are well known in the art. For example, the compound disclosed herein can be tested for P2X₃ inhibitory activity, pharmacokinetic activity, and/or liver microsomal stability, etc., by an appropriate conventional method. The detection methods provided by the present invention are presented as examples only and do not limit the present invention. The compound disclosed herein is active in at least one of the detection methods provided by the present invention.

In this specification, terms such as "some embodiments," "examples," or "a preferred embodiment" mean that a particular feature, structure, material or characteristic described in reference to the embodiment or example is included in at least one embodiment or example of

the present invention. In this specification, the schematic descriptions of the terms used above do not necessarily refer to the same embodiment or example. Furthermore, the particular feature, structure, material or characteristic described may be combined in any one or more embodiments or examples in an appropriate manner. Moreover, various embodiments or examples and features of various embodiments or examples described in this specification can be combined by one skilled in the art to the extent that they do not contradict each other.

The above preferred conditions may be combined arbitrarily to obtain preferred embodiments of the present invention without departing from the general knowledge in the art.

The reagents and starting materials used in the present invention are commercially available.

The benefits of the present invention: the heterocyclic compound features high antagonistic activity against P2X₃, good selectivity, low toxicity, good metabolic stability and small influence on taste, showing a good prospect of drug development.

BRIEF DESCRIPTION OF THE DRAWINGS

FIG. 1 shows the number of citric acid-induced coughs for the inventive compounds vs. vehicle;

FIG. 2 shows the latency of citric acid-induced coughs for the inventive compounds vs. vehicle;

FIG. 3 shows the number of citric acid-induced coughs for the inventive compounds vs. their baseline;

FIG. 4 shows the number of coughs, cough suppression rate and cough latency obtained *in vivo* for a portion of the inventive compounds;

FIG. 5 shows the number of coughs for the inventive compounds in the ATP-citric acid model;

FIG. 6 shows the cough latency for the inventive compounds in the ATP-citric acid model;

FIG. 7 shows the ratio of quinine/tap water consumed in rats after the administration of the inventive compounds; and

FIG. 8 shows the ratio of quinine/tap water consumed in rats after the administration of the inventive compounds at gradient concentrations.

DETAILED DESCRIPTION

The present invention is further illustrated by the following examples, which are not intended to limit the invention thereto. Experimental procedures without specifying specific

conditions in the following examples are conducted in accordance with conventional procedures and conditions, or in accordance with commercial instructions.

The following abbreviations are used throughout the present invention:

TEMPO (2,2,6,6-tetramethyl-1-piperidone), LDA (lithium diisopropylamide), DMF (*N,N*-dimethylformamide), DMA (*N,N*-dimethylacetamide), DCM (dichloromethane), DME (dimethoxyethane), PE (petroleum ether), EA (ethyl acetate), DIPEA (*N,N*-diisopropylethylamine), THF (tetrahydrofuran), Ac (acetyl), MeOH (methanol), Boc (*tert*-butoxycarbonyl), B2Pin2 (bis(pinacolato)diboron), rt (room temperature), HATU (2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate), reflux (refluxing conditions), eq (equivalent), R_f (retardation factor), g (gram), mg (milligram), mol (mole), mmol (millimole), h (hour), min (minute), mL (milliliter), μ L (microliter).

Overnight refers to 8-15 h, for example 12 h; the room temperature refers to 10-30 °C; solvent ratio such as PE/EA refers to the volume ratio.

Unless otherwise indicated, all temperatures in the examples described below are given in Celsius degrees. Unless otherwise indicated, reagents are purchased from commercial suppliers such as Aldrich Chemical Company, Arco Chemical Company and Alfa Chemical Company, and used without further purification. General reagents are purchased from Shantou Xilong Chemical Plant Co. Ltd., Guangdong Guanghua Chemical Reagent Factory, Guangzhou Chemical Reagent Factory, Tianjin Haoyuyu Chemical Co., Ltd., Qingdao Tenglong Chemical Reagent Co., Ltd., and Qingdao Haiyang Chemical Co., Ltd.

Anhydrous tetrahydrofuran, dioxane, toluene and diethyl ether are obtained by refluxing and drying with metal sodium. Anhydrous dichloromethane and chloroform are obtained by refluxing and drying with calcium hydride. Ethyl acetate, petroleum ether, *n*-hexane, *N,N*-dimethylacetamide and *N,N*-dimethylformamide are pre-dried over anhydrous sodium sulfate.

The following reactions are generally conducted under a positive pressure of nitrogen or argon or by placing a drying tube over an anhydrous solvent (unless otherwise indicated), the reaction flask is stoppered with a suitable rubber stopper and the substrate is driven in by a syringe. Each piece of glassware is dried.

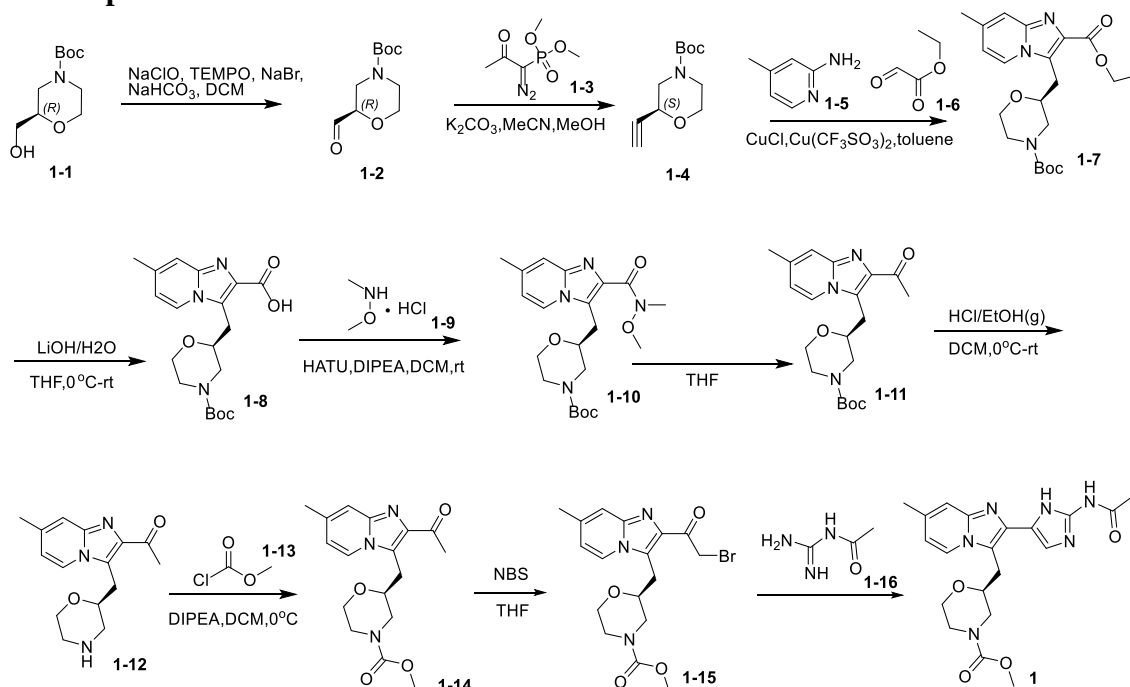
Chromatographic column is a silica gel column. Silica gel (300-400 mesh) is purchased from Qingdao Haiyang Chemical Co., Ltd. NMR spectral data are measured on a Bruker Avance 400 NMR spectrometer or a Bruker Avance IIIHD 600 NMR spectrometer using CDCl₃, DMSO-*d*₆, CD₃OD or Acetone-*d*₆ as solvents (reported in ppm) and TMS (0ppm) or chloroform (7.25ppm) as reference standards. When multiple peaks are present, the following abbreviations will be used: s (singlet), d (doublet), t (triplet), m (multiplet), br (broadened), dd (doublet of

doublets), dt (doublet of triplets), ddd (doublet of doublet of doublets), ddt (doublet of doublet of triplets), dddd (doublet of doublet of doublet of doublets). Coupling constants are expressed in hertz (Hz).

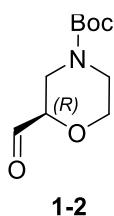
Low-Resolution Mass Spectrometry (MS) data are determined on an Agilent 6320 series LC-MS spectrometer equipped with a G1312A binary pump and an aG1316ATCC (column temperature maintained at 30 °C), with a G1329A autosampler and G1315BDAD detector applied to the analysis and an ESI source applied to the LC-MS spectrometer.

The above two spectrometers are equipped with an Agilent ZorbaxSB-C18 column (2.1×30 mm, 5 μm). The injection volume is determined by the sample concentration; the flow rate is 0.6 mL/min; peak values of HPLC are recorded at UV-Vis wavelength of 210 nm and 254 nm.

Example 1



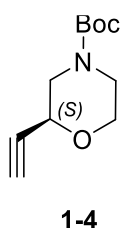
Step (1) Preparation of *tert*-butyl (*R*)-2-formyl-morpholine-4-carboxylate



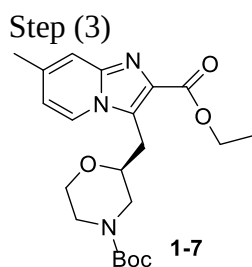
A mixture of **compound 1-1** (10 g, 460 mmol), TEMPO (0.073 g, 0.44 mmol), aqueous sodium bromide solution (0.5 M, 10 mL, 41 mmol) and dichloromethane (100 mL) was cooled to 0-5 °C. To a sodium hypochlorite solution (1.5 M, 34 mL, 58 mmol) was added sodium bicarbonate (2.3 g, 23 mmol), and the solution was adjusted to pH 9.3, and added dropwise to

the reaction system over 30 min. After the completion of the dropwise addition, the reaction system was stirred for 0.5 h, heated to 20 °C, and added with water (50 mL). The aqueous phase was extracted with dichloromethane multiple times. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated under reduced pressure. The residue was purified by column chromatography to give **compound 1-2** (6 g, 60% yield) in the form of an orange-yellow oily liquid. LC-MS: $[M+H]^+ = 216.4$.

Step (2) Preparation of *tert*-butyl (*S*)-2-ethynyl-morpholine-4-carboxylate



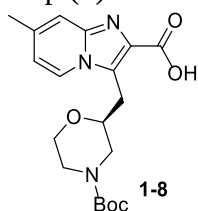
Dimethyl(1-diazo-2-oxopropyl)phosphonate (6 g, 31.3 mmol) was dissolved in a mixture of acetonitrile and methanol (5:1), and K_2CO_3 (59 mmol) was added. The reaction system was stirred at 20 °C for 15 min, and added dropwise with a dissolved solution of **compound 1-2** (6 g, 28 mmol) in a mixed solvent (10 mL) of acetonitrile and methanol (5:1) slowly. After the completion of the dropwise addition, the reaction system was incubated at this temperature overnight. The suspension was filtered, and the filtrate was concentrated under reduced pressure to give an oil, which was then added to water (about 100 mL) to precipitate a solid. The mixture was filtered, and the filter cake was washed with water. The residue was purified by silica gel column chromatography to give **compound 1-4** (3.4 g, 56.7% yield) in the form of a solid. LC-MS: $[M+H]^+ = 212.1$.



To a solution of **compound 1-4** (300 mg, 1.4 mmol), **compound 1-5** (145 mg, 1.4 mmol) and **compound 1-6** (145 mg, 1.4 mmol) in toluene (3 mL) were added CuCl (42 mg, 0.4 mmol) and $Cu(CF_3SO_3)_2$ (154 mg, 0.4 mmol) at room temperature. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with DME (0.1 mL), and reacted at this temperature for 5 h. After the reaction was completed, the reaction system was added with water (15 mL) to quench the reaction, extracted with dichloromethane (10 mL $\times$ 3), washed with saturated sodium chloride solution (10 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under

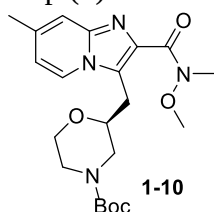
reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 1/1) to give **intermediate 1-7** (500 mg) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 404.2$.

Step (4)



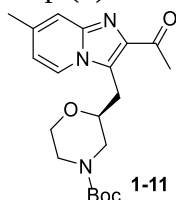
- 5 To a solution of **intermediate 1-7** (3 g, 7.44 mmol) in THF (20 mL) was added LiOH (936 mg, 22.3 mmol) at 0 °C. The reaction system was stirred at room temperature for 2 h. After the reaction was completed, the reaction system was adjusted to pH 6 with HCl (1 M) at 0 °C, extracted with DCM (10 mL $\times$ 3), washed with saturated sodium chloride solution (10 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure to give
- 10 **intermediate 1-8** (2 g) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 376.1$.

Step (5)



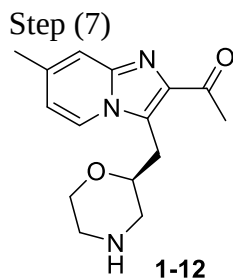
- Intermediate 1-8** (375 mg) was dissolved in DCM (10 mL), and HATU (2 mmol) and DIPEA (4 mmol) were added. The reaction system was stirred at room temperature for 15 min,
- 15 added with *N,O*-dimethylhydroxylamine hydrochloride (1.2 mmol), and stirred at room temperature for another 6-8 h. After the reaction was completed as monitored by TLC, the residue was purified by column chromatography to give **intermediate 1-10** (275 mg, 65.8% yield), LC-MS: $[M+H]^+ = 419.3$.

Step (6)

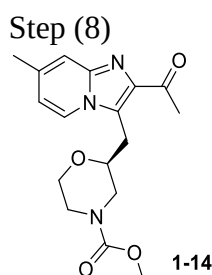


- 20 **Intermediate 1-10** (418 mg, 1 mmol) was dissolved in anhydrous THF (10 mL), and methyl Grignard reagent was added dropwise at 0 °C. After the completion of the dropwise addition, the reaction system was stirred overnight. After the reaction was completed, the reaction was quenched, and the reaction system was extracted with ethyl acetate, washed, dried, and purified

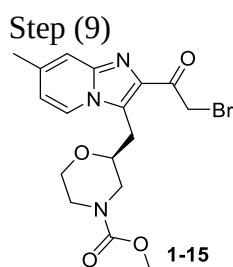
by column chromatography to give **intermediate 1-11** (236 mg, 63.3% yield). LC-MS: $[M+H]^+ = 374.1$.



5 **Intermediate 1-11** (373 mg) was dissolved in DCM (5 mL), and a solution of HCl in EtOH (1 N, 1.5 mL) was added dropwise at 0 °C. After the completion of the dropwise addition, the reaction system was stirred for 2 h, and poured into NaHCO₃ solution to adjust the pH to 8. The aqueous phase was extracted with DCM. The organic phase was washed with NaCl solution, dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under
10 reduced pressure. The residue was purified by column chromatography to give **intermediate 1-12** (160 mg, 58.6% yield). LC-MS: $[M+H]^+ = 274.6$.



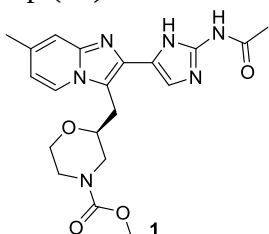
15 **Intermediate 1-12** (273 mg, 1 mmol) and DIPEA (258 mg, 2 mmol) were dissolved in DCM (5 mL), and methyl chloroformate (**compound 1-13**) (282 mg, 3 mmol) was added. The reaction system was stirred at 0 °C for 3 h, poured into H₂O and extracted with DCM. The organic phase was washed with NaCl solution, dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by column chromatography to give **intermediate 1-14** (300 mg, 91% yield) in the form of a yellow oil. LC-
20 MS: $[M+H]^+ = 332.2$.



Intermediate 1-14 (331 mg, 1 mmol) was dissolved in tetrahydrofuran (5 mL) and water (2 mL), and *N*-bromosuccinimide (214 mg, 1.2 mmol) was added. The reaction system was

reacted at room temperature for 1 h under nitrogen atmosphere. Then the reaction system was concentrated to half volume, diluted with water (10 mL), and extracted with ethyl acetate (10 mL $\times$ 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated to give **intermediate 1-15** (300 mg, 73.3% yield) in the form of a solid. LC-MS: $[M+H]^+ = 410.2$.

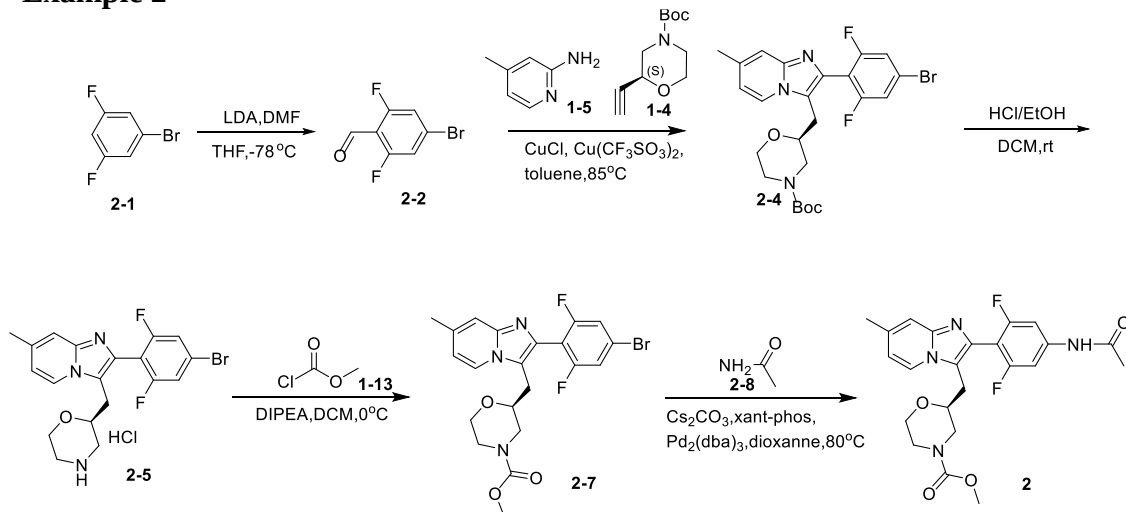
Step (10)



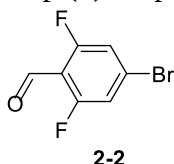
Intermediate 1-15 (0.82 g, 2 mmol) was dissolved in absolute ethanol (10 mL), and 1-acetylguanidine (242 mg, 2.4 mmol) and triethylamine (404 mg, 4 mmol) were added. The reaction system was heated to reflux. After the reaction was completed as detected by TLC, the reaction system was concentrated under reduced pressure, extracted with EA, and washed. The residue was purified by column chromatography to give **compound 1** (531 mg, 64.4% yield).

LC-MS: $[M+H]^+ = 413.2$.

Example 2



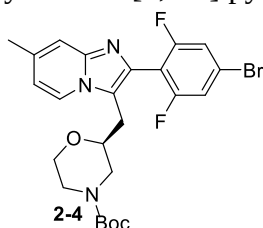
Step (1) Preparation of 2,6-difluoro-4-bromobenzaldehyde



3,5-difluorobromobenzene (10 g, 51.8 mmol) was dissolved in anhydrous THF (80 mL), and LDA (31 mL, 62.5 mmol) was added dropwise at -78°C under N_2 atmosphere. After the completion of the dropwise addition, the reaction system was stirred for 1 h, added with DMF

(4 mL, 51.9 mmol), and stirred for another 30 min. The reaction system was poured into NH₄Cl solution and extracted with ethyl acetate. The organic phase was washed with NaCl solution, dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **compound 2-2** (8 g, 70% yield) in the form of a yellow solid.

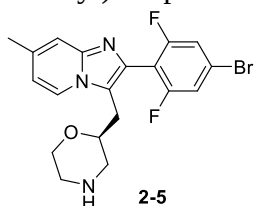
Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-bromophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)-methyl)morpholine-4-carboxylate



Compound 2-2 (200 mg, 0.9 mmol), 2-amino-4-methylpyridine (100 mg, 0.9 mmol), **compound 1-4** (191 mg, 0.9 mmol), CuCl (27 mg, 0.27 mmol), and Cu(CF₃SO₃)₂ (100 mg, 0.27 mmol) were dissolved in toluene (3 mL). Under N₂ atmosphere, the reaction system was heated to 85 °C, added with DMA (0.05 mL), and stirred for 5 h. Then the reaction system was cooled to room temperature, and incubated overnight. The resulting reaction system was poured into water (5 mL) and extracted with DCM (5 mL × 3). The organic phase was washed with saturated NaCl solution, dried over MgSO₄ filtered, and concentrated under reduced pressure. The crude product was purified by column chromatography (PE:EA = 1:1) to give **intermediate 2-4** (100 mg, 21% yield) in the form of a yellow oil. LC-MS: [M+H]⁺ = 522.3.

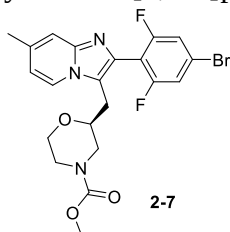
Step (3) Preparation of (S)-2-((2-(2,6-difluoro-4-bromophenyl)-7-methylimidazo[1,2-*a*]pyridin-

3-yl)-methyl)morpholine



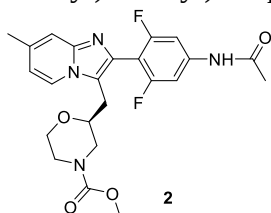
Intermediate 2-4 (100 mg) was dissolved in DCM (1 mL), and a solution of HCl in EtOH (0.5 mL) was added dropwise at 0 °C. After the completion of the dropwise addition, the reaction system was stirred for 2 h, and poured into NaHCO₃ solution to adjust the pH to 8. The aqueous phase was extracted with DCM. The organic phase was washed with NaCl solution, dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by column chromatography to give **intermediate 2-5** (60 mg, 75% yield) in the form of a yellow oil. LC-MS: [M+H]⁺ = 422.6.

Step (4) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-bromophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate



Intermediate 2-5 (320 mg, 0.76 mmol) and DIPEA (86 mg, 0.91 mmol) were dissolved in DCM (5 mL), and methyl chloroformate (**compound 2-6**) (86 mg, 0.91 mmol) was added. The reaction system was stirred at 0 °C for 3 h, poured into H₂O and extracted with DCM. The organic phase was washed with NaCl solution, dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under reduced pressure. The residue was purified by column chromatography to give **intermediate 2-7** (330 mg, 91% yield) in the form of a yellow oil. LC-MS: [M+H]⁺ = 480.6.

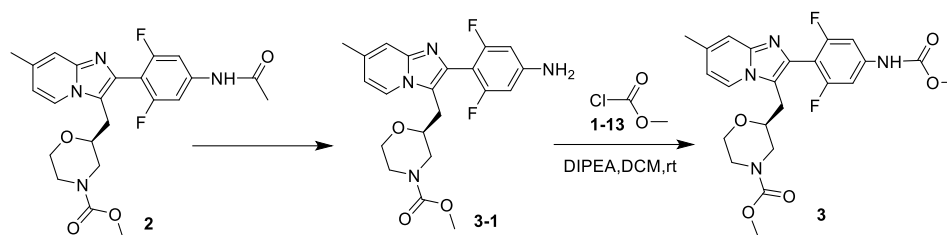
Step (5) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-acetamidophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate



To a solution of **intermediate 2-7** (50 mg, 0.10 mmol), acetamide (**compound 2-8**) (10 mg, 0.17 mmol) in 1,4-dioxane (2 mL) were added CS₂CO₃ (51 mg, 0.16 mmol), Xantphos (1.2 mg, 0.002 mmol) and Pd₂(dba)₃ (2 mg, 0.002 mmol) at room temperature. Under nitrogen atmosphere, the reaction system was heated to 80 °C and reacted overnight. After the reaction was completed, the reaction system was cooled to room temperature, filtered, and concentrated under reduced pressure. The residue was purified by preparative HPLC to give **compound 2** (18 mg, 38% yield) in the form of a white solid.

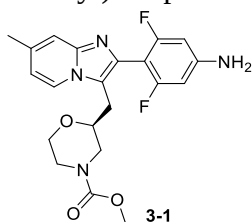
LC-MS: [M + H]⁺ = 459.7. ¹H NMR (400 MHz, MeOD): δ 8.41 (d, *J* = 7.1 Hz, 1H), 7.43 (d, *J* = 9.7 Hz, 2H), 7.35 (s, 1H), 6.88 (dd, *J* = 7.1, 1.5 Hz, 1H), 3.80 (d, *J* = 12.3 Hz, 3H), 3.67 (s, 3H), 3.59 (m, 1H), 3.43 – 3.34 (m, 1H), 3.15 – 3.01 (m, 2H), 2.91 (m, 1H), 2.63 (m, 1H), 2.47 (s, 3H), 2.19 (s, 3H).

Example 3



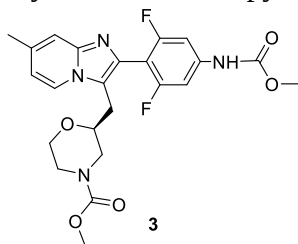
Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-aminophenyl)-7-methylimidazo[1,2-a]pyridin-

5 3-yl)-methyl)morpholine-4-carboxylate



To a solution of **compound 2** (145 mg, 0.32 mmol) in ethanol (3 mL) was added hydrochloric acid (1.5 mL) at room temperature. The reaction system was reacted at 100 °C for 1 h. After the reaction was completed, the reaction system was cooled to room temperature, added with NaHCO₃ solution (1 N) to quench the reaction, extracted with DCM (5 mL × 3), washed with saturated sodium chloride solution (5 mL × 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure to give a yellow oily liquid (90 mg). LC-MS: [M + H]⁺ = 417.3.

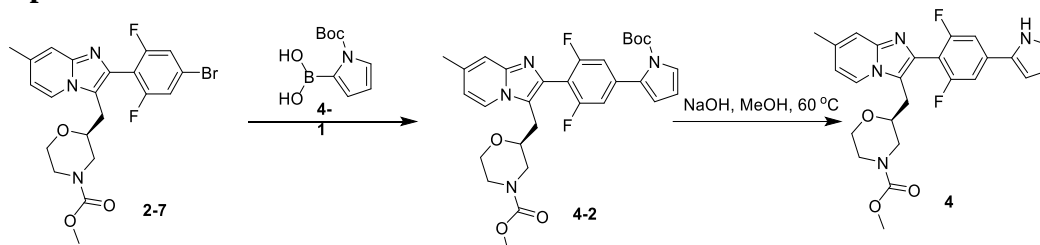
Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methoxycarbonylamino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate



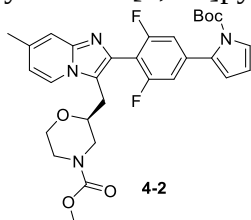
To a solution of **intermediate 3-1** (70 mg, 0.17 mmol) and DIPEA (65 mg, 0.5 mmol) in DCM (5 mL) was added methyl chloroformate (20 mg, 0.21 mmol) at 0 °C. The reaction system was reacted at this temperature for 3 h. After the reaction was completed, the reaction system was added with water (3 mL), extracted with DCM (3 mL × 3), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (DCM/MeOH, 40/1) to give **compound 3** (20 mg, 25% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 475.6$. ^1H NMR (400 MHz, MeOD): δ 8.42 (d, $J = 7.1$ Hz, 1H), 7.35 (s, 1H), 7.30 (d, $J = 9.9$ Hz, 2H), 6.90 (d, $J = 7.1$ Hz, 1H), 3.88 – 3.81 (m, 2H), 3.80 (s, 3H), 3.79 – 3.76 (m, 1H), 3.67 (s, 3H), 3.63 – 3.55 (m, 1H), 3.38 (t, $J = 10.8$ Hz, 1H), 3.15 – 3.00 (m, 2H), 2.90 (dt, $J = 20.6, 7.7$ Hz, 1H), 2.64 (t, $J = 11.5$ Hz, 1H), 2.47 (s, 3H).

5 Example 4

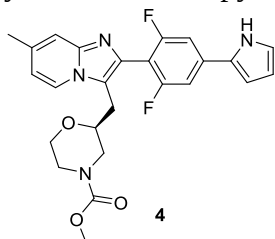


Step (1) Preparation of methyl (S)-2-((2-(4-(1-*tert*-butoxycarbonylpyrrol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)-methyl)morpholine-4-carboxylate



To a solution of **intermediate 2-7** (50 mg, 0.1 mmol), *N-tert*-butoxycarbonylpyrrolidine-2-boronic acid (**compound 4-1**) (26 mg, 0.12 mmol) and K_2CO_3 (36 mg, 0.26 mmol) in H_2O (0.3 mL) and 1,4-dioxane (2 mL) was added $\text{Pd}(\text{dppf})\text{Cl}_2$ (2 mg, 0.016 mmol) at room temperature. Under nitrogen atmosphere, the mixture was heated to 80 °C and reacted for 3 h. After the reaction was completed, the reaction mixture was cooled to room temperature, filtered, added with water (3 mL), extracted with ethyl acetate (5 mL $\times$ 3), washed with saturated sodium chloride solution (3 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 5/1) to give a yellow oily liquid (50 mg, 93% yield). LC-MS: $[M + H]^+ = 567.4$.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1H-pyrrol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



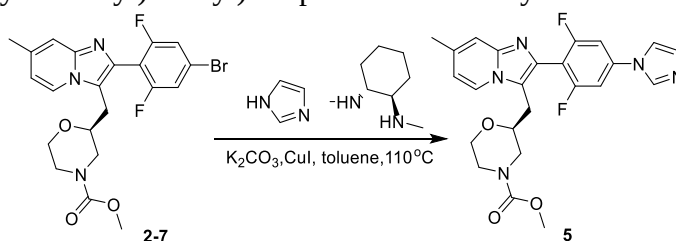
To a solution of **intermediate 4-2** (20 mg, 0.035 mmol) in methanol (1 mL) was added 1 M NaOH (4 mg, 0.1 mmol). The reaction mixture was reacted at 60 °C for 2 h. After the reaction

was completed, the reaction mixture was added with water (1 mL), extracted with DCM (1 mL $\times$ 3), washed with saturated sodium chloride solution (1 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by preparative HPLC to give **compound 4** (17.3 mg, 92.9% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 467.6$. ^1H NMR (400 MHz, MeOD): δ 8.41 (d, $J = 7.1$ Hz, 1H), 7.34 (d, $J = 9.2$ Hz, 3H), 6.91 (dd, $J = 2.6, 1.4$ Hz, 1H), 6.88 (dd, $J = 7.1, 1.5$ Hz, 1H), 6.69 (dd, $J = 3.6, 1.4$ Hz, 1H), 6.24 (dd, $J = 3.5, 2.7$ Hz, 1H), 3.81 (d, $J = 12.4$ Hz, 3H), 3.66 (s, 3H), 3.63 – 3.56 (m, 1H), 3.39 (t, $J = 10.6$ Hz, 1H), 3.17–3.04 (m, 2H), 2.96–2.85 (m, 1H), 2.71–2.59 (m, 1H), 2.47 (s, 3H).

Example 5

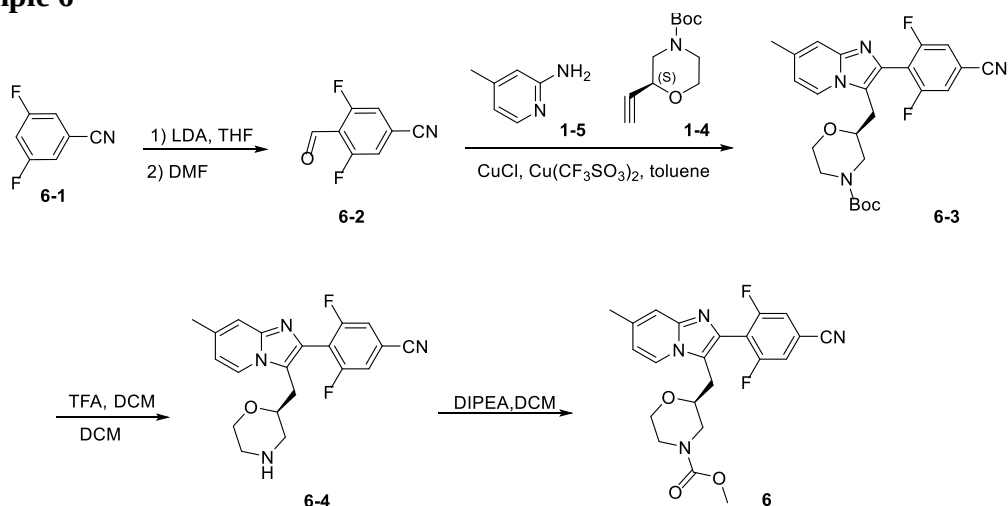
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1H-imidazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



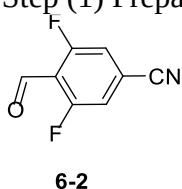
Intermediate 2-7 (50 mg, 0.1 mmol) was dissolved in toluene (2 mL), and imidazole (21 mg, 0.31 mmol), (1R,2R)-N1,N2-dimethylcyclohexyl-1,2-diamine (1.5 mg, 0.01 mmol) and K_2CO_3 (15 mg, 0.11 mmol) was added. Under N_2 atmosphere, the reaction system was added with CuI (2 mg, 0.01 mmol), and heated to 110 $^\circ\text{C}$ overnight. Then the reaction system was filtered, and the filtrate was purified to give **compound 5** (8.4 mg, 17.2% yield) in the form of a yellow solid.

LC-MS: $[M+H]^+ = 468.6$. ^1H NMR (400 MHz, MeOD) : δ 8.44(d, 1H), 8.42(s, 1H), 7.78(s, 1H), 7.56(d, 2H), 7.36(s, 1H), 7.22(s, 1H), 6.91(dd, 1H), 3.82(m, 3H), 3.67(s, 3H), 3.60(m, 1H), 3.38(m, 1H), 3.12(m, 2H), 2.90(m, 2H), 2.64(s, 3H).

Example 6

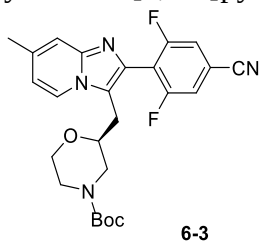


Step (1) Preparation of 2,6-difluoro-4-cyanobenzaldehyde



To a solution of **compound 6-1** (2 g, 14 mmol) in THF (20 mL) was added LDA (8.6 mL, 17.2 mmol) dropwise at -78 °C under nitrogen atmosphere. Then the reaction system was stirred at -78 °C for 1 h under nitrogen atmosphere, and added dropwise with DMF (1.26 g, 17.2 mmol) over 30 min. After the reaction was completed, the reaction system was added with AcOH (10%) to quench the reaction, extracted with ethyl acetate (50 mL × 3), washed with saturated sodium chloride solution (50 mL × 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 8/1) to give **compound 6-2** (1.5 g, 60% yield) in the form of a yellow solid.

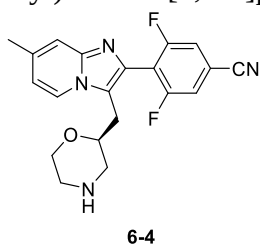
Step (2) Preparation of *tert*-butyl (S)-2-((2-(4-cyano-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



To a solution of **compound 6-2** (500 mg, 3 mmol), **compound 1-5** (330 mg, 3 mmol) and **compound 1-4** (630 mg, 3 mmol) in toluene (15 mL) were added CuCl (90 mg, 0.9 mmol) and Cu(CF₃SO₃)₂ (330 mg, 0.9 mmol) at room temperature. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with DME (0.1 mL), and reacted at this temperature for 5 h. After the reaction was completed, the reaction system was added with water (15 mL) to quench

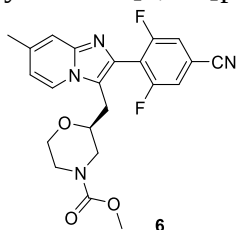
the reaction, extracted with dichloromethane (10 mL $\times$ 3), washed with saturated sodium chloride solution (10 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 1/1) to give **intermediate 6-3** (450 mg, 32% yield) in the form of a yellow oily liquid.

5 Step (3) Preparation of (S)-3,5-difluoro-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)benzonitrile



To a solution of **intermediate 6-3** (400 mg, 1.1 mmol) in DCM (6 mL) was added TFA (2 mL) dropwise at 0 °C. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added with saturated NaHCO₃ to quench the reaction at 0 °C, extracted with dichloromethane (3 mL $\times$ 3), washed with saturated sodium chloride solution (3 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (DCM/MeOH, 40/1) to give **intermediate 6** (280 mg, 79% yield) in the form of a brown solid.

15 Step (4) Preparation of methyl (S)-2-((2-(4-cyano-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

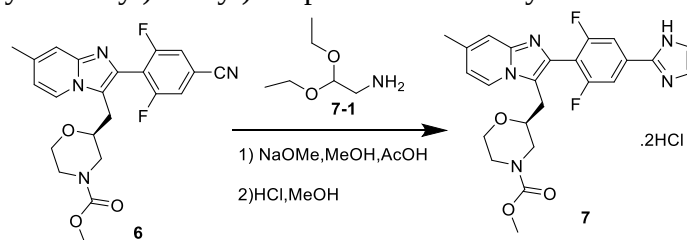


To a solution of **intermediate 6-4** (230 mg, 0.6 mmol) and DIPEA (230 mg, 1.8 mmol) in DCM (2 mL) was added methyl chloroformate (66 mg, 0.72 mmol) at 0 °C. The reaction system was reacted for 1 h. After the reaction was completed, the reaction system was added with water (3 mL), extracted with dichloromethane (3 mL $\times$ 3), washed with saturated sodium chloride solution (3 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 2/1) to give **compound 6** (230 mg, 83% yield) in the form of a yellow solid.

25 LC-MS: [M + H]⁺ = 427.7. ¹H NMR (400 MHz, CDCl₃): δ 8.19 (d, *J* = 7.1 Hz, 1H), 7.41 (s, 1H), 7.37 (d, *J* = 6.2 Hz, 2H), 6.72 (d, *J* = 7.0 Hz, 1H), 4.03 – 3.74 (m, 3H), 3.71 (s, 3H), 3.61 – 3.51 (m, 1H), 3.38 (t, *J* = 11.8 Hz, 1H), 3.09 – 2.82 (m, 3H), 2.68 - 2.53 (m, 1H), 2.44 (s, 3H).

Example 7

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1*H*-imidazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

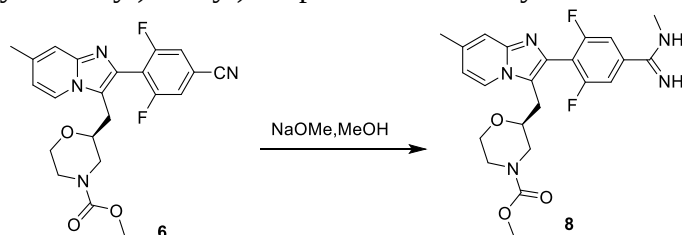


A solution of **compound 6** (120 mg, 0.3 mmol) and NaOMe (3 mg, 0.05 mmol) in MeOH (2 mL) was stirred at room temperature for 15 min, and aminoacetaldehyde diethyl acetal (49 mg, 0.4 mmol) and AcOH (34 mg, 0.6 mmol) were added. The reaction system was reacted at 50 °C for 1 h, then cooled to room temperature, added with HCl (0.5 mL) and MeOH (2 mL), and reacted at 65 °C for 18 h. After the reaction was completed, the reaction system was concentrated under reduced pressure. The crude product was purified by preparative HPLC, added with 1 M hydrochloric acid, and lyophilized to give **compound 7** (32 mg, 24% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 468.1$. ^1H NMR (400 MHz, CDCl_3): δ 16.00 (s, 2H), 15.16 (s, 1H), 8.68 (s, 3H), 8.00 (s, 1H), 7.49 (s, 2H), 7.22 (s, 1H), 3.94 (s, 3H), 3.70 (m, 3H), 3.61 (m, 1H), 3.34 (m, 1H), 2.97 (m, 4H), 2.62 (s, 3H).

Example 8

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(*N*-methylcarbamimidoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

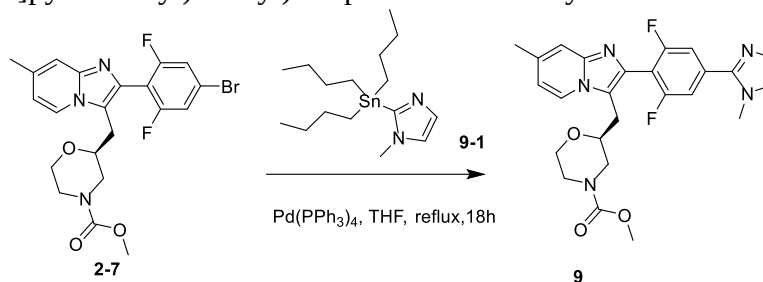


A solution of **compound 6** (50 mg, 0.12 mmol) and NaOMe (2 mg, 0.03 mmol) in MeOH (2 mL) was stirred at room temperature for 30 min, then a solution of MeNH_2/THF (2 M, 0.1 mL, 0.15 mmol) and AcOH (14 mg, 0.24 mmol) were added at 50 °C. The reaction system was reacted for 1 h. After the reaction was completed, the reaction system was concentrated under reduced pressure. The residue was purified by silica gel column chromatography to give **compound 8** (17 mg, 31% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 458.1$. ^1H NMR (400 MHz, CDCl_3): δ 8.54 (s, 1H), 8.25 (s, 1H), 7.57 (d, $J = 6.8$ Hz, 2H), 7.39 (s, 1H), 6.74 (d, $J = 6.8$ Hz, 1H), 3.82 (m, 3H), 3.68 (s, 3H), 3.56 (m, 1H), 3.36 (m, 1H), 3.22 (s, 2H), 3.07 – 2.75 (m, 4H), 2.61 (m, 1H), 2.45 (s, 3H).

Example 9

- 5 Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1-methyl-1*H*-imidazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

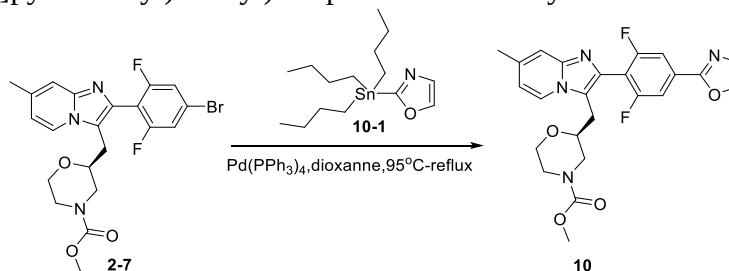


- 10 **Intermediate 2-7** (50 mg, 0.1 mmol) was dissolved in toluene (3 mL), and the reaction system was added with $\text{Pd}(\text{PPh}_3)_4$ (12 mg, 0.01 mmol) and 1-methyl-2-(tri-*n*-butylstannyl)imidazole (78 mg, 0.2 mmol). Under N_2 atmosphere, the reaction system was heated to 85 °C and incubated at this temperature for 5 h. Then the reaction system was filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **compound 9** (24 mg, 49% yield) in the form of a white solid.

- 15 LC-MS: $[M+H]^+ = 482.2$. ^1H NMR (400 MHz, MeOD): δ 8.69(s, 1H), 8.55(d, 1H), 7.99(s, 1H), 7.55(s, 1H), 7.22(d, 2H), 7.15(d, 1H), 4.06(s, 1H), 3.96(s, 3H), 3.86(m, 2H), 3.73(s, 3H), 3.68(s, 1H), 3.43(m, 1H), 3.09(d, 2H), 2.99(m, 1H), 2.73(m, 1H), 2.61(s, 3H).

Example 10

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(oxazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

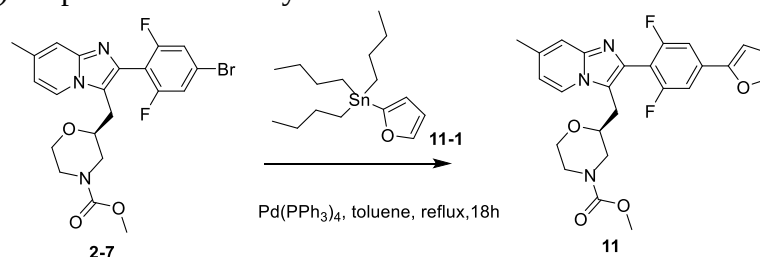


- 20 To a solution of **intermediate 2-7** (50 mg, 0.1 mmol) in 1,4-dioxane (2 mL) were added $\text{Pd}(\text{Ph}_3\text{P})_4$ (12 mg, 0.01 mmol) and 2-(tri-*n*-butylstannyl)oxazole (119 mg, 0.33 mmol) at room temperature. Under nitrogen atmosphere, the reaction system was heated to 95 °C and reacted for 3 h. After the reaction was completed, the reaction solution was cooled to room temperature, 25 filtered, and concentrated under reduced pressure. The residue was purified by preparative HPLC to give **compound 10** (12 mg, 25% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 469.7$. ^1H NMR (400 MHz, MeOD): δ 8.43 (d, $J = 7.1$ Hz, 1H), 8.10 (d, $J = 0.8$ Hz, 1H), 7.80 (d, $J = 8.0$ Hz, 2H), 7.42 (d, $J = 0.8$ Hz, 1H), 7.36 (s, 1H), 6.89 (dd, $J = 7.2, 1.6$ Hz, 1H), 3.93–3.74 (m, 3H), 3.66 (s, 3H), 3.64–3.56 (m, 1H), 3.42–3.34 (m, 1H), 3.13 (d, $J = 6.3$ Hz, 2H), 2.97–2.85 (m, 1H), 2.73–2.60 (m, 1H), 2.48 (s, 3H).

5 Example 11

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(furan-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

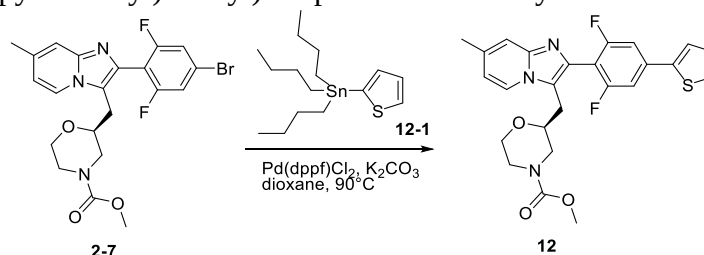


Intermediate 2-7 (50 mg, 0.1 mmol) was dissolved in toluene (3 mL), and the reaction system was added with $\text{Pd(PPh}_3)_4$ (12 mg, 0.01 mmol) and 2-(tri-*n*-butylstannyl)furan (37 mg, 0.1 mmol). Under N_2 atmosphere, the reaction system was heated to 110 °C and incubated at this temperature for 18 h. Then the reaction system was filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **compound 11** (12 mg, 24% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 468.1$. ^1H NMR (400 MHz, CDCl_3): δ 8.55 (d, 1H), 8.08 (s, 1H), 7.58(d, 1H), 7.40(d, 2H), 7.14(d, 1H), 6.86(d, 1H), 6.58(s, 1H), 3.82(m, 3H), 3.73(s, 3H), 3.68(m, 1H), 3.43(m, 1H), 3.05(m, 2H), 2.96(m, 1H), 2.71(m, 1H), 2.61(s, 3H).

Example 12

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(thiophen-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

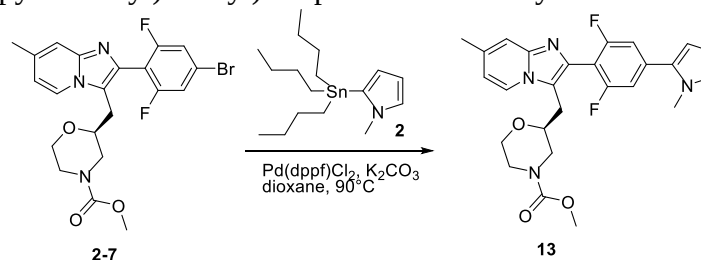


Intermediate 2-7 (50 mg, 0.1 mmol) was dissolved in dioxane (3 mL), and the reaction system was added with $\text{Pd(PPh}_3)_4$ (12 mg, 0.01 mmol) and 2-(tri-*n*-butylstannyl)thiophene (117 mg, 0.31 mmol). Under N_2 atmosphere, the reaction system was heated to 100 °C and incubated at this temperature for 8 h. Then the reaction system was filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **compound 12** (18 mg, 35% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 484.6$. $^1\text{H NMR}$ (400 MHz, CDCl_3): δ 8.22 (d, 1H), 7.41(m, 3H), 7.31(m, 2H), 7.14(dd, 1H), 6.71(d, 1H), 3.82(m, 3H), 3.68(s, 3H), 3.62(m, 1H), 3.41(m, 1H), 3.03(m, 2H), 2.92(m, 1H), 2.64(m, 1H), 2.46(s, 3H).

Example 13

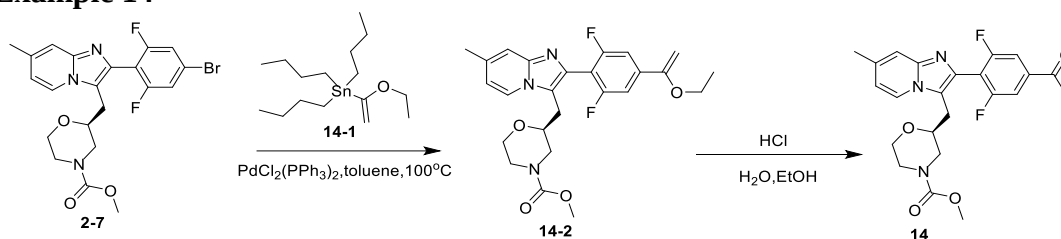
- 5 Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1-methyl-1H-pyrrol-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



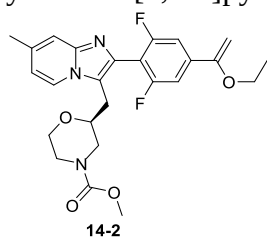
- Intermediate 2-7** (50 mg, 0.1 mmol) was dissolved in dioxane (3 mL), and the reaction system was added with $\text{Pd(PPh}_3)_4$ (12 mg, 0.01 mmol) and 2-(tri-*n*-butylstannyl)methylpyrrole (115 mg, 0.31 mmol). Under N_2 atmosphere, the reaction system was heated to 100 °C and incubated at this temperature for 8 h. Then the reaction system was filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **compound 13** (28 mg, 54% yield) in the form of a white solid.

- LC-MS: $[M+H]^+ = 481.4$. $^1\text{H NMR}$ (400 MHz, CD_3OD): δ 8.86 (d, 1H), 7.72(s, 1H), 7.42(dd, 1H), 7.39(d, 2H), 6.91(t, 1H), 6.49(dd, 1H), 6.20(dd, 1H), 4.02(d, 1H), 3.84(s, 3H) 3.75(m, 3H), 3.69(s, 3H), 3.38(m, 1H), 3.27(m, 2H), 2.96(m, 1H), 2.79(m, 1H), 2.66(s, 3H).

Example 14



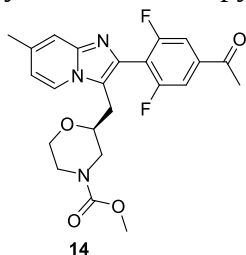
- Step (1) Preparation of methyl (S)-2-((2-(4-(1-ethoxyethenyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 2-7 (300 mg, 0.62 mmol) was dissolved in toluene (3 mL), and the reaction system was added with $\text{PdCl}_2(\text{PPh}_3)_2$ (44 mg, 0.062 mmol), tri-*tert*-butyl(1-ethoxyethenyl)tin

(680 mg, 1.88 mmol). Under N₂ atmosphere, the reaction system was heated to 95 °C and incubated at this temperature for 3 h. Then the reaction system was filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified by column chromatography to give **intermediate 14-2** (215 mg, 73% yield) in the form of a yellow oil. LC-MS: [M+H]⁺ = 472.7.

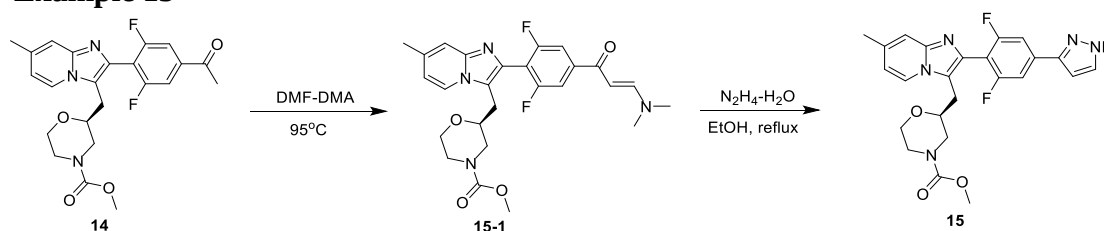
Step (2) Preparation of methyl (S)-2-((2-(4-acetyl-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



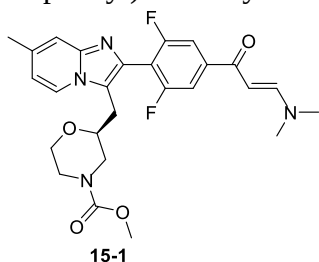
Intermediate 14-2 (250 mg, 0.53 mmol) was dissolved in EtOH (3 mL), and HCl (2 N, 1.7 mL) was added. The reaction system was stirred at room temperature for 2 h. After the reaction was completed, the reaction system was poured into NaHCO₃ solution to adjust the pH to 8. The aqueous phase was extracted with DCM. The organic phase was washed with NaCl solution 2-3 times, followed by liquid separation. The organic phase was dried over anhydrous magnesium sulfate and filtered, and the filtrate was concentrated under reduced pressure. The crude product was purified to give **compound 14** (25 mg, 38% yield) in the form of a white solid.

LC-MS: [M+H]⁺ = 444.7. ¹H NMR (400 MHz, MeOD): δ 8.43(d, 1H), 7.76(d, 2H), 7.36(s, 1H), 6.90(dd, 1H), 3.79(m, 3H), 3.67(s, 3H), 3.59(m, 1H), 3.36(m, 1H), 3.11(m, 2H), 2.86(m, 1H), 2.68(s, 3H), 2.63(m, 1H), 2.47(s, 3H).

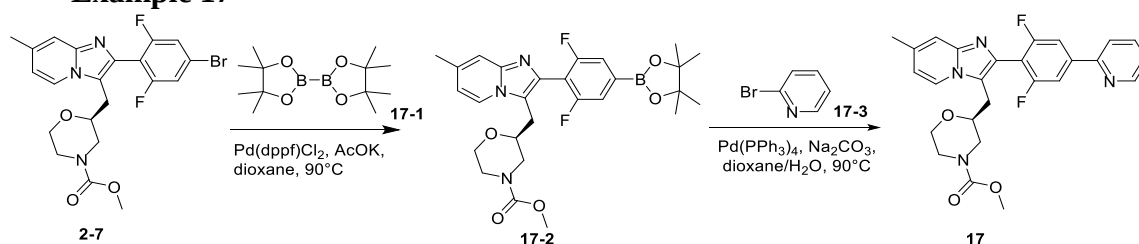
Example 15



Step (1) Preparation of methyl (S,E)-2-((2-(4-(3-(dimethylamino)acryloyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate

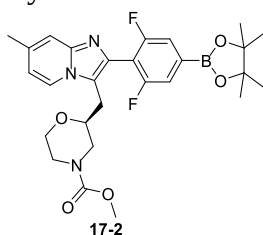


Example 17



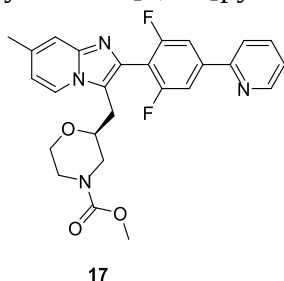
Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-

5 carboxylate



Intermediate 2-7 (100 mg, 0.21 mmol), boronate (64 mg, 0.25 mmol), Pd(dppf)Cl₂ (8 mg, 0.01 mmol) and AcOK (6.2 mg, 0.625 mmol) were dissolved in dioxane (3 mL). Under N₂ atmosphere, the reaction system was heated to 90 °C and reacted overnight. Then the reaction system was cooled to room temperature to give **intermediate 17-2**. LC-MS: [M+H]⁺ = 528.2.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(pyridin-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

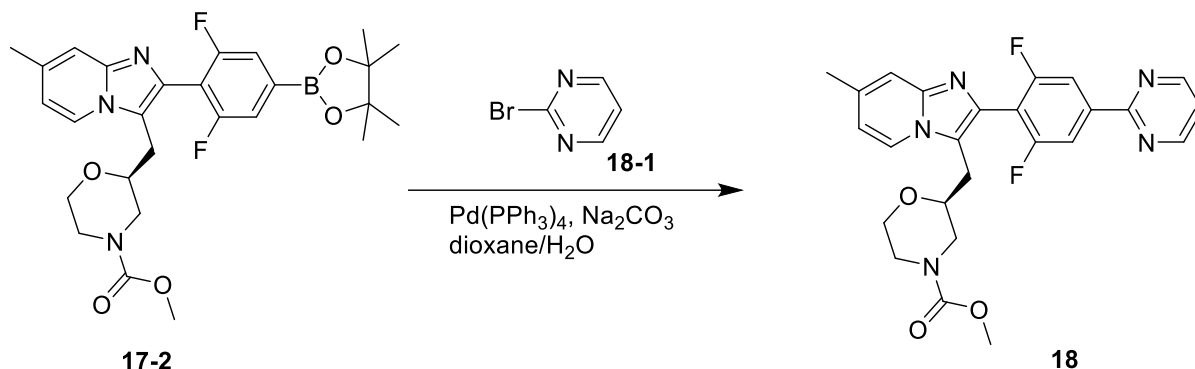


Intermediate 17-2 (55 mg, 0.1 mmol), 2-bromopyridine (18 mg, 0.11 mmol), Na₂CO₃ (33 mg, 0.31 mmol), Pd(PPh₃)₄ (6 mg, 0.005 mmol) were dissolved in dioxane (3 mL) and a small amount of water. Under N₂ atmosphere, the reaction system was heated to 100 °C. After the reaction was completed, the reaction system was filtered, and the filtrate was diluted with water and extracted with EA. The organic phase was washed with NaCl, dried, filtered, and concentrated under reduced pressure. The residue was purified to give **compound 17** (25 mg, 50% yield) in the form of a white solid.

LC-MS: [M+H]⁺ = 479.7. ¹H NMR (400 MHz, CDCl₃): δ 8.75 (d, 1H), 8.27 (d, 1H), 7.85(t, 1H), 7.76(dd, 3H), 7.56(m, 1H), 7.35(m, 1H), 6.76(d, 1H), 3.82(s, 3H), 3.68(s, 3H), 3.62(s, 1H), 3.41(s, 1H), 3.03(dd, 2H), 2.92(s, 1H), 2.64(dd, 1H), 2.47(s, 3H).

Example 18

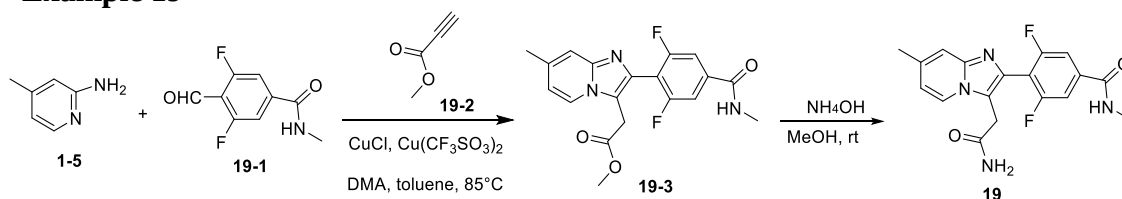
Preparation of methyl (S)-2-((2-(3-fluoro-4-(methylcarbamoyl)-1H-pyrrol-2-yl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



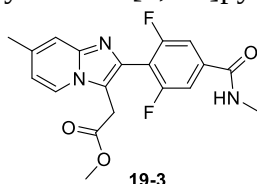
Intermediate 17-2 (55 mg, 0.1 mmol), 2-bromopyrimidine (19 mg, 0.11 mmol), Na₂CO₃ (27 mg, 0.31 mmol), Pd(PPh₃)₄ (4 mg, 0.005 mmol) were dissolved in dioxane (3 mL) and a small amount of water. Under N₂ atmosphere, the reaction system was heated to 90 °C. After the reaction was completed, the reaction system was filtered, and the filtrate was diluted with water and extracted with EA. The organic phase was washed with NaCl, dried, filtered, and concentrated under reduced pressure. The residue was purified to give **compound 18** (25 mg, 50% yield) in the form of a white solid.

LC-MS: [M+H]⁺ = 480.2. ¹HNMR (400 MHz, CDCl₃): δ 8.87 (d, 2H), 8.24 (t, 1H), 8.18(d, 2H), 7.54(s, 1H), 7.30(m, 1H), 6.76(d, 1H), 3.82(s, 3H), 3.72-3.77(m, 3H), 3.68(s, 3H), 3.62(s, 1H), 3.41(s, 1H), 3.03(t, 2H), 2.92(s, 1H), 2.64(dd, 1H), 2.46(s, 3H).

Example 19



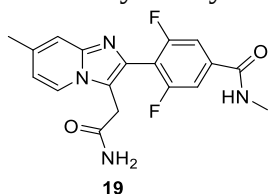
Step (1) Preparation of methyl 2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)acetate



3,5-difluoro-4-formyl-N-methylbenzamide (200 mg, 1 mmol), 2-amino-4-methylpyridine (108 mg, 1 mmol), methyl propiolate (84 mg, 1 mmol), CuCl (30 mg, 0.3 mmol), and Cu(CF₃SO₃)₂ (109 mg, 0.3 mmol) were dissolved in toluene (3 mL). Under N₂ atmosphere, the

reaction system was heated to 85 °C, added with DMA (0.05 mL), and stirred for 5 h. Then the reaction system was cooled to room temperature, and incubated overnight. The resulting reaction system was poured into H₂O (5 mL) and extracted with DCM (5 mL × 3). The organic phase was washed with saturated NaCl, dried over MgSO₄, filtered, and concentrated under reduced pressure. The crude product was purified by column chromatography (PE:EA = 1:1) to give **intermediate 19-3** (172 mg, 25% yield) in the form of a yellow oil. LC-MS: [M+H]⁺ = 374.4.

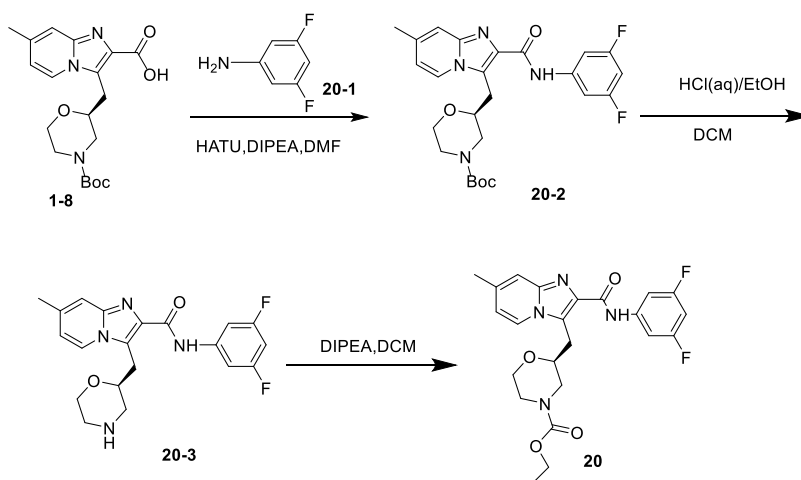
Step (2) Preparation of 4-(3-(2-amino-2-oxoethyl)-7-methylimidazo[1,2-*a*]pyridine)-3,5-difluorobenzoylmethylamine



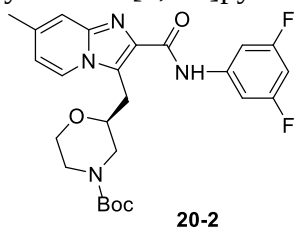
Intermediate 19-3 (82 mg, 0.22 mmol) was dissolved in methanol (2 mL). The reaction system was added with NH₃·H₂O (1 mL) and stirred overnight. Then the reaction system was concentrated under reduced pressure. The crude product was purified to give **compound 19** (25 mg, 32% yield) in the form of a white solid.

LC-MS: [M+H]⁺ = 359.8. ¹HNMR (400 MHz, CD₃OD): δ 8.36 (d, 1H), 7.63(d, 2H), 7.46(s, 1H), 7.02(d, 1H), 3.93(s, 2H), 2.97(s, 3H), 2.52(s, 3H).

Example 20

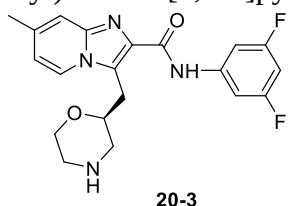


Step (1) Preparation of *tert*-butyl (S)-2-((2-(3,5-difluorophenyl)carbamoyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)-methyl)morpholine-4-carboxylate



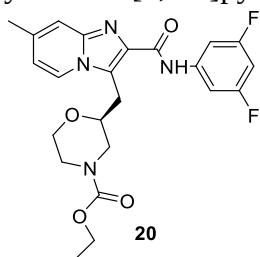
A solution of **intermediate 1-8** (400 mg, 1.07 mmol) and HATU (608 mg, 1.6 mmol) in DMF (5 mL) was stirred at room temperature for 10 min, and DIPEA (418 mg, 3.24 mmol) and **compound 20-1** (168 mg, 1.3 mmol) were added at room temperature. The reaction system was reacted for 1 h. After the reaction was completed, the reaction system was added with water (15 mL) to quench the reaction, extracted with dichloromethane (10 mL $\times$ 3), washed with saturated sodium chloride solution (10 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (DCM/MeOH, 10/1) to give **intermediate 20-2** (450 mg) in the form of a yellow oily liquid.

Step (2) Preparation of (S)-N-(3,5-difluorophenyl)-7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-a]pyridine-2-carboxamide



To a solution of **intermediate 20-2** (330 mg, 0.7 mmol) in DCM (4 mL) was added a solution of HCl in ethanol (1 mL) dropwise at 0 °C. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added with saturated NaHCO₃ to quench the reaction at 0 °C, extracted with dichloromethane (3 mL $\times$ 3), washed with saturated sodium chloride solution (3 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (DCM/MeOH, 10/1) to give **intermediate 20-3** (80 mg, 31% yield) in the form of a brown solid.

Step (3) Preparation of ethyl (S)-2-((2-(3,5-difluorophenyl)carbamoyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-acetate

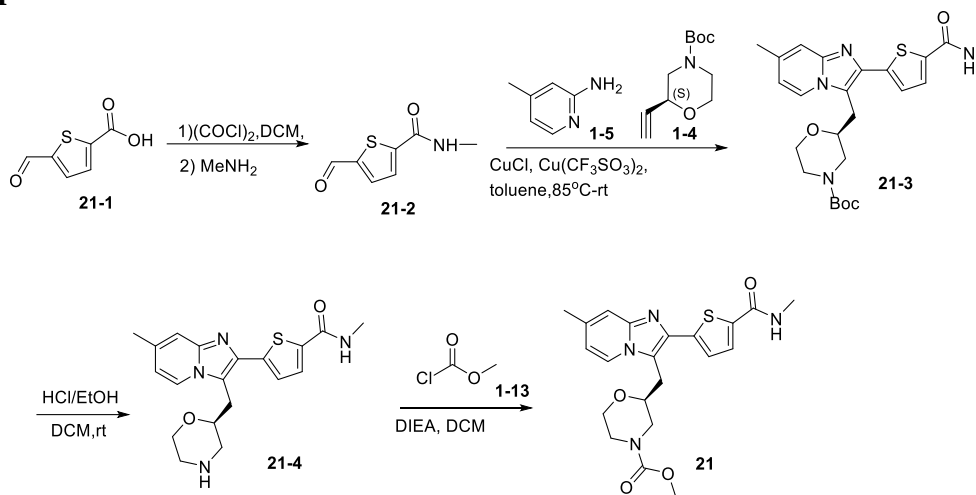


To a solution of **intermediate 20-3** (80 mg, 0.2 mmol) and DIPEA (25 mg, 0.6 mmol) in DCM (2 mL) was added methyl chloroformate (1 mL) at 0 °C. The reaction system was reacted for 1 h. After the reaction was completed, the reaction system was added with water (3 mL), extracted with dichloromethane (3 mL $\times$ 3), washed with saturated sodium chloride solution (3 mL $\times$ 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The

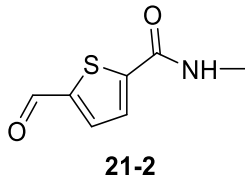
residue was purified by preparative HPLC to give **compound 20** (15 mg, 16% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 459.7$. ^1H NMR (400 MHz, $CDCl_3$): δ 8.30 (s, 1H), 7.52 (s, 3H), 6.90 (s, 1H), 6.59 (t, $J = 8.7$ Hz, 1H), 4.23 (d, $J = 9.8$ Hz, 1H), 4.16 (dd, $J = 14.1, 7.1$ Hz, 2H), 3.96 - 3.79 (m, 3H), 3.75 (d, $J = 12.2$ Hz, 1H), 3.41 (dd, $J = 11.9, 9.2$ Hz, 1H), 3.35 - 3.19 (m, 1H), 2.99 - 2.85 (m, 1H), 2.84 - 2.73 (m, 1H), 2.50 (s, 3H), 1.35 - 1.18 (m, 3H).

Example 21

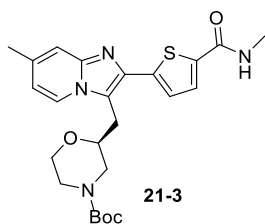


Step (1) Preparation of 5-formylthiophene-2-carboxamide



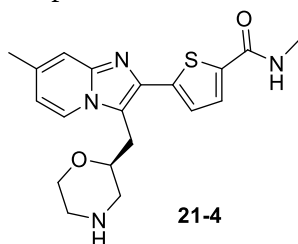
To a solution of 5-formylthiophene-2-carboxylic acid (500 mg, 3.2 mmol) and DMF (12 mg) was added oxalyl chloride (447 mg, 1.1 mmol) at 0 °C. The reaction system was reacted at this temperature for 2 h, and added with aqueous methylamine solution (40%, 746 mg, 9.6 mmol) over 10 min. After the reaction was completed, the reaction system was added with water (3 mL) to quench the reaction, extracted with dichloromethane (5 mL $\times$ 3), washed with saturated sodium chloride solution (5 mL $\times$ 2), dried over anhydrous sodium sulfate and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (PE/EA, 1/1) to give **compound 21-2** in the form of a white solid (300 mg, 56% yield), LC-MS: $[M+H]^+ = 170.9$.

Step (2) Preparation of *tert*-butyl (S)-2-((7-methyl-2-(5-(methylcarbamoyl)thiophen-2-yl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



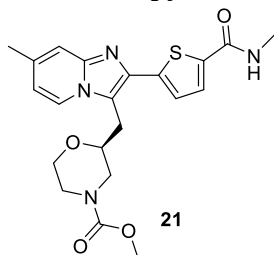
To a solution of **compound 21-2** (500 mg, 2.96 mmol), 4-methylpyridin-2-amine (320 mg, 2.96 mmol) and **compound 1-4** (625 mg, 2.96 mmol) in toluene (5 mL) were added copper(I) chloride (88 mg, 0.89 mmol) and copper(II) trifluoromethanesulfonate (321 mg, 0.89 mmol) at room temperature. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with DMA (0.05 mL), and reacted at this temperature for 5 h. After the reaction was completed, the reaction system was added with water (5 mL) to quench the reaction, extracted with dichloromethane (5 mL × 3), washed with saturated sodium chloride solution (5 mL × 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (DCM/MeOH, 40/1) to give **intermediate 21-3** (300 mg, 56% yield) in the form of a yellow oily liquid, LC-MS: $[M+H]^+ = 471.7$.

Step (3) Preparation of (S)-5-(7-methyl-3-(morpholin-2-yl-methyl)imidazo[1,2-a]pyridin-2-yl)thiophene-2-carboxamide



To a solution of **intermediate 21-3** (400 mg, 0.85 mmol) in DCM (4 mL) was added a solution of HCl in ethanol (0.5 mL, 33%) dropwise at 0 °C. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added with saturated NaHCO_3 to quench the reaction at 0 °C, extracted with dichloromethane (5 mL × 3), washed with saturated sodium chloride solution (5 mL × 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure to give **intermediate 21-4** (300 mg) in the form of a yellow oily liquid, LC-MS: $[M+H]^+ = 371.8$.

Step (4) Preparation of methyl (S)-2-((7-methyl-2-(5-(methylcarbamoyl)thiophen-2-yl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

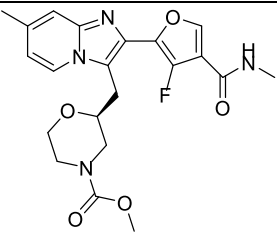
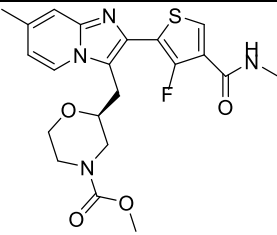
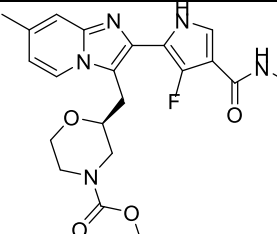


To a solution of **intermediate 21-4** (100 mg, 0.27 mmol) and DIPEA (105 mg, 0.81 mmol) in DCM (2 mL) was added methyl chloroformate (31 mg, 0.33 mol) at 0 °C. The reaction system was reacted for 3 h. After the reaction was completed, the reaction system was added with water (3 mL), extracted with dichloromethane (3 mL × 2), washed with saturated sodium chloride solution (3 mL × 2), dried over anhydrous sodium sulfate, and concentrated under reduced pressure. The residue was purified by preparative HPLC to give **compound 21** (28.5 mg, 28% yield) in the form of a yellow solid.

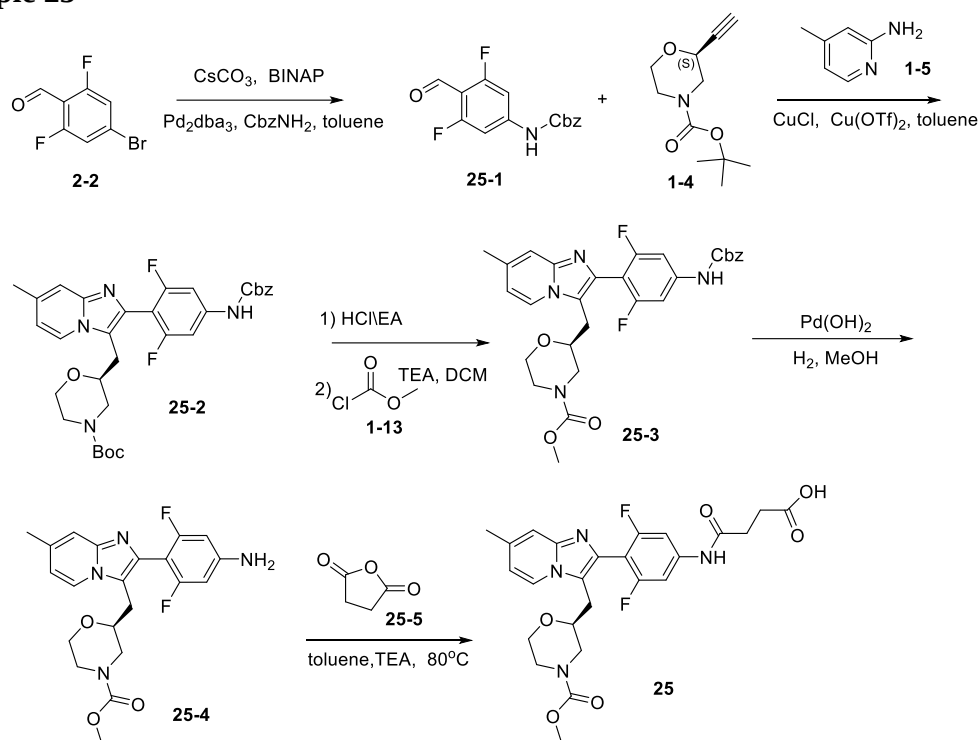
LC-MS: $[M + H]^+ = 429.7$. ^1H NMR (400 MHz, MeOD): δ 8.32 (d, $J = 7.0$ Hz, 1H), 7.67 (d, $J = 3.9$ Hz, 1H), 7.52 (d, $J = 3.8$ Hz, 1H), 7.31 (s, 1H), 6.88- 6.80 (m, 1H), 4.13 (d, $J = 12.8$ Hz, 1H), 3.92 - 3.75 (m, 3H), 3.72 (s, 3H), 3.46 - 3.28 (m, 3H), 3.01 (m, 1H), 2.94 (s, 3H), 2.91 (m, 1H), 2.44 (s, 3H).

Examples 22-24

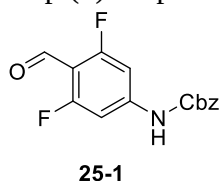
The compounds were prepared using the corresponding starting materials according to the method as described in Example 21.

Number	Compound structure	LC-MC (RT, m/z)
Example 22	 22	431.4
Example 23	 23	447.1
Example 24	 24	430.1

Example 25

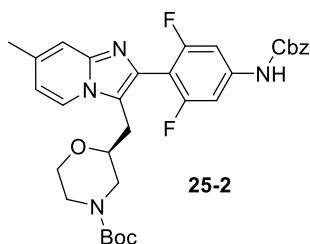


Step (1) Preparation of benzyl (3,5-difluoro-4-formylphenyl)-carbamate



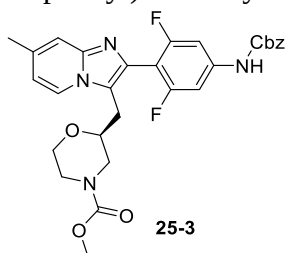
- 5 To a solution of **compound 2-2** (1 g, 4.5 mmol) in toluene (20 mL) were added cesium carbonate (2 g, 6.3 mmol), benzyl carbamate (0.8 g, 5.4 mmol), ($\pm$)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (0.28 g, 0.45 mmol) and tris(dibenzylideneacetone)dipalladium (0.2 g, 0.2 mmol) at room temperature. After the addition, the reaction system was purged with nitrogen three times, heated to 100°C and stirred overnight
- 10 (16 h) under nitrogen atmosphere. Then the reaction system was filtered and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (EA/PE = 1/10) to give **intermediate 25-1** (0.72 g, 55.2% yield) in the form of a yellow solid, LC-MS: $[\text{M}+\text{H}]^+ = 292.1$.

- 15 Step (2) Preparation of *tert*-butyl (S)-2-((2-(4-(((benzyloxy)-carbonyl)amino)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



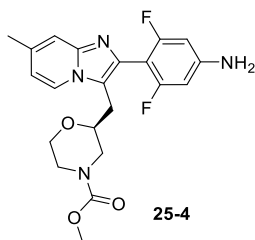
To a 20 mL microwave tube (Biotage) were added **compound 25-1** (0.2 g, 0.68 mmol), **compound 1-4** (0.22 g, 1.03 mmol) and **compound 1-5** (74 mg, 0.68 mmol), copper(I) chloride (20 mg, 0.2 mmol), copper(II) trifluoromethanesulfonate (74 mg, 0.2 mmol) and toluene (6 mL). After the addition, the reaction system was purged with nitrogen for 2 min, and reacted at 120 °C overnight (16 h) after the tube was sealed. A new spot was generated by developing the TLC plate (PE/EA = 1/1) with R_f value being about 0.3, and was identified as the target product by liquid chromatography. Then the reaction system was cooled to room temperature and concentrated under reduced pressure. The residue was mixed with silica gel and purified by silica gel column chromatography (EA/PE = 1/1) to give **intermediate 25-2** (184 mg, 45.5% yield) in the form of a yellow solid. LC-MS: [M+H]⁺ = 593.6.

Step (3) Preparation of methyl (S)-2-((2-(4-(((benzyloxy)-carbonyl)amino)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



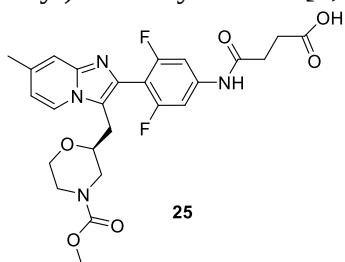
To **intermediate 25-2** (0.8 g, 1.35 mmol) was added a solution of HCl in ethyl acetate (about 4 M, 5 mL) at room temperature. After the addition, the reaction system was stirred at room temperature for 2 h. After the reaction was completed as monitored by liquid chromatography, the reaction system was concentrated under reduced pressure to give a crude product, to which were added dichloromethane (5 mL), triethylamine (0.4 g, 4 mmol) and methyl chloroformate (0.19 g, 2 mmol). After the addition, the reaction system was stirred at room temperature for 3 h. After the reaction was completed as monitored by liquid chromatography, the reaction system was concentrated under reduced pressure. The residue was purified by silica gel column chromatography (EA) to give **intermediate 25-3** (675 mg, 90.8% yield) in the form of a brown oil. LC-MS: [M+H]⁺ = 551.2.

Step (4) Preparation of methyl (S)-2-((2-(4-amino-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



To a solution of **intermediate 25-3** (200 mg, 0.363 mmol) in methanol (10 mL) was added palladium hydroxide (about 40 mg, 20% w/w) at room temperature. After the addition, the reaction system was purged with hydrogen balloon 3 times, and stirred at room temperature for 4 h under hydrogen atmosphere. After the reaction was completed as monitored by liquid chromatography, the reaction system was filtered and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (EA) to give **intermediate 25-4** (111 mg, 74% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 417.2$.

Step (5) Preparation of (S)-4-((3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)-phenyl)-amino)-4-oxobutanoic acid

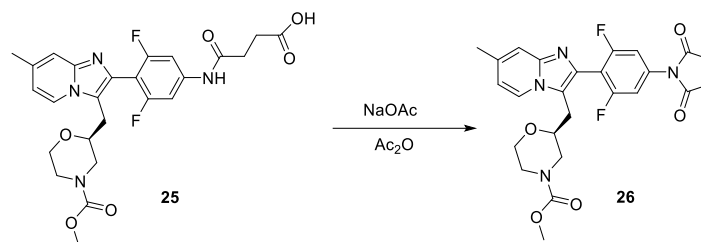


To a solution of **intermediate 25-4** (100 mg, 0.24 mmol) in toluene (5 mL) were added triethylamine (24 mg, 0.24 mmol) and succinic anhydride (28.8 mg, 0.288 mmol) at room temperature. After the addition, the reaction system was heated to 80 °C and reacted for 4 h. After the reaction was completed as monitored by liquid chromatography, the reaction system was concentrated under reduced pressure. The residue was purified by preparative high performance liquid chromatography (ammonium bicarbonate method), and lyophilized to give **compound 25** (90 mg, 72.5% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 517.1$. ^1H NMR (400 MHz, CD_3OD): δ 8.39 (d, $J = 7.2$ Hz, 1H), 7.46-7.38 (m, 2H), 7.33 (s, 1H), 6.87-6.85 (m, 1H), 3.85-3.58 (m, 3H), 3.67 (s, 3H), 3.60-3.56 (m, 1H), 3.39-3.33 (m, 1H), 3.13-3.02 (m, 2H), 2.93-2.87 (m, 1H), 2.74-2.65 (m, 4H), 2.64-2.60 (m, 1H), 2.46 (s, 3H).

Example 26

Preparation of methyl (S)-2-((2-(4-(2,5-dioxopyrrolidin-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

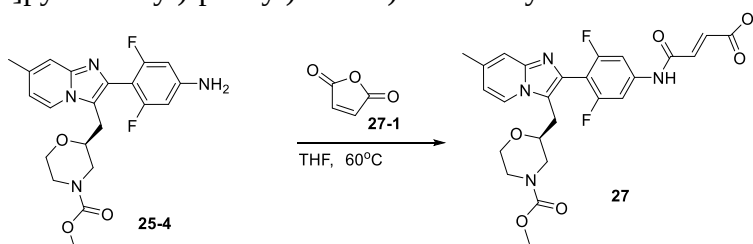


To a solution of **compound 25** (77.6 mg, 0.15 mmol) in acetic anhydride (2 mL) was added sodium acetate (9.2 mg, 0.11 mmol) at room temperature. The reaction system was stirred at room temperature for 4 h. Then the reaction system was concentrated under reduced pressure. The residue was purified by preparative high performance liquid chromatography (trifluoroacetic acid method), and lyophilized to give **compound 26** (30 mg, 40% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 499.1$. ^1H NMR (400 MHz, CD_3OD): δ 8.84 (d, $J = 6.8$ Hz, 1H), 7.72 (s, 1H), 7.44-7.40 (m, 3H), 4.02-3.99 (m, 1H), 3.82-3.69 (m, 2H), 3.67 (s, 3H), 3.35-3.24 (m, 5H), 2.92-2.88 (m, 4H), 2.76-2.71 (m, 1H), 2.64 (s, 3H).

Example 27

(*S,E*)-4-((3,5-difluoro-4-(3-((4-(methoxycarbonyl)-morpholin-2-yl)-methyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)-phenyl)-amino)-4-oxobutyl-2-enoic acid

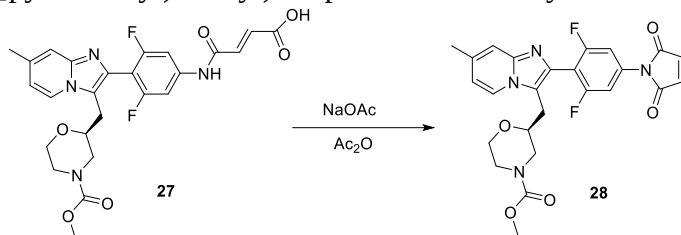


To a solution of **intermediate 25-4** (20 mg, 0.048 mmol) in tetrahydrofuran (5 mL) was added maleic anhydride (7 mg, 0.071 mmol) at room temperature. After the addition, the reaction system was heated to 60 °C and reacted overnight (16 h). After the reaction was completed as monitored by liquid chromatography, the reaction system was concentrated under reduced pressure. The residue was purified by preparative high performance liquid chromatography (trifluoroacetic acid method), and lyophilized to give **compound 27** (16 mg, 65% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 515.9$. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 12.9 (br, 1H), 10.88 (s, 1H), 8.78 (d, $J = 2.8$ Hz, 1H), 7.68 (s, 1H), 7.56 (d, $J = 10.4$ Hz, 2H), 7.31-7.30 (m, 1H), 6.53 (d, $J = 12$ Hz, 1H), 6.37 (d, $J = 12$ Hz, 1H), 3.90-3.87 (m, 1H), 3.76-3.62 (m, 3H), 3.57-3.54 (m, 3H), 3.25-3.07 (m, 4H), 2.85-2.80 (m, 1H), 2.52 (overlap, 3H).

Example 28

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(2,5-dioxo-2,5-dihydropyrrol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

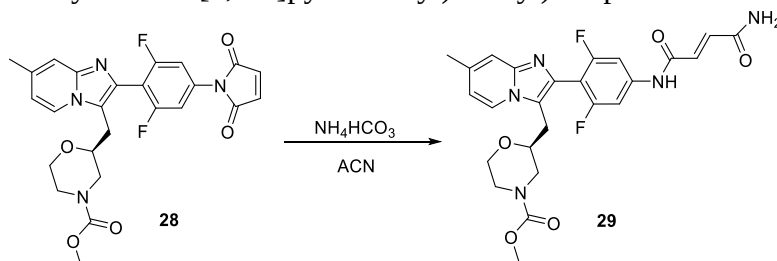


To a solution of **compound 27** (100 mg, 0.25 mmol) in acetic anhydride (3 mL) was added sodium acetate (32.7 mg, 0.4 mmol) at room temperature. The reaction system was stirred at room temperature for 4 h. Then the reaction system was concentrated under reduced pressure. The residue was purified by preparative high performance liquid chromatography (trifluoroacetic acid method), and lyophilized to give **compound 28** (40 mg, 30% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 497.1$. ^1H NMR (400 MHz, CD_3OD): δ 8.83 (d, $J = 7.2$ Hz, 1H), 7.71 (s, 1H), 7.56-7.52 (m, 2H), 7.43-7.39 (m, 1H), 7.68 (s, 2H), 4.02-3.99 (m, 1H), 3.83-3.69 (m, 3H), 3.67 (s, 3H), 3.36-3.33 (m, 1H), 3.30-3.22 (m, 2H), 2.92-2.90 (m, 1H), 2.87-2.73 (m, 1H), 2.64 (s, 3H).

Example 29

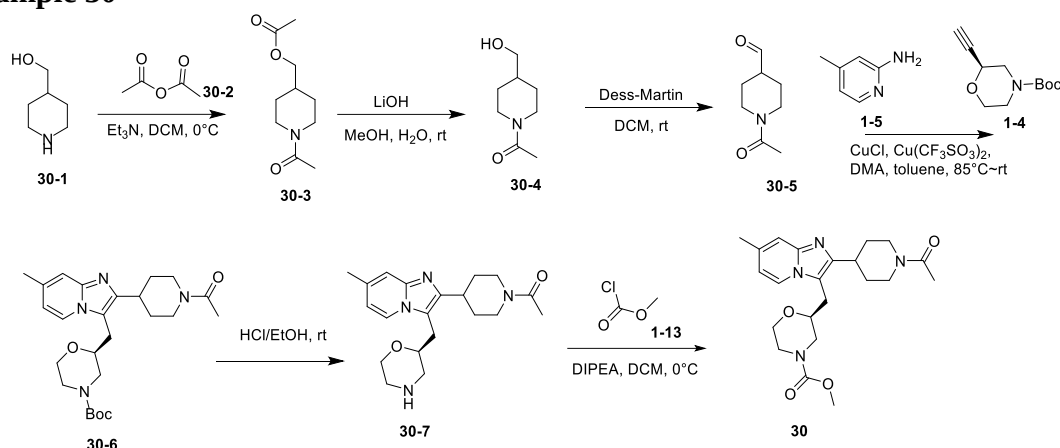
Preparation of methyl (S,E)-2-((2-(2,6-difluoro-4-(4-amino-4-oxobutyl-2-enamido)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



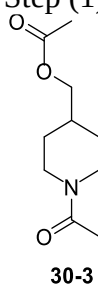
To a solution of **compound 28** (20 mg, 0.04 mmol) in acetonitrile (3 mL) was added aqueous ammonium bicarbonate solution (5%, 2 mL) at room temperature. The reaction system was reacted at room temperature for 2 h. Then the reaction system was concentrated under reduced pressure. The residue was purified by preparative high performance liquid chromatography (ammonium bicarbonate method), and lyophilized to give **compound 29** (10 mg, 40% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 514.1$. ^1H NMR (400 MHz, $\text{DMSO}-d_6$): δ 11.36 (s, 1H), 8.38 (d, $J = 7.2$ Hz, 1H), 7.93 (s, 1H), 7.45-7.42 (m, 3H), 7.33 (s, 1H), 6.82-6.80 (m, 1H), 6.35-6.28 (m, 2H), 3.73-3.62 (m, 3H), 3.55 (s, 3H), 3.51-3.46 (m, 1H), 3.25-3.22 (m, 2H), 3.02-3.00 (m, 2H), 2.83-2.79 (m, 1H), 2.37 (s, 3H).

Example 30

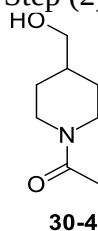


Step (1) Preparation of (1-acetylpiperidin-4-yl)methyl acetate



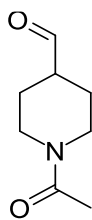
- 5 4-hydroxymethylpiperidine (5.2 g, 45.15 mmol) was dissolved in dichloromethane (60 mL), and triethylamine (13.5 g, 133.61 mmol) was added. The reaction system was cooled to 0 °C, added with acetic anhydride (13.8 g, 135.4 mmol), and reacted at room temperature for 15 h. Then the reaction system was washed with diluted hydrochloric acid (1 N, 40 mL). The organic phase was dried over anhydrous sodium sulfate and concentrated to give **compound 30-3** (6.2 g) in the form of a yellow oily liquid.
- 10

Step (2) Preparation of 1-(4-(hydroxymethyl)piperidin-1-yl)ethan-1-one



- 15 **Compound 30-3** (6.2 g) was dissolved in methanol (50 mL), and aqueous lithium hydroxide solution (3.75 N, 25 mL) was added. The reaction system was stirred at room temperature for 12 h. Then the reaction system was concentrated. The residue was separated by column chromatography (EA/MeOH = 50/1) to give **compound 30-4** (7.5 g) in the form of a colorless oily liquid. LC-MS: [M+H]⁺ = 158.0.

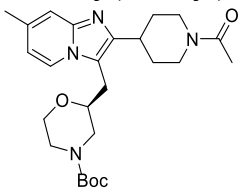
Step (3) Preparation of 1-acetylpiperidine-4-carbaldehyde



30-5

Compound 30-4 (3.2 g, 20.35 mmol) was dissolved in dichloromethane (40 mL), and Dess-Martin periodinane (12.9 g, 30.52 mmol) was added. The reaction system was reacted at room temperature for 1.5 h. Then the reaction system was added with saturated sodium thiosulfate solution (20 mL), diluted with water (40 mL), and extracted with dichloromethane (40 mL × 2). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (EA-EA/MeOH = 50/1) to give **compound 30-5** (1.0 g, 31% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 156.1$.

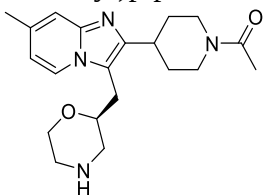
Step (4) Preparation of *tert*-butyl (S)-2-((2-(1-acetylpiperidin-4-yl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



30-6

Compound 30-5 (200 mg, 1.29 mmol) was dissolved in anhydrous toluene (40 mL), and 4-methyl-2-aminopyridine (140 mg, 1.29 mmol), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (272 mg, 1.29 mmol), copper(I) chloride (40 mg, 0.4 mmol), and copper(II) trifluoromethanesulfonate (140 mg, 0.39 mmol) were added. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with two drops of *N,N*-dimethylacetamide, heated and stirred for 6 h. Then the reaction system was cooled to room temperature and stirred overnight. The resulting reaction system was filtered, and the filtrate was concentrated. The residue was separated by column chromatography (DCM/MeOH = 20/1) to give **intermediate 30-6** (227 mg, 38% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 456.8$.

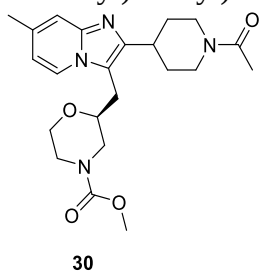
Step (5) Preparation of (S)-1-(4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)piperidin-1-yl)acetamide



30-7

Intermediate 30-6 (227 mg, 0.49 mmol) was dissolved in ethanol (2 mL), and a solution of HCl in ethanol (1 mL, 33%) was added. The reaction system was stirred at room temperature for 1.5 h. Then the reaction system was concentrated to give **intermediate 30-7** (250 mg, crude product), which was used directly in the next step. LC-MS $[M+H]^+ = 357.8$.

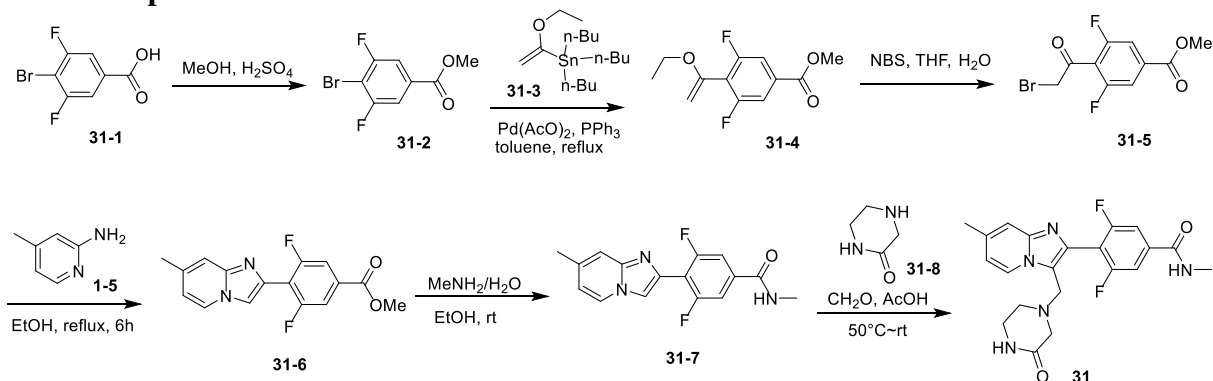
- 5 Step (6) Preparation of methyl (S)-2-((2-(1-acetylpiperidin-4-yl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



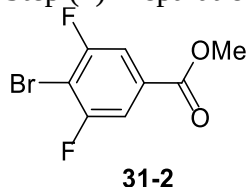
- 10 **Intermediate 30-7** (356 mg) was dissolved in dichloromethane (5 mL), and pyridine (244 mg, 3.09 mmol) was added, followed by the addition of methyl chloroformate (62 mg, 0.65 mmol). The reaction system was stirred at room temperature for 1.5 h. Then the reaction system was concentrated to give **compound 30** (360 mg, 87% yield).

LC-MS: $[M+H]^+ = 415.6$. $^1\text{H NMR}$ (400MHz, MeOD): δ 8.42(s, 1H), 7.56(d, 1H), 6.91(d, 1H), 3.82(m, 3H), 3.67(s, 3H), 3.60(m, 1H), 3.54(t, 4H) 3.38(dd, 1H), 3.12(m, 2H), 2.90(m, 2H), 2.83(m, 2H), 2.64(m, 2H), 2.10(s, 3H), 1.69(m, 4H).

15 Example 31



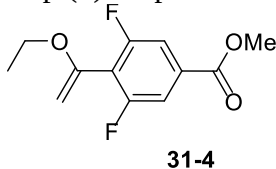
Step (1) Preparation of methyl 4-bromo-3,5-difluorobenzoate



- 20 4-bromo-3,5-difluorobenzoic acid (11.5 g, 48.5 mmol) was dissolved in methanol (100 mL), and concentrated sulfuric acid (1.0 mL) was added. The reaction system was reacted under reflux for 15 h. Then the reaction system was concentrated, diluted with water (50 mL) and

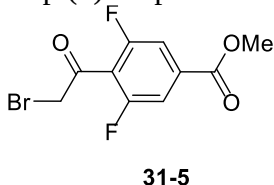
extracted with ethyl acetate (50 mL $\times$ 2). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE) to give **compound 31-2** (10 g, 82% yield) in the form of a colorless oily liquid.

Step (2) Preparation of methyl 4-(1-ethoxyethenyl)-3,5-difluorobenzoate



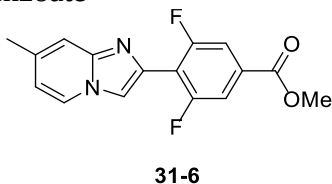
Compound 31-2 (10.0 g, 39.83 mmol) was dissolved in anhydrous toluene (90 mL), and palladium acetate (449 mg, 2 mmol), triphenylphosphine (2.0 g, 8.0 mmol) and tributyl(1-ethoxyethenyl)stannane (15.9 g, 44 mmol) were added. The reaction system was refluxed overnight under nitrogen atmosphere. Then the reaction system was concentrated, diluted with water (100 mL) and extracted with ethyl acetate (50 mL $\times$ 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated. The residue was separated by column chromatography (PE) to give **compound 31-4** (7.4 g, 76% yield) in the form of a colorless liquid. LC-MS: $[M+H]^+ = 214.8/242.8$.

Step (3) Preparation of methyl 4-(2-bromoacetyl)-3,5-difluorobenzoate



Compound 31-4 (7.4 g, 30.46 mmol) was dissolved in tetrahydrofuran (60 mL) and water (20 mL), and *N*-bromosuccinimide (5.4 g, 30.46 mmol) was added. The reaction system was reacted at room temperature for 1 h under nitrogen atmosphere. Then the reaction system was concentrated to half volume, diluted with water (60 mL), and extracted with ethyl acetate (30 mL $\times$ 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated to give **intermediate 31-5** (9.0 g) in the form of a yellow solid. LC-MS: $[M+H]^+ = 292.5$.

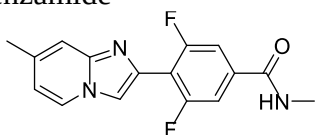
Step (4) Preparation of methyl 3,5-difluoro-4-(7-methylimidazo[1,2-*a*]pyridin-2-yl)benzoate



Intermediate 31-5 (9.0 g) was dissolved in ethanol (90 mL), and 4-methyl-2-aminopyridine (3.3 g, 30.47 mmol) was added. The reaction system was reacted under reflux for

12 h under nitrogen atmosphere. Then the reaction system was concentrated, diluted with water (60 mL), adjusted to pH 7-8 with saturated sodium bicarbonate solution, and extracted with ethyl acetate (40 mL $\times$ 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by column chromatography (PE/EA = 1/1) to give **intermediate 31-6** (4.2 g, 56% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 303.8$.

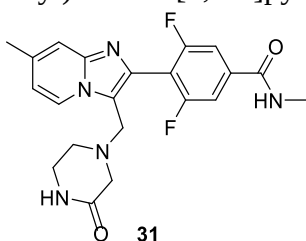
Step (5) Preparation of 3,5-difluoro-*N*-methyl-4-(7-methylimidazo[1,2-*a*]pyridin-2-yl)benzamide



31-7

Intermediate 31-6 (1.5 g, 4.96 mmol) was dissolved in ethanol (15 mL), and aqueous methylamine solution (6 mL, 25-30%) was added. The reaction system was stirred at room temperature for 12 h. Then the reaction system was concentrated. The residue was separated by column chromatography (EA-EA/MeOH = 50/1) to give **intermediate 31-7** (1.1 g, 73% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 302.8$.

Step (6) Preparation of 3,5-difluoro-4-(7-methyl-3-((3-oxopiperazin-1-yl)methyl)imidazo[1,2-*a*]pyridin-2-yl)benzamide

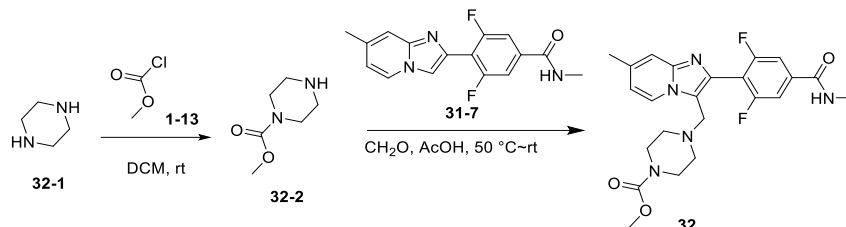


31

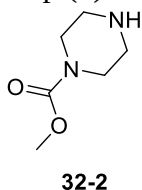
Intermediate 31-7 (150 mg, 0.49 mmol) and piperazin-2-one (50 mg, 0.49 mmol) were dissolved in acetic acid (1 mL), and aqueous formaldehyde solution (40 mg, 37%) was added. The reaction system was heated to 50 °C and reacted for 3 h. Then the reaction system was cooled to room temperature and stirred for 12 h. The resulting reaction system was diluted with dichloromethane (10 mL), and adjusted to pH 8 with aqueous sodium hydroxide solution (2 N). The aqueous phase was extracted with dichloromethane (10 mL $\times$ 2). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was purified by preparative thin-layer chromatography (DCM/MeOH = 20/1) to give **compound 31** (39.2 mg, 19% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 414.8$. ^1H NMR (400 MHz, MeOD): δ 8.52 (d, $J = 7.1$ Hz, 1H), 7.62 (d, $J = 8.0$ Hz, 2H), 7.40 (s, 1H), 6.94 (dd, $J = 7.1, 1.5$ Hz, 1H), 3.90 (s, 2H), 3.21 (t, $J = 5.4$ Hz, 2H), 3.05 (s, 2H), 2.97 (s, 3H), 2.66 – 2.54 (m, 2H), 2.48 (s, 3H).

Example 32

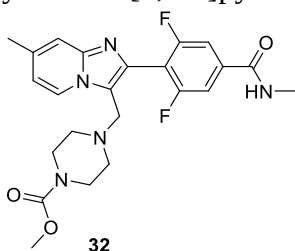


Step (1) Preparation of methyl piperazine-1-carboxylate



Piperazine (3 g, 34.8 mmol) was dissolved in dichloromethane (50 mL), and methyl chloroformate (658 mg, 6.96 mmol) was added in portions. The reaction system was reacted at room temperature for 30 min. Then the reaction system was added with aqueous potassium hydroxide solution (50 mL, 1 N) to quench the reaction, and extracted with dichloromethane (50 mL $\times$ 3). The organic phases combined, dried over anhydrous sodium sulfate, and concentrated to give a white solid (1.8 g, 36% yield). LC-MS: $[M+H]^+ = 145.7$.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)piperazine-4-carboxylate

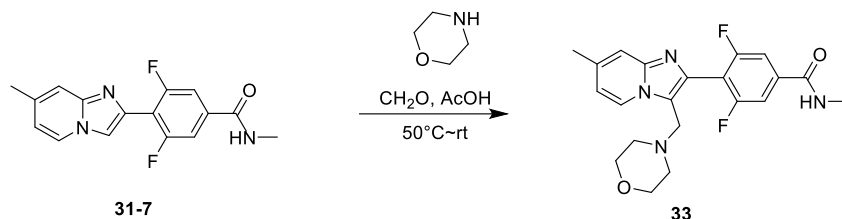


Intermediate 31-7 (150 mg, 0.49 mmol) and **compound 32-2** (180 mg) were dissolved in acetic acid (1 mL), and aqueous formaldehyde solution (40 mg, 37%) was added. The reaction system was heated to 50 °C and reacted for 3 h. Then the reaction system was cooled to room temperature and stirred for 12 h. The resulting reaction system was diluted with dichloromethane (10 mL), and adjusted to pH 8 with aqueous sodium hydroxide solution (2 N). The aqueous phase was extracted with dichloromethane (15 mL $\times$ 2). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was separated by HPLC to give **compound 32** (98.2 mg, 43% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 458.7$. $^1\text{H NMR}$: $^1\text{H NMR}$ (400 MHz, MeOD): δ 8.55 (d, $J = 7.0$ Hz, 1H), 7.60 (d, $J = 7.9$ Hz, 2H), 7.39 (s, 1H), 6.94 (d, $J = 7.1$ Hz, 1H), 3.80 (s, 2H), 3.66 (s, 3H), 3.47 – 3.36 (m, 4H), 2.97 (s, 3H), 2.49 (s, 3H), 2.37 – 2.29 (m, 4H).

Example 33

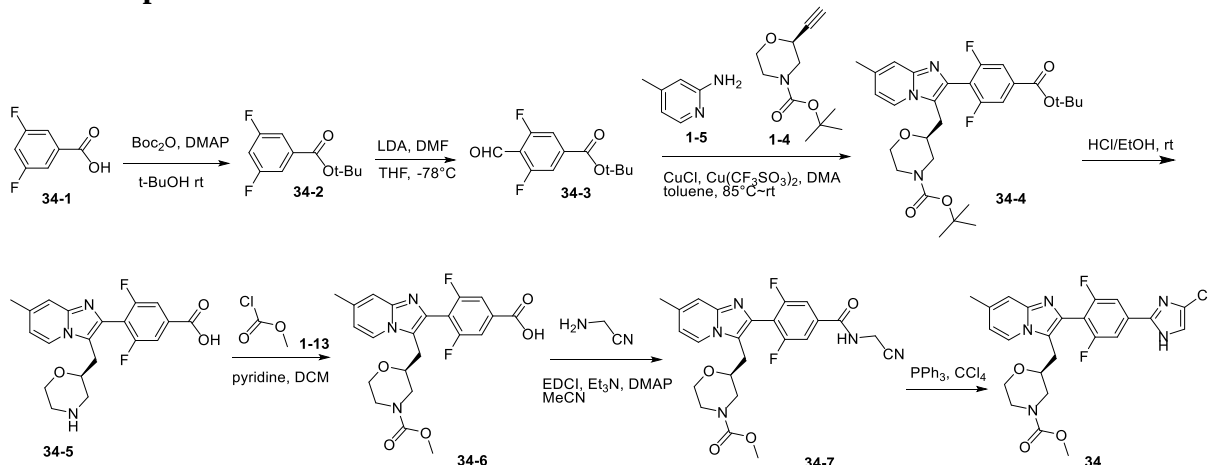
- 5 Preparation of 3,5-difluoro-4-(7-methyl-3-morpholinomethylimidazo[1,2-*a*]pyridin-2-yl)benzamide



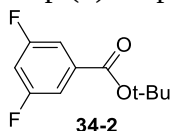
- Intermediate 31-7** (100 mg, 0.33 mmol) and morpholine (29 mg, 0.33 mmol) were dissolved in acetic acid (0.5 mL), and aqueous formaldehyde solution (28 mg, 37%) was added. The reaction system was heated to 50 °C and reacted for 3 h. Then the reaction system was cooled to room temperature and stirred for 12 h. The resulting reaction system was diluted with dichloromethane (15 mL), and adjusted to pH 8 with aqueous sodium hydroxide solution (2 N). The aqueous phase was extracted with dichloromethane (10 mL $\times$ 2). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was purified by HPLC to give **compound 33** (50.2 mg, 38% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 401.7$. $^1\text{H NMR}$ (400 MHz, CDCl_3): δ 8.44 (d, $J = 7.0$ Hz, 1H), 7.58 – 7.37 (m, 4H), 6.80 (d, $J = 6.9$ Hz, 1H), 3.71 (s, 2H), 3.67 – 3.60 (m, 4H), 3.03 (d, $J = 4.6$ Hz, 3H), 2.48 (s, 3H), 2.39 – 2.31 (m, 4H).

Example 34

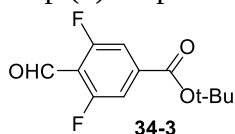


Step (1) Preparation of *tert*-butyl 3,5-difluorobenzoate



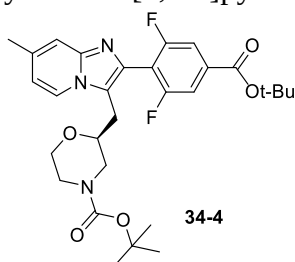
3,5-difluorobenzoic acid (10.0 g, 63.25 mmol) was dissolved in *tert*-butanol (100 mL), and DMAP (2.3 g, 18.98 mmol) and di-*tert*-butyl dicarbonate (27.6 g, 126.50 mmol) were added. The reaction system was reacted at room temperature for 15 h. Then the reaction system was concentrated, diluted with water (80 mL) and extracted with ethyl acetate (50 mL × 2). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 20/1) to give **compound 34-2** (12.0 g, 88% yield) in the form of a colorless oily liquid.

Step (2) Preparation of *tert*-butyl 3,5-difluoro-4-formylbenzoate



LDA (2 N, 8.4 mL) was dissolved in tetrahydrofuran (10 mL). The reaction system was cooled to -78 °C under nitrogen atmosphere. **Compound 34-2** (3.0 g, 14.0 mmol) was dissolved in tetrahydrofuran (20 mL), and the solution was added to the reaction system. The reaction system was stirred at -78 °C for 1.5 h, added with DMF (1.2 g, 16.8 mmol), and stirred for another 1 h with the temperature maintained not higher than -70 °C. Then the reaction system was added with saturated ammonium chloride solution (60 mL) to quench the reaction, and extracted with ethyl acetate (30 mL × 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated. The residue was separated by column chromatography (PE/EA = 10/1) to give **compound 34-3** (1.5 g, 44% yield) in the form of a yellow solid.

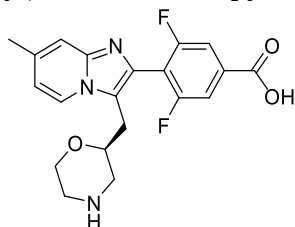
Step (3) Preparation of *tert*-butyl (S)-2-((2-(4-(*tert*-butoxycarbonyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-carboxylate



Compound 34-3 (480 mg, 1.98 mmol) was dissolved in anhydrous toluene (4 mL), and 4-methyl-2-aminopyridine (215 mg, 1.98 mmol), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (419 mg, 1.98 mmol), copper(I) chloride (62 mg, 0.63 mmol), and copper(II) trifluoromethanesulfonate (216 mg, 0.60 mmol) were added. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with two drops of *N,N*-dimethylacetamide, heated and stirred for 6 h. Then the reaction system was cooled to room temperature and stirred overnight. The resulting reaction system was filtered, and the filtrate was concentrated. The

residue was separated by column chromatography (PE/EA = 2/1) to give **intermediate 34-4** (340 mg, 31% yield) in the form of a brown solid. LC-MS: $[M+H]^+ = 544.7$.

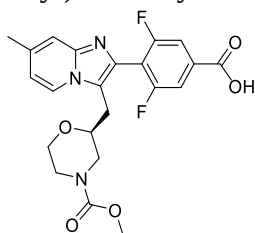
Step (4) Preparation of (S)-3,5-difluoro-4-(7-methyl-3-(morpholinyl-2-methyl)imidazo[1,2-a]pyridin-2-yl)benzoic acid



34-5

Intermediate 34-4 (340 mg, 0.62 mmol) was dissolved in ethanol (3 mL), and a solution of HCl in ethanol (1 mL, 33%) was added. The reaction system was stirred at room temperature for 1.5 h. Then the reaction system was concentrated to give **intermediate 34-5** (400 mg) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 388.7$.

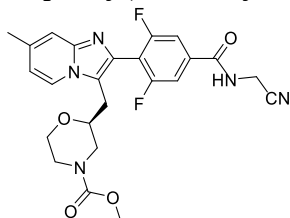
Step (5) Preparation of (S)-3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)benzoic acid



34-6

Intermediate 34-5 (400 mg) was dissolved in dichloromethane (5 mL), and pyridine (244 mg, 3.09 mmol) was added, followed by the addition of methyl chloroformate (62 mg, 0.65 mmol). The reaction system was stirred at room temperature for 1.5 h, and concentrated to give **intermediate 34-6** (190 mg). LC-MS: $[M+H]^+ = 446.6$.

Step (6) Preparation of methyl (S)-2-((2-(4-((cyanomethyl)carbamoyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

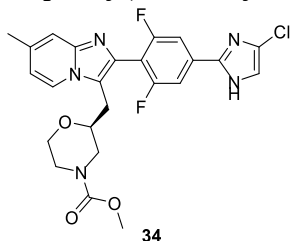


34-7

Intermediate 34-6 (445 mg, 1 mmol) was dissolved in CH_3CN (10 mL), and triethylamine (1.2 mmol), DMAP (1.2 mmol) and EDCI (1.2 mmol) were added. The reaction system was stirred at room temperature overnight. After the reaction was completed, the reaction system was

purified by column chromatography to give **intermediate 34-7** (362 mg, 75% yield). LC-MS: $[M+H]^+ = 484.6$.

Step (7) Preparation of methyl (S)-2-((2-(4-(4-chloro-1H-imidazol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

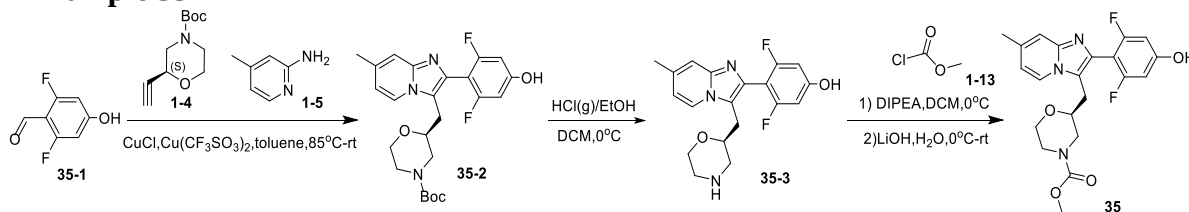


5

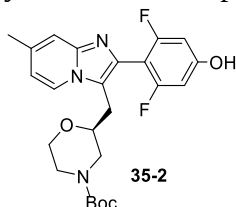
Intermediate 34-7 (483 mg, 1 mmol) was dissolved in THF (10 mL), and a certain amount of CCl_4 and PPh_3 was added. The reaction system was heated to 50 °C, and detected by LC-MS for completion. After the reaction was completed, the reaction system was purified by column chromatography to give **compound 34** (50 mg, 10% yield). LC-MS: $[M+H]^+ = 502.1$.

10

Example 35



Step (1) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-hydroxyphenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



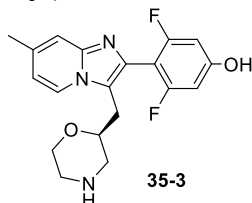
15

2,6-difluoro-4-hydroxybenzaldehyde (2.0 g, 9.05 mmol), 4-methylpyridin-2-amine (0.97 g, 9.05 mmol), **compound 1-4** (1.9 g, 9.05 mmol), copper(I) chloride (0.27 g, 2.70 mmol) and copper(II) trifluoromethanesulfonate (0.98 g, 2.70 mmol) were dissolved in toluene solution (20 mL) under nitrogen atmosphere. The reaction system was reacted at 85 °C for 10 min, added with *N,N*-dimethylacetamide (0.5 mL), and reacted at 85 °C for another 5 h. Then the reaction system was reacted at room temperature overnight. After the reaction was completed, the reaction system was added with water (40 mL), and extracted with dichloromethane (50 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography (40 g of silica gel, petroleum ether:ethyl acetate =

20

1:1) to give **intermediate 35-2** (1.2 g, 29% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 460.1$.

Step (2) Preparation of (S)-3,5-difluoro-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)phenol



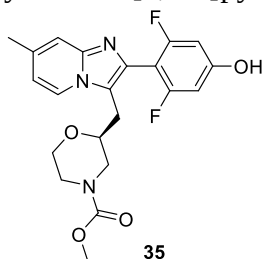
5

Intermediate 35-2 (1.0 g, 1.91 mmol) was dissolved in dichloromethane (10 mL), and a solution of HCl in ethanol (2 mL, 33%) was added dropwise. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. After the reaction was completed, the reaction system was added dropwise with saturated sodium bicarbonate solution to adjust the pH to 8, added with water (30 mL), and extracted with dichloromethane (40 mL $\times$ 3). The organic phases were combined, washed with saturated brine (40 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 35-3** (800 mg) in the form of a yellow oil. LC-MS $[M+H]^+ = 360.7$.

10

Step (3) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-hydroxyphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

15



Intermediate 35-3 (350 mg, 0.97 mmol) and *N,N*-diisopropylethylamine (378 mg, 2.92 mmol) were dissolved in dichloromethane (10 mL), and methyl chloroformate (110 mg, 1.16 mmol) was added dropwise with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 3 h. After the reaction was completed, the reaction system was added with lithium hydroxide (1.0 mL, 3 N) solution, and reacted at room temperature for 2 h. After the reaction was completed, the reaction system was adjusted to pH 5 with diluted hydrochloric acid, diluted with water (10 mL), and extracted with dichloromethane (15 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, and dried. The organic phase was filtered and concentrated by rotary evaporation. The residue was separated by HPLC to give **compound 35** (150 mg, 37% yield) in the form of a yellow solid.

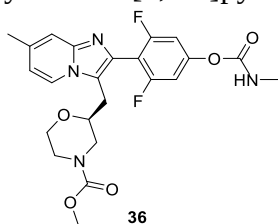
20

25

LC-MS: $[M+H]^+ = 418.7$. ^1H NMR (400 MHz, MeOD): δ 8.57 (d, $J = 6.9$ Hz, 1H), 7.47 (s, 1H), 7.08 (d, $J = 6.9$ Hz, 1H), 6.60 (d, $J = 9.6$ Hz, 2H), 3.92 - 3.76 (m, 3H), 3.68 (s, 3H), 3.69 - 3.57 (m, 1H), 3.39 (dd, $J = 11.9, 2.7$ Hz, 1H), 3.16 - 3.10 (m, 2H), 2.98 - 2.86 (m, 1H), 2.74 - 2.63 (m, 1H), 2.53 (s, 3H).

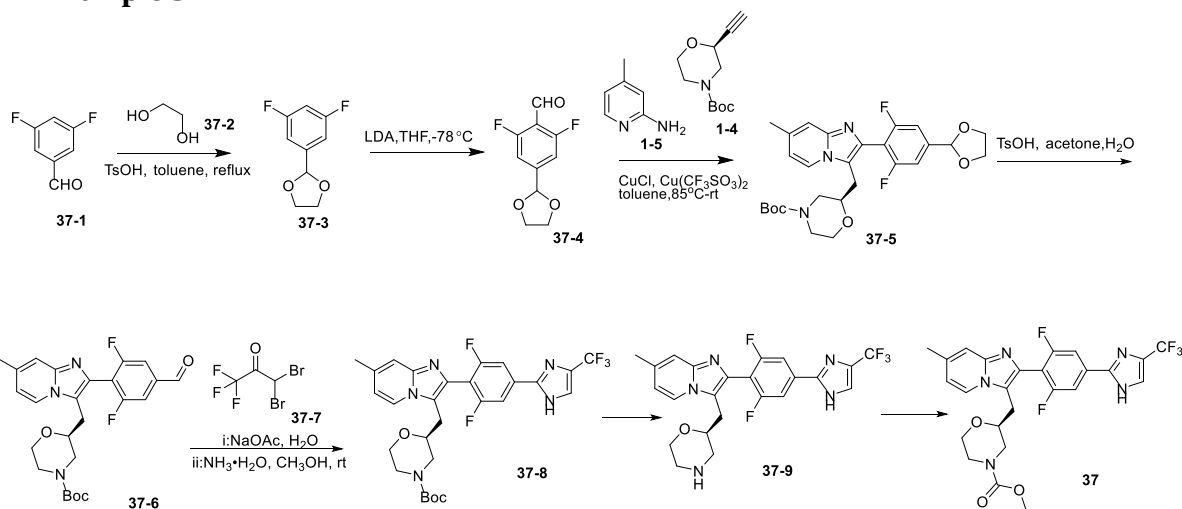
5 Example 36

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyloxy)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

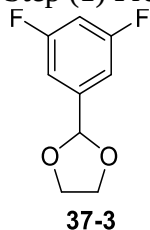


Compound 35 (359 mg, 1 mmol) was dissolved in CH_2Cl_2 (10 mL), and triethylamine (2 mmol) was added. Methylaminoformyl chloride (186 mg, 2 mmol) was dissolved in anhydrous CH_2Cl_2 (2 mL) and the solution was added dropwise to the reaction system. After the reaction was completed, the reaction system was neutralized and washed, dried, filtered, and purified by column chromatography to give **compound 36** (62 mg, 13% yield). LC-MS $[M+H]^+ = 475.1$.

Example 37



Step (1) Preparation of 2-(3,5-difluorophenyl)-1,3-dioxolane

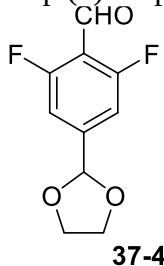


3,5-difluorobenzaldehyde (5.68 g, 40 mmol), *p*-toluenesulfonic acid (0.69 g, 4 mmol) and ethylene glycol (7.45 g, 120 mmol) were dissolved in toluene (80 mL). The reaction system was

refluxed for separating water overnight. Then the reaction system was cooled to room temperature, diluted with ethyl acetate (40 mL) and washed with NaHCO₃ solution (40 mL × 2). The organic phase was dried over Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The residue was purified by silica gel column chromatography (PE:EA = 20:1) to give **compound 37-3** (7.0 g, 94% yield) in the form of a white solid.

¹H NMR (400 MHz, CDCl₃) : δ 7.03-7.00 (m, 2H), 6.82-6.76 (m, 1H), 5.79 (s, 1H), 4.09-4.05 (m, 2H), 4.07-3.99 (m, 2H).

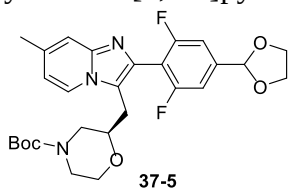
Step (2) Preparation of 4-(1,3-dioxolan-2-yl)-2,6-difluorobenzaldehyde



Compound 37-3 (7.0 g, 37.8 mmol) was dissolved in tetrahydrofuran (50 mL). The reaction system was cooled to -78 °C under nitrogen atmosphere, and added dropwise with LDA (22.4 mL, 44.8 mmol). After the completion of the dropwise addition, the reaction system was reacted at this temperature for 30 min, then added with anhydrous DMF (3.3 g, 44.3 mmol). After the completion of the dropwise addition, the reaction system was reacted at this temperature for 2.5 h. Then the reaction system was added with saturated NH₄Cl (20 mL) to quench the reaction, diluted with water (50 mL), and extracted with ethyl acetate (50 mL × 3). The organic phases were combined, washed with saturated sodium chloride solution (50 mL), dried over anhydrous sodium sulfate and filtered, and the filtrate was concentrated by rotary evaporation. The resulting crude product was purified by silica gel column chromatography (PE:EA = 10:1) to give **intermediate 37-4** (0.90 g, 12% yield) in the form of a white solid.

¹H NMR (300 MHz, DMSO) : δ 10.18 (s, 1H), 7.28 (d, J = 9.8 Hz, 2H), 5.82 (s, 1H), 4.06-4.00 (m, 2H), 4.01-3.93 (m, 2H).

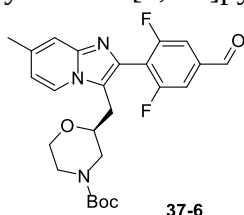
Step (3) Preparation of *tert*-butyl (S)-2-((2-(4-(1,3-dioxolan-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 37-4 (100 mg, 0.47 mmol), **compound 1-4** (100 mg, 0.47 mmol), 4-methylpyridin-2-amine (51 mg, 0.47 mmol), copper(I) chloride (25 mg, 0.14 mmol), and copper(II) trifluoromethanesulfonate (51 mg, 0.14 mmol) were dissolved in toluene (3 mL) in a

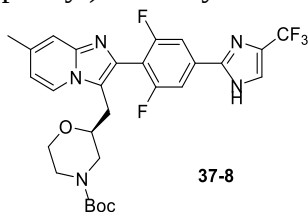
single-necked flask. The reaction system was heated to 85 °C under nitrogen atmosphere, added dropwise with two drops of DMA and reacted at this temperature for 5 h. Then the reaction system was cooled to room temperature and stirred overnight. The resulting reaction system was diluted with water (20 mL) and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The resulting crude product was purified by silica gel column chromatography (PE:EA = 5:1) to give **intermediate 37-5** (150 mg, 62% yield). LC-MS: [M+H]⁺ = 516.7.

Step (4) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-formylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 37-5 (150 mg, 0.29 mmol) was dissolved in acetone (10 mL) and water (1 mL), and TsOH (90 mg, 0.3 mmol) was added. The reaction system was heated to 50 °C and reacted overnight. Then the reaction system was diluted with water (50 mL), added with Na₂CO₃ (92 mg, 0.87 mmol) and (Boc)₂O (75 mg, 0.35 mmol), stirred for 1.5 h and extracted with ethyl acetate (15 mL × 2). The organic phases were combined, dried over anhydrous Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The resulting crude product was purified by silica gel column chromatography (PE:EA = 5:1) to give **intermediate 37-6** (60 mg, 44.8% yield) in the form of a white solid. LC-MS: [M+H]⁺ = 472.7.

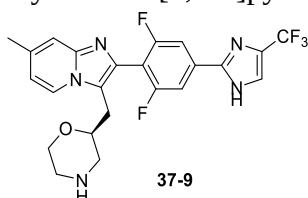
Step (5) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(4-trifluoromethyl-1H-imidazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



3,3-dibromo-1,1,1-trifluoropropan-2-one (35 mg, 0.13 mmol) was dissolved in sodium acetate (2 mL, 15 weight%) solution. The reaction system was stirred at 90 °C for 30 min and cooled to room temperature. **Intermediate 37-6** (50 mg, 0.11 mmol) was dissolved in methanol (3 mL) and strong ammonia (0.6 mL), and the solution was added to the reaction system, and stirred at room temperature overnight. Then the reaction system was concentrated by rotary evaporation. The residual liquid was extracted with ethyl acetate (5 mL × 3) and washed with

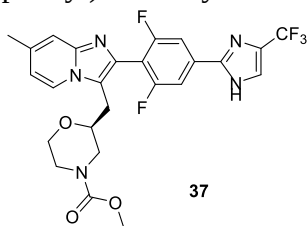
saturated brine (10 mL). The organic phase was dried over Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography (dichloromethane:methanol = 50:1) to give **intermediate 37-8** (45 mg, 72.2% yield) in the form of a white solid. LC-MS: [M+H]⁺ = 578.6.

- 5 Step (6) Preparation of (S)-2-((2-(2,6-difluoro-4-(4-trifluoromethyl-1H-imidazole)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine



- Intermediate 37-8** (45 mg, 0.08 mmol) was dissolved in dichloromethane (3 mL), and a solution of HCl in ethanol (0.5 mL) was added. The reaction system was stirred at room temperature for 3 h. Then the reaction system was concentrated by rotary evaporation to give **intermediate 37-9** (80 mg, crude product). LC-MS: [M+H]⁺ = 478.7.

Step (7) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-trifluoromethyl-1H-imidazol-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

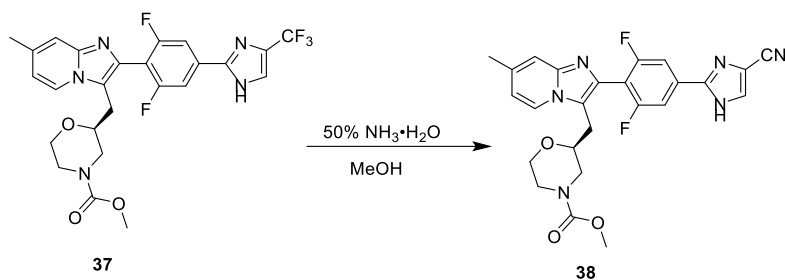


- 15 **Intermediate 37-9** (80 mg, crude product) and DIPEA (30 mg, 0.23 mmol) were dissolved in dichloromethane (3 mL). The reaction system was cooled to 0 °C and stirred for 10 min. Then the reaction system was added dropwise with a solution of methyl chloroformate in dichloromethane (0.1 N, 0.8 mL), stirred for 3 h, added with water (10 mL) to quench the reaction and extracted with ethyl acetate (10 mL × 3). The organic phase was dried over Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography (dichloromethane:methanol = 20:1), prepared and lyophilized to give **compound 37** (7.0 mg, 16.3% yield) in the form of a white solid product.

- LC-MS: [M +H]⁺ = 536.7. ¹H NMR (400 MHz, CDCl₃): δ 13.18 (s, 1H), 8.58 (brs, 1H), 7.97 – 7.93 (m, 3H), 7.53 (s, 1H), 7.16 (d, J = 6.0 Hz, 1H), 4.05 – 3.70 (m, 3H), 3.71 (s, 3H), 3.64 – 3.56 (m, 1H), 3.45 – 3.30 (m, 1H), 3.09 – 2.85 (m, 3H), 2.74 – 2.64 (m, 1H), 2.60 (s, 3H).

Example 38

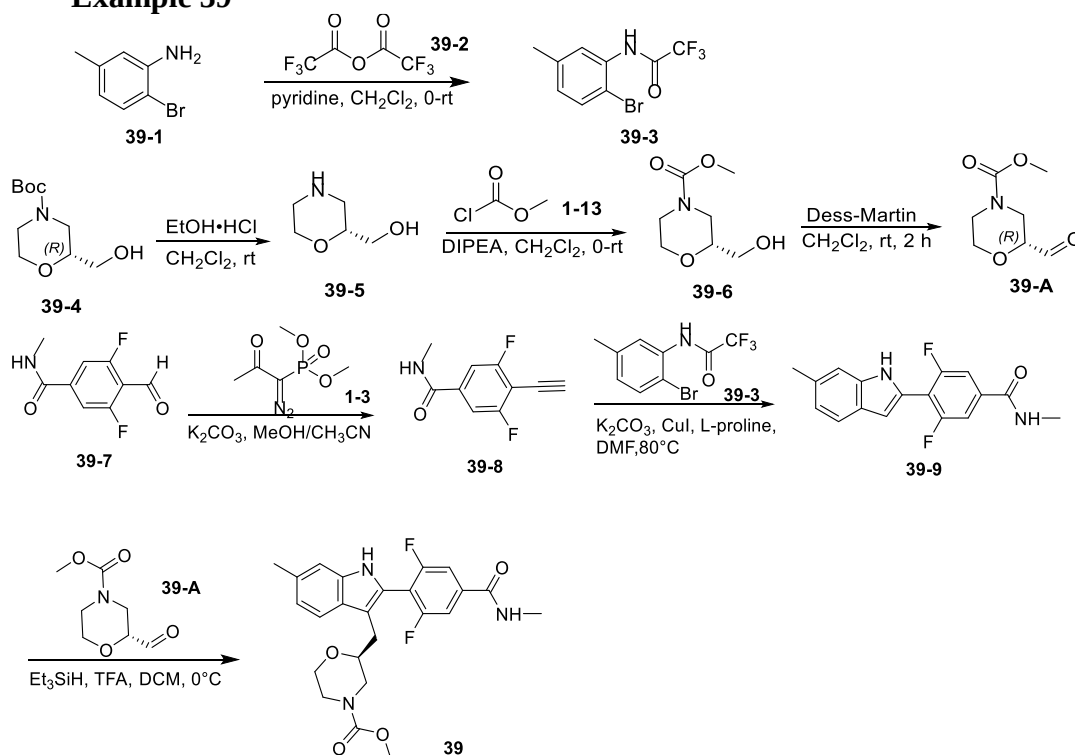
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-cyano-1H-imidazol-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



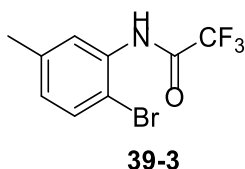
Compound 37 (30 mg, 0.05 mmol) was dissolved in methanol (2 mL), and strong ammonia (0.8 mL) and water (0.8 mL) were added. The reaction system was stirred at room temperature for 36 h. Then the reaction system was diluted with water (10 mL) and extracted with ethyl acetate (10 mL × 3). The organic phase was dried over Na₂SO₄ and filtered, and the filtrate was concentrated by rotary evaporation. The crude product was purified by preparative thin-layer chromatography (dichloromethane:methanol = 20:1) and lyophilized to give **compound 38** (11 mg, 44.6% yield) in the form of a white solid product.

LC-MS: [M + H]⁺ = 493.6. ¹H NMR (400 MHz, CDCl₃): δ 15.61 (brs, 1H), 8.31 (d, J = 7.0 Hz, 1H), 7.74 (s, 1H), 7.47 (s, 1H), 7.33 (d, J = 7.2 Hz, 2H), 6.85 (d, J = 7.0 Hz, 1H), 3.97 – 3.72 (m, 3H), 3.66 (s, 3H), 3.62 – 3.52 (m, 1H), 3.42 – 3.30 (m, 1H), 3.03 – 2.85 (m, 3H), 2.65 – 2.55 (m, 1H), 2.54 (s, 3H).

Example 39



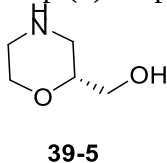
Step (1) Preparation of *N*-(2-bromo-5-methylphenyl)-2,2,2-trifluoroacetamide



2-bromo-5-methylbenzenamine (2.5 g, 13.44 mmol) and pyridine (3.3 mL, 40.32 mmol) were dissolved in dichloromethane (30 mL). The reaction system was added dropwise with trifluoroacetic anhydride (3.4 g, 16.12 mmol) at 0 °C and reacted at room temperature overnight.

- 5 The reaction mixture was washed with water (10 mL × 2) and diluted hydrochloric acid (30 mL, 1 mol/L) sequentially, dried over anhydrous magnesium sulfate, filtered, and concentrated under reduced pressure to give **compound 39-3** (3.4 g, 90% yield) in the form of a gray solid, LC-MS: $[M+H]^+ = 281.7$.

Step (2) Preparation of *R*-(morpholin-2-yl) methanol

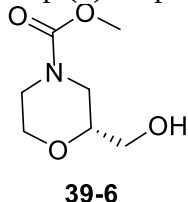


10

(*R*)-*N*-Boc-2-hydroxymethylmorpholine (3.5 g, 16.13 mmol) was dissolved in dichloromethane (20 mL), and a solution of HCl in ethanol (3.5 mL, 33%) was added dropwise at 0 °C. The reaction system was stirred at room temperature overnight, and concentrated under reduced pressure to give **compound 39-5** (1.8 g, 95% yield, crude product) in the form of a yellow oil. LC-MS: $[M+H]^+ = 118.1$.

15

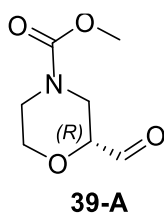
Step (3) Preparation of methyl (*R*)-2-(hydroxymethyl)morpholine-4-carboxylate



Compound 39-5 (4.0 g, 16.13 mmol) was dissolved in dichloromethane (40 mL), and DIPEA (12.5 g, 96.78 mmol) was added dropwise at 0 °C. Then methyl chloroformate (1.97 g, 20.97 mmol) was dissolved in dichloromethane (10 mL) and added dropwise to the reaction system at 0 °C. The reaction system was stirred at room temperature overnight, added with dichloromethane (100 mL), washed with brine (35 mL × 2), and concentrated under reduced pressure. The residue was purified by silica gel column chromatography (20 g of silica gel, ethyl acetate:petroleum ether = 1:2) to give **intermediate 39-6** (1.0 g, 35% yield) in the form of a colorless oil. LC-MS: $[M+H]^+ = 176.9$.

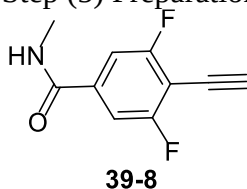
25

Step (4) Preparation of methyl (*R*)-2-formylmorpholine-4-carboxylate



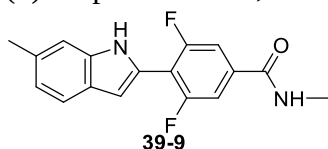
Intermediate 39-6 (600 mg, 3.43 mmol) was dissolved in dichloromethane (10 mL), and Dess-Martin periodinane (1.74 g, 4.11 mmol) was added. The reaction system was reacted at room temperature for 2 h, added with saturated sodium thiosulfate (2 mL) and stirred for 30 min. Then the reaction system was extracted with dichloromethane (30 mL × 2), washed with brine (15 mL), and purified by silica gel column chromatography (12 g of silica gel, ethyl acetate:petroleum ether = 1:1) to give **intermediate 39-A** (180 mg, 30% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 174.5$.

Step (5) Preparation of 4-ethynyl-3,5-difluoro-*N*-methylbenzamide



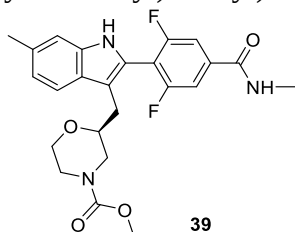
Dimethyl (1-diazo-2-oxopropyl)phosphonate (2.2 g, 11.3 mmol) was dissolved in acetonitrile (15 mL) and methanol (3 mL), then potassium carbonate (2.3 g, 16.56 mmol) was added. The reaction system was stirred at room temperature for 30 min, added with 3,5-difluoro-4-formyl-*N*-methylbenzamide (1.5 g, 7.53 mmol), and stirred at room temperature overnight. Then the reaction system was filtered and concentrated. The residue was purified by silica gel column chromatography (20 g of silica gel, ethyl acetate:petroleum ether = 1:5) to give **intermediate 39-8** (1.2 g, 81% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 196.1$.

Step (6) Preparation of 3,5-difluoro-4-(6-methyl-1*H*-indol-2-yl)benzamide



Intermediate 39-8 (1.4 g, 7.18 mmol), **intermediate 39-3** (2.43 g, 8.61 mmol), potassium carbonate (1.97 g, 15.09 mmol), copper(I) iodide (0.14 g, 1.43 mmol) and *L*-proline (0.25 g, 2.87 mmol) were added to *N,N*-dimethylformamide (20 mL). The reaction system was reacted at 80 °C overnight under nitrogen atmosphere. Then the reaction system was cooled, poured into water (60 mL), extracted with ethyl acetate (50 mL × 2), washed with saturated brine (30 mL × 3), and purified by silica gel column chromatography (20 g of silica gel, developing agent: ethyl acetate:petroleum ether = 1:2) to give **intermediate 39-9** (800 mg, 44% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 301.8$.

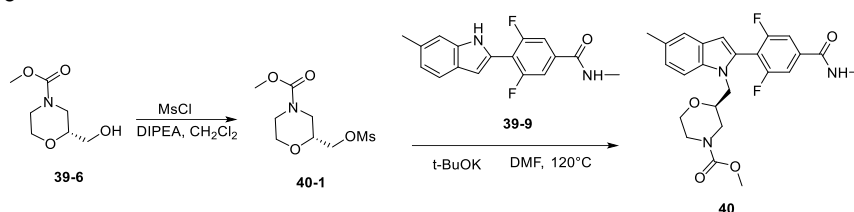
Step (7) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylindol-3-yl)methyl)morpholine-4-carboxylate



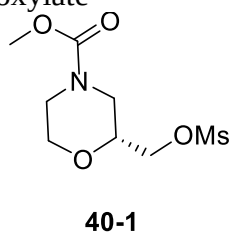
Trifluoroacetic acid (304 mg, 2.66 mmol) and triethylsilane (617 mg, 5.32 mmol) were dissolved in dichloromethane (3 mL) in a round-bottom flask (25 mL). **Intermediate 39-9** (160 mg, 0.53 mmol) and **intermediate 39-A** (110 mg, 0.64 mmol) were then dissolved in dichloromethane (3 mL), and the solution was added dropwise to the reaction system at 0 °C. The reaction system was stirred at room temperature overnight. Then the reaction system was added with water (20 mL), extracted with dichloromethane (20 mL × 2), and separated by preparative reverse phase chromatography to give **compound 39** (18 mg, 7.4% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 458.2$. ^1H NMR (400 MHz, CDCl_3): δ 8.18 (brs, 1H), 7.62 (d, $J = 8.2$ Hz, 1H), 7.43 (d, $J = 7.5$ Hz, 2H), 7.21 (s, 1H), 7.02 (d, $J = 8.1$ Hz, 1H), 6.29 (brs, 1H), 3.88 – 3.75 (m, 3H), 3.64 (s, 3H), 3.60 – 3.55 (m, 1H), 3.45–3.39 (m, 1H), 3.07 (d, $J = 3.6$ Hz, 3H), 3.04 – 2.98 (m, 1H), 2.90–2.88 (m, 1H), 2.78 (dd, $J = 14.9, 6.6$ Hz, 1H), 2.50 (s, 3H), 2.49 – 2.41 (m, 1H).

Example 40



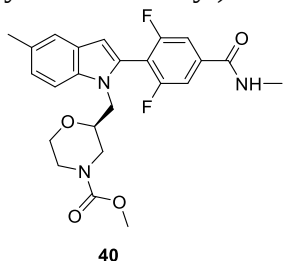
Step (1) Preparation of methyl (R)-2-(((methylsulfonyl)oxo)methyl)morpholine-4-carboxylate



Intermediate 39-6 (400 mg, 2.28 mmol) and *N,N*-diisopropylethylamine (737 mg, 5.71 mmol) were dissolved in dichloromethane (10 mL), and methylsulfonyl chloride (313 mg, 2.74 mmol) was added dropwise at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at room temperature overnight. Then the reaction system was added with

dichloromethane (20 mL), washed with saturated brine (15 mL × 2), dried over anhydrous magnesium sulfate, filtered, and concentrated under reduced pressure to give **intermediate 40-1** (600 mg, crude product) in the form of a brown oil. LC-MS: $[M+H]^+ = 254.9$.

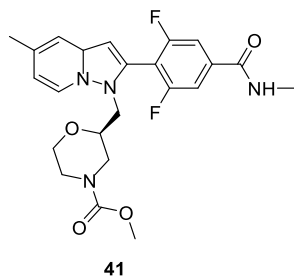
Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-5-methyl-1*H*-indol-1-yl)methyl)morpholine-4-carboxylate



Intermediate 39-9 (250 mg, 0.83 mmol) was dissolved in *N,N*-dimethylformamide (5 mL), and intermediate **40-1** (421 mg, 1.67 mmol) and potassium *tert*-butoxide (205 mg, 1.83 mmol) were added at room temperature. The reaction system was stirred at 120 °C overnight under nitrogen atmosphere. Then the reaction system was cooled to room temperature, added with water (20 mL), and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous magnesium sulfate, filtered and concentrated. The crude product was purified by preparative reverse phase chromatography to give **compound 40** (35 mg, 9.2% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 458.8$. ^1H NMR (400 MHz, CDCl_3) : δ 7.57 (d, $J = 8.1$ Hz, 1H), 7.43 (d, $J = 7.7$ Hz, 1H), 7.22 (s, 1H), 7.03 (d, $J = 8.1$ Hz, 1H), 6.61 (s, 1H), 6.22 (brs, 1H), 4.15 – 3.99 (m, 2H), 3.85 – 3.60 (m, 3H), 3.67 (s, 3H), 3.60 – 3.50 (m, 1H), 3.32 (t, 1H), 3.08 (d, $J = 4.7$ Hz, 3H), 2.81 (t, $J = 11.6$ Hz, 1H), 2.54 (s, 3H), 2.39 (dd, $J = 13.0, 10.8$ Hz, 1H).

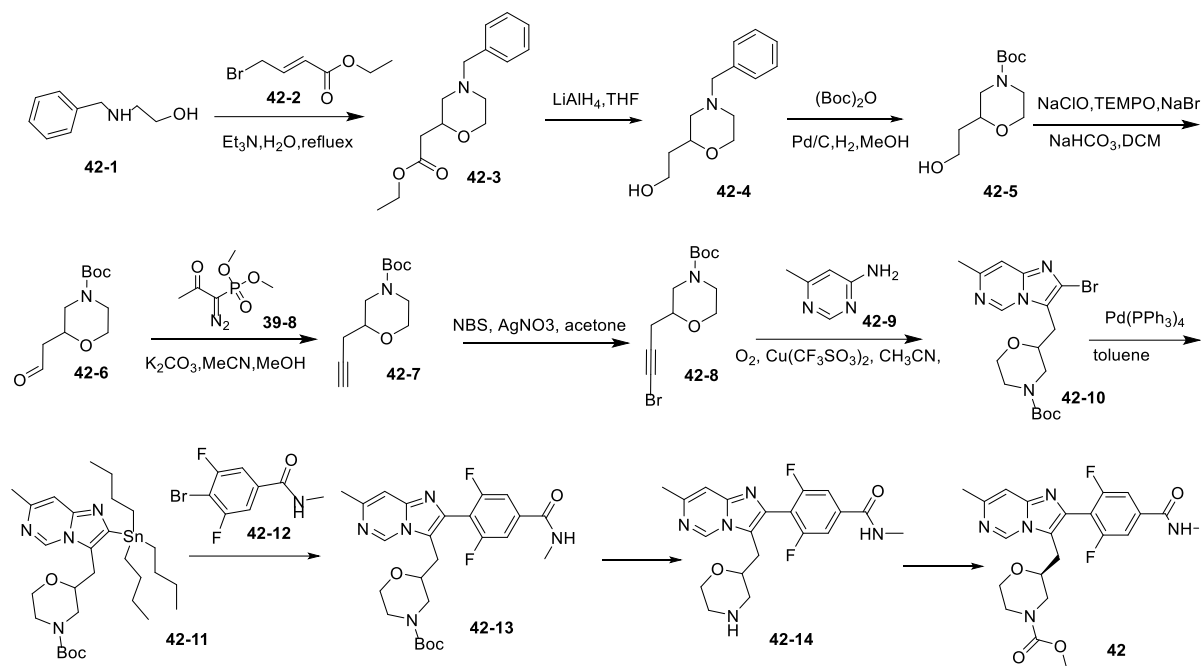
Example 41



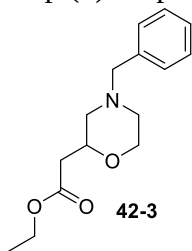
The compound was prepared according to the preparation method as described in Example 40.

LC-MS: $[M+H]^+ = 461.2$.

Example 42

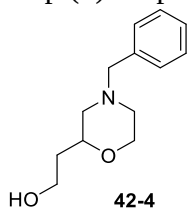


Step (1) Preparation of ethyl 2-(4-phenylmethylmorpholin-2-yl)acetate



2-(phenylmethylamino)ethanol (45.0 g, 298 mmol) and triethylamine (30.0 g, 298 mmol) were dissolved in water (450 mL). The reaction system was heated to 100 °C, added with (*E*)-ethyl-4-bromobut-2-enoate (57.5 g, 298 mmol), reacted overnight, and extracted with ethyl acetate (200 mL × 3). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 10/1) to give **compound 42-3** (27.0 g, 35% yield) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 264.2$.

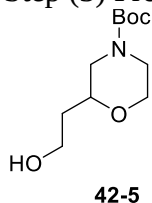
Step (2) Preparation of 2-(4-phenylmethylmorpholin-2-yl)ethanol



Compound 42-3 (27.0 g, 103 mmol) was dissolved in tetrahydrofuran (300 mL). The reaction system was cooled to 0 °C under nitrogen atmosphere. added with lithium aluminum hydride (3.9 g, 103 mmol) in portions, and warmed to room temperature and stirred for 3 h. Then the reaction system was added with sodium sulfate decahydrate to quench the reaction, filtered,

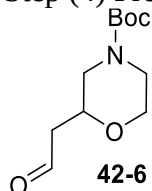
and washed with ethyl acetate, and the filtrate was concentrated. The residue was separated by column chromatography (PE/EA = 2/1) to give **compound 42-4** (17.4 g, 76% yield) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 222.1$.

Step (3) Preparation of *tert*-butyl 2-(2-hydroxyethyl)morpholine-4-carboxylate



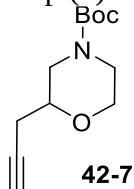
Compound 42-4 (17.6 g, 78.4 mmol) and di-*tert*-butyl dicarbonate (51.3 g, 235 mmol) were dissolved in absolute methanol (180 mL), and palladium on carbon (1.74 g, 10%) was added. The reaction system was purged with hydrogen and stirred at room temperature overnight. Then the reaction system was filtered, and the filtrate was concentrated. The residue was separated by column chromatography (PE/EA = 3/1) to give **compound 42-5** (12.5 g, 68% yield) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 232.3$.

Step (4) Preparation of *tert*-butyl 2-(2-carbonylethyl)morpholine-4-carboxylate



Compound 42-5 (2.0 g, 1.44 mmol) was dissolved in dichloromethane (40 mL), and Dess-Martin periodinane (5.5 g, 2.15 mmol) was added. The reaction system was stirred at room temperature for 2 h. Then the reaction system was added with sodium thiosulfate to quench the reaction, and extracted with dichloromethane (40 mL $\times$ 3). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 2/1) to give **compound 42-6** (1.1 g, 67% yield) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 230.1$.

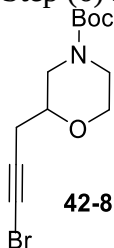
Step (5) Preparation of *tert*-butyl 2-(prop-2-yn-1-yl)morpholine-4-carboxylate



Dimethyl (1-diazo-2-oxopropyl)phosphonate (4.0 g, 20.9 mmol) was dissolved in dichloromethane (40 mL) and acetonitrile (8 mL). The reaction system was stirred at room temperature for 15 min, added with **compound 42-6** (4.0 g, 17.5 mmol) and potassium carbonate (2.4 g, 17.5 mmol), and stirred at room temperature overnight. Then the reaction system was

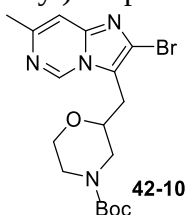
filtered, and the filtrate was concentrated. The residue was separated by column chromatography (PE/EA = 10/1) to give **compound 42-7** (2.1 g, 53% yield) in the form of a colorless oily liquid. LC-MS: $[M-55]^+ = 226.2$.

Step (6) Preparation of *tert*-butyl 2-(3-bromoprop-2-yn-1-yl)morpholine-4-carboxylate



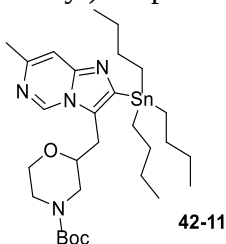
Compound 42-7 (2.1 g, 9.3 mmol) and silver nitrate (1.9 g, 11.1 mmol) were dissolved in acetone (20 mL), and *N*-bromosuccinimide (1.8 g, 10.2 mmol) was added. The reaction system was stirred at room temperature for 4 h, and extracted with ethyl acetate (60 mL $\times$ 3). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 20/1) to give **intermediate 42-8** (2.0 g, 71% yield) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 304.2$.

Step (7) Preparation of *tert*-butyl 2-((2-bromo-7-methylimidazo[1,2-*c*]pyrimidin-3-yl)methyl)morpholine-4-carboxylate



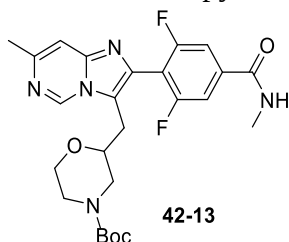
Intermediate 42-8 (2.0 g, 6.5 mmol), 6-methylpyrimidin-4-amine (0.85 g, 7.9 mmol) and copper(II) trifluoromethanesulfonate (0.24 g, 0.65 mmol) were dissolved in acetonitrile (20 mL). The reaction system was stirred at room temperature overnight under oxygen atmosphere, diluted with water (50 mL) and extracted with ethyl acetate (30 mL $\times$ 3). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 2/1) to give **intermediate 42-10** (400 mg, 15% yield) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 411.1$.

Step (8) Preparation of *tert*-butyl 2-((7-methyl-2-(tributylstannyl)imidazo[1,2-*c*]pyrimidin-3-yl)methyl)morpholine-4-carboxylate



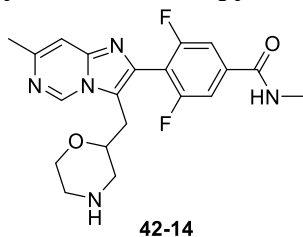
Intermediate 42-10 (40 mg, 0.1 mmol), 1,1,1,2,2,2-hexabutyldistannane (170 mg, 0.3 mmol) and tetrakis(triphenylphosphine)palladium (11 mg, 0.01 mmol) were dissolved in 1,4-dioxane (2 mL). The reaction system was stirred at 100 °C overnight under nitrogen atmosphere, and detected by LC-MS for completion to give **intermediate 42-11** (5% yield). LC-MS: $[M+H]^+$ = 623.5.

Step (9) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*c*]pyrimidin-3-yl)methyl)morpholine-4-carboxylate



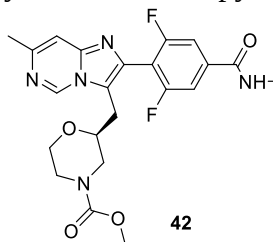
The compound was prepared according to the preparation method for compound 37-9.

Step (10) Preparation of (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*c*]pyrimidin-3-yl)methyl)morpholine



The compound was prepared according to the preparation method as described in step (6) of Example 37.

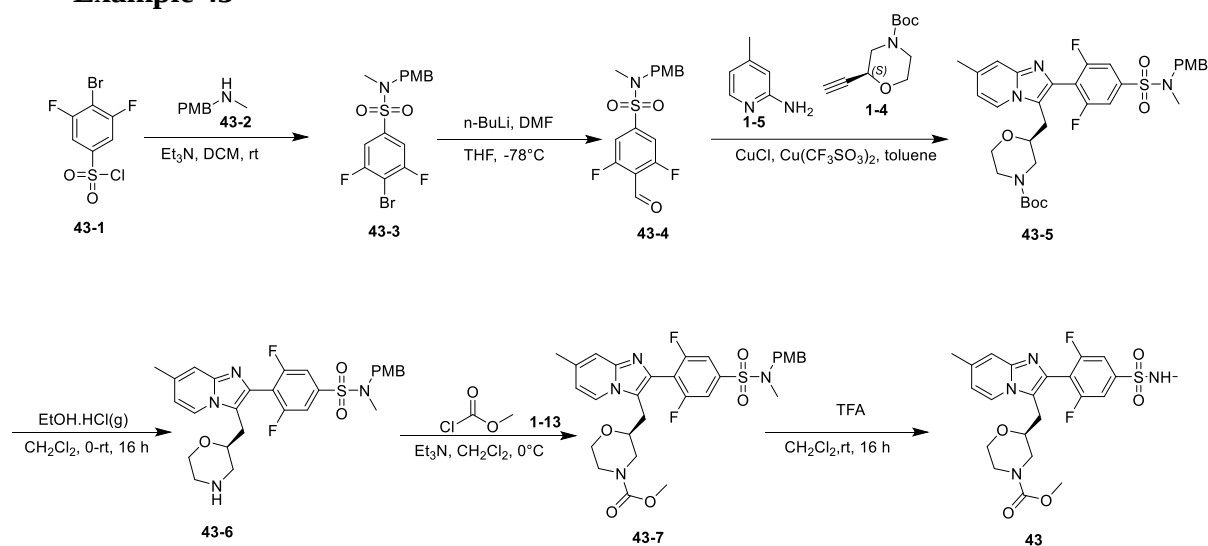
Step (11) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*c*]pyrimidin-3-yl)methyl)morpholine-4-carboxylate



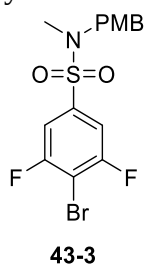
The compound was prepared according to the preparation method as described in step (7) of Example 37.

LC-MS: $[M + H]^+ = 460.2$.

Example 43



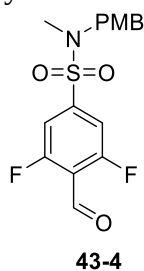
Step (1) Preparation of 4-bromo-3,5-difluoro-*N*-(4-methoxybenzyl)-*N*-methylbenzenesulfonamide



5

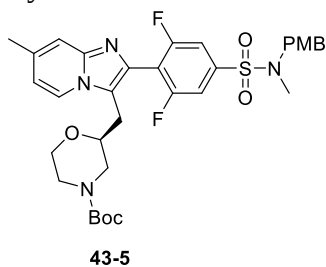
1-(4-methoxyphenyl)-*N*-methylmethanamine (1.30 g, 8.24 mmol) and triethylamine (2.10 g, 20.61 mmol) were dissolved in dichloromethane (16 mL), and a solution of 4-bromo-3,5-difluorobenzenesulfonyl chloride (2.00 g, 6.87 mmol) in dichloromethane (4 mL) was added dropwise at 0 °C. The reaction system was warmed to room temperature and reacted for 4 h. Then the reaction system was added with water (50 mL) to quench the reaction, extracted with dichloromethane (50 mL × 2), washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 43-3** (2.30 g, 78% yield) in the form of a white solid. MS [M+Na] = 427.6, 429.6.

Step (2) Preparation of 3,5-difluoro-4-formyl-*N*-(4-methoxybenzyl)-*N*-methylbenzenesulfonamide



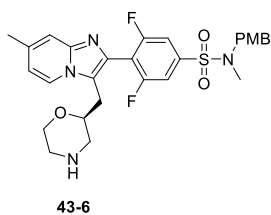
Intermediate 43-3 (1.70 g, 4.19 mmol) was dissolved in anhydrous tetrahydrofuran (17 mL) under nitrogen atmosphere, and *n*-butyllithium (2.4 mol/L, 1.75 mL) was added dropwise at 78 °C. After the completion of the dropwise addition, the reaction system was stirred at this temperature for 0.5 h, added dropwise with *N,N*-dimethylformamide (367 mg, 5.03 mmol), and reacted for 3 h. Then the reaction system was added with diluted hydrochloric acid (0.5 mol/L, 10 mL) to quench the reaction, diluted with water (50 mL) and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 43-4** (600 mg, 32% yield) in the form of a white solid. MS [M+Na] = 377.8.

Step (3) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(*N*-(4-methoxybenzyl)-*N*-methylsulfamoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



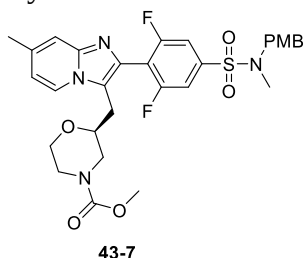
Intermediate 43-4 (600 mg, 1.69 mmol), 4-methylpyridin-2-amine (183 mg, 1.69 mmol) and *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (359 mg, 1.69 mmol) were dissolved in toluene (6 mL) under nitrogen atmosphere, and copper(I) chloride (50 mg, 0.51 mmol) and copper(II) trifluoromethanesulfonate (183 mg, 0.51 mmol) were added. The reaction system was reacted at 85 °C for 10 min, added with dimethylacetamide (0.1 mL), and reacted at 85 °C for another 5 h. Then the reaction system was stirred at room temperature overnight. The resulting reaction system was filtered, and the filtrate was added with water (30 mL) and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 43-5** (298 mg, 27% yield) in the form of a light yellow solid. MS [M+H]⁺ = 656.7.

Step (4) Preparation of (S)-3,5-difluoro-*N*-(4-methoxybenzyl)-*N*-methyl-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)benzenesulfonamide hydrochloride



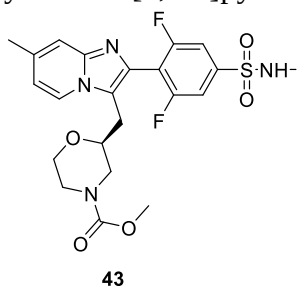
Intermediate 43-5 (298 mg, 0.45 mmol) was dissolved in dichloromethane (5 mL), and a solution of HCl in ethanol (33%, 0.45 mL) was added dropwise. The reaction system was reacted at room temperature for 16 h. Then the reaction system was concentrated under reduced pressure to give **intermediate 43-6** (300 mg, crude product as a hydrochloride) in the form of a light yellow solid. MS $[M+H]^+ = 556.8$.

Step (5) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(N-(4-methoxybenzyl)-N-methylsulfamoyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 43-6 (300 mg, 0.50 mmol) was dissolved in dichloromethane (5 mL) under nitrogen atmosphere, and triethylamine (202 mg, 2.00 mmol) was added at 0 °C, followed by the dropwise addition of a solution of methyl chloroformate (57 mg, 0.60 mmol) in dichloromethane (1 mL). The reaction system was reacted at 0 °C for 4 h. Then the reaction system was added with water (20 mL) to quench the reaction, and extracted with dichloromethane (20 mL $\times$ 2). The organic phases were combined, washed with saturated brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 43-7** (200 mg, 88% purity, 64.5% yield) in the form of a brown solid. MS $[M+H]^+ = 615.2$.

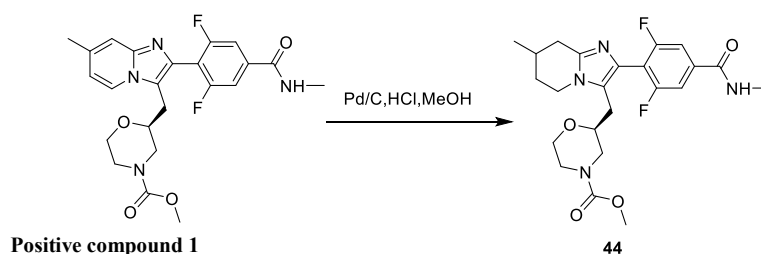
Step (6) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(N-methylsulfamoyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



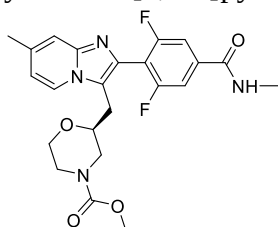
Intermediate 43-7 (200 mg, 0.32 mmol) was dissolved in dichloromethane (3 mL), and trifluoroacetic acid (1 mL) was added at 0 °C. The reaction system was reacted at room temperature for 16 h. Then the reaction system was concentrated under reduced pressure. The residue was dissolved in dichloromethane (40 mL), washed with aqueous sodium bicarbonate solution (10%, 15 mL × 2) and saturated brine (15 mL) sequentially, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative chromatography to give **compound 43** (80.2 mg, 49.8% yield) in the form of a white solid.

MS [M + H]⁺ = 494.7. ¹H NMR (400 MHz, CDCl₃) δ 8.58 (d, J = 5.2 Hz, 1H), 7.93 (s, 1H), 7.61 (d, J = 6.7 Hz, 2H), 7.20 (d, J = 7.1 Hz, 1H), 5.11 - 4.95 (m, 1H), 4.22 - 4.06 (m, 1H), 3.96 - 3.90 (m, 1H), 3.86 - 3.77 (m, 1H), 3.74 (s, 3H), 3.69 - 3.63 (m, 1H), 3.45 - 3.39 (m, 1H), 3.13 - 3.02 (m, 2H), 3.01 - 2.92 (m, 1H), 2.79 (s, 3H), 2.75 - 2.70 (m, 1H), 2.63 (s, 3H).

Example 44



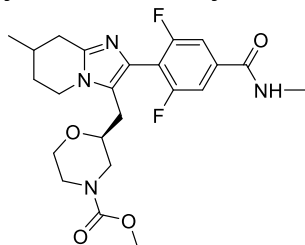
Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-methylcarbamoyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Positive compound 1

The compound was prepared according to patent literature CN105246888A.

Step (2) Preparation of methyl (2S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methyl-5,6,7,8-tetrahydroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



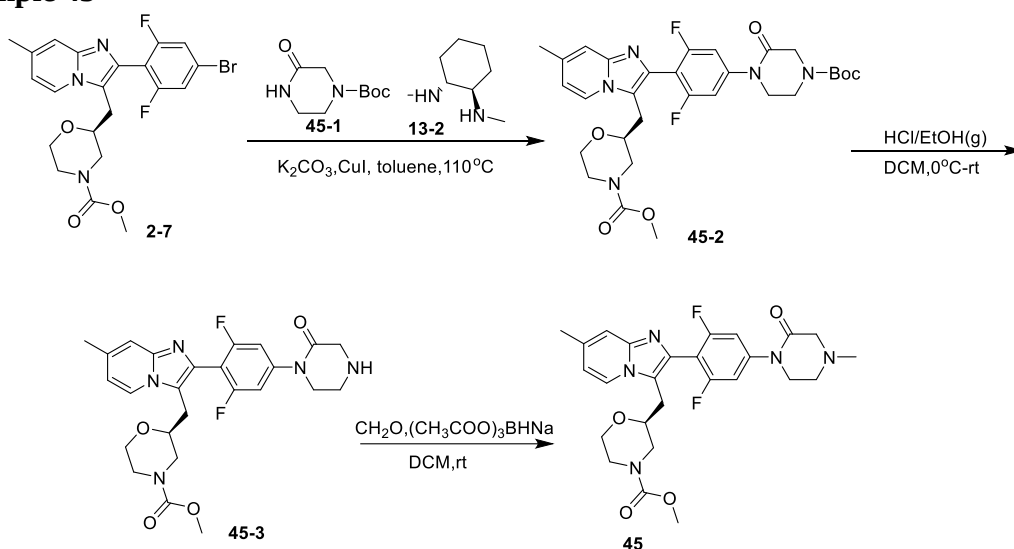
44

Positive compound 1 (120 mg, 0.3 mmol) and concentrated hydrochloric acid (0.1 mL) were dissolved in absolute methanol (3 mL). The reaction system was purged with hydrogen and

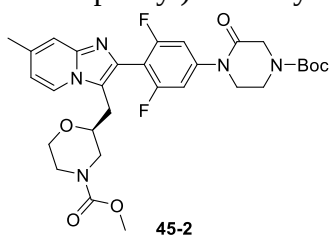
stirred at room temperature for 6 h. Then the reaction system was filtered and concentrated. The residue was diluted with ethyl acetate (50 mL), washed with saturated sodium bicarbonate (15 mL) and saturated brine (15 mL) sequentially, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography (DCM/MeOH = 40/1) to give **compound 44** (61 mg, 50% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 463.2$. ^1H NMR (400 MHz, CDCl_3): δ 7.48 (brs, 2H), 4.30 – 4.23 (m, 1H), 4.12 – 4.04 (m, 1H), 3.99 – 3.69 (m, 4H), 3.68 (s, 3H), 3.40 – 3.30 (s, 2H), 3.20 – 3.10 (m, 1H), 3.03 (d, $J = 3.5$ Hz, 3H), 2.92 – 2.80 (m, 1H), 2.74 – 2.62 (m, 2H), 2.60 – 2.44 (m, 2H), 2.18 – 2.06 (m, 2H), 1.80 – 1.68 (m, 1H), 1.20 (d, $J = 5.8$ Hz, 3H).

Example 45



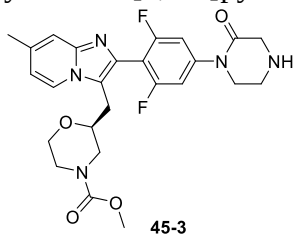
Step (1) Preparation of methyl (S)-2-((2-(4-(4-(*tert*-butoxycarbonyl)-2-oxopiperazin-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)-methyl)morpholine-4-carboxylate



Intermediate 2-7 (250 mg, 0.52 mmol) was dissolved in anhydrous toluene (3 mL) and *tert*-butyl 3-carboxylpiperazine-1-carboxylate (125 mg, 0.62 mmol), potassium carbonate (216 mg, 1.56 mmol), *trans*-(1*R*,2*R*)-*N,N'*-dimethyl-1,2-cyclohexanediamine (7.4 mg, 0.052 mmol) and copper(I) iodide (10 mg, 0.052 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 110 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature, and filtered, and the filtrate was added with water (10 mL), and extracted with dichloromethane (30 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL $\times$ 1), dried over magnesium sulfate, filtered and concentrated by rotary

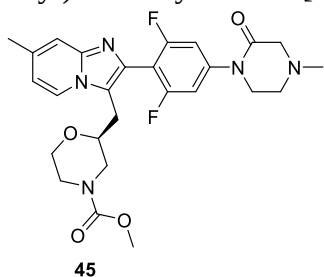
evaporation to give a crude product. The crude product was purified by column chromatography (4 g of silica gel column, petroleum ether:ethyl acetate = 1:1) to give **intermediate 45-2** in the form of a yellow oil (200 mg, 64% yield), LC-MS: $[M+H]^+ = 600.9$.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(2-oxopiperazin-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

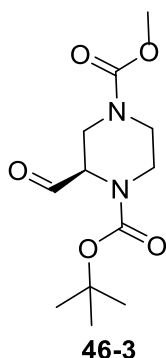


Intermediate 45-2 (200 mg, 0.33 mmol) was dissolved in dichloromethane (5 mL), and a solution of HCl in ethanol (33%, 0.5 mL) was added dropwise with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. After the reaction was completed, the reaction system was added dropwise with saturated sodium bicarbonate solution to adjust the pH to 8 with the temperature maintained at 0 °C, added with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 1), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 45-3** (175 mg) in the form of a yellow oil, LC-MS: $[M+H]^+ = 500.9$.

Step (3) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-methyl-2-oxopiperazin-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

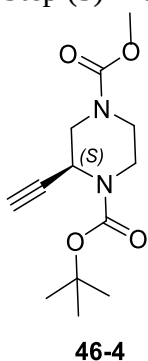


Intermediate 45-3 (175 mg, 0.35 mmol) and aqueous formaldehyde solution (37%, 30 mg, 0.364 mmol) were dissolved in dichloromethane (3 mL), and sodium triacetoxyborohydride (287 mg, 1.36 mmol) was slowly added. The reaction system was reacted at room temperature for 3 h. After the reaction was completed, the reaction system was added with water (10 mL) and extracted with dichloromethane (15 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 1), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by column chromatography to give **compound 45** (74 mg, 41% yield) in the form of a yellow solid.



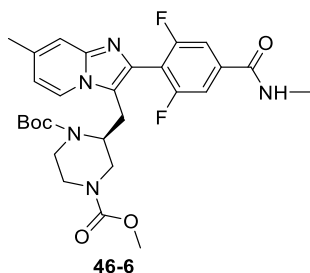
Intermediate 46-2 (1.7 g, 6.0 mmol) was dissolved in dichloromethane (30 mL). Under nitrogen atmosphere, the reaction system was cooled to 0 °C, added with Dess-Martin periodinane (3.9 g, 9.0 mmol), and warmed to room temperature and stirred for 2 h. Then the reaction system was extracted with dichloromethane (30 mL × 3). The organic phase was washed with saturated brine (50 mL × 2), dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 2/1) to give **intermediate 46-3** (360 mg, 21% yield) in the form of a colorless oily liquid. LC-MS: [M-55]⁺ = 217.1.

Step (3) Preparation of 1-*tert*-butyl-4-methyl (S)-2-ethynylpiperazine-1,4-dicarboxylate



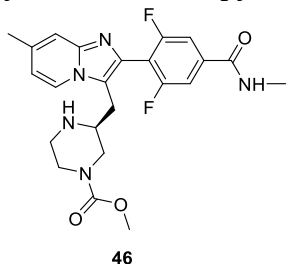
Compound 1-3 (305 mg, 1.6 mmol) and potassium carbonate (365 mg, 2.6 mmol) were dissolved in acetonitrile (2.5 mL) and absolute methanol (0.5 mL). The reaction system was stirred at room temperature for 15 min, added with a dissolved solution of **intermediate 46-3** (360 mg, 1.3 mmol) in acetonitrile (2.5 mL) and absolute methanol (0.5 mL), and stirred at room temperature overnight. Then the reaction system was filtered and concentrated. The residue was separated by column chromatography (PE/EA = 10/1) to give **intermediate 46-4** (165 mg, 46% yield) in the form of a white oily liquid. LC-MS: [M-55]⁺ = 213.7.

Step (4) Preparation of methyl (S)-2-((2-(2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)-1-*tert*-butyl carboxylate-piperazine-4-carboxylate



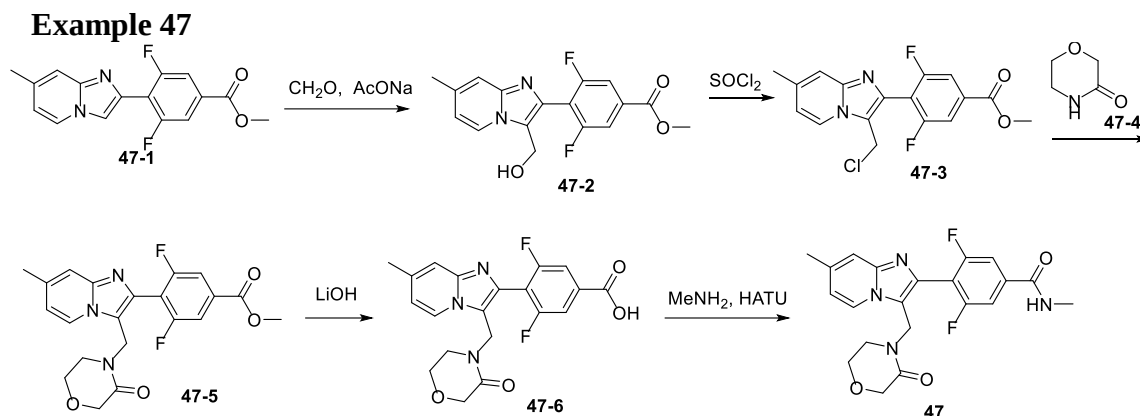
Intermediate 46-4 (165 mg, 0.6 mmol), 4-methylpyridin-2-amine (66 mg, 0.6 mmol), 3,5-difluoro-4-formyl-*N*-methylbenzamide (123 mg, 0.6 mmol), copper(I) chloride (18 mg, 0.2 mmol) and copper(II) trifluoromethanesulfonate (67 g, 0.2 mmol) were dissolved in anhydrous toluene (3 mL). The reaction system was heated to 85 °C, added with *N,N*-dimethylacetamide (0.2 mL), and reacted at 85 °C for 5 h. Then the reaction system was cooled to room temperature and reacted overnight, and extracted with dichloromethane (15 mL × 3). The organic phase was washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (DCM/MeOH = 40/1) to give **intermediate 46-6** (105 mg) in the form of a brown solid. LC-MS: $[M+H]^+ = 558.2$.

Step (5) Preparation of methyl (S)-2-((2-(2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)piperazine-4-carboxylate

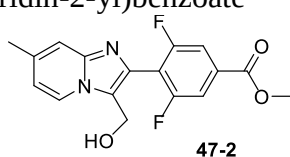


Intermediate 46-6 (105 mg, 0.2 mmol) was dissolved in anhydrous dichloromethane (2 mL). The reaction system was cooled to 0 °C, added dropwise with a solution of HCl in ethanol (0.5 mL), and warmed to room temperature and stirred for 1 h. Then the reaction system was concentrated. The residue was separated by preparative reverse phase chromatography to give **compound 46** (3.5 mg) in the form of a white solid.

LC-MS: $[M+H]^+ = 458.5$. ^1H NMR (400 MHz, MeOD) : δ 8.75 (brs, 1H), 7.84 (m, 3H), 7.42 (brs, 1H), 4.10 – 4.02 (m, 2H), 3.70 - 3.66 (m, 1H), 3.66 (s, 3H), 3.57 – 3.37 (m, 3H), 3.20 – 3.10 (m, 2H), 2.98(s, 3H), 2.95 -2.85 (m, 1H), 2.64(s, 3H).

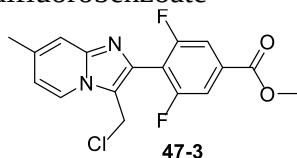


Step (1) Preparation of methyl 3,5-difluoro-4-(3-(hydroxymethyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)benzoate



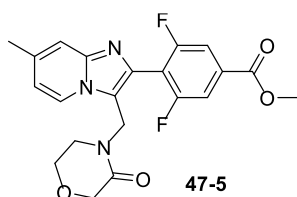
Intermediate 47-1 (synthesized according to the preparation method for intermediate 31-7, LC-MS $[M+H]^+$: 303.18) (1.2 g, 4.0 mmol) was dissolved in acetonitrile (8 mL), and acetic acid (8 mL), aqueous formaldehyde solution (0.25 mL, 24 mmol) and sodium acetate (1.31 g, 16 mmol) were added. The reaction system was heated to 50 °C and reacted for 16 h. Then the reaction system was concentrated by rotary evaporation to remove the solvent. The residue was diluted with dichloromethane (50 mL), and washed with saturated aqueous sodium bicarbonate solution (20 mL $\times$ 2). The organic phases were combined, dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (DCM/MeOH = 50/1) to give **intermediate 47-2** (945 mg, 71% yield) in the form of a yellow solid. LC-MS: $[M+H]^+$ = 333.2.

Step (2) Preparation of methyl 4-(3-(chloromethyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)-3,5-difluorobenzoate



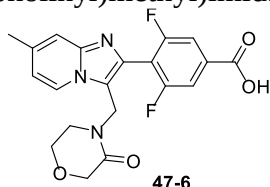
Intermediate 47-2 (1.2 g, 4.0 mmol) was dissolved in dichloromethane (15 mL), and thionyl chloride (4 mL) was added at 0 °C. The reaction system was reacted at 0 °C for 3 h. Then the reaction system was concentrated by rotary evaporation to remove the solvent to give **intermediate 47-2** (900 mg) in the form of a yellow solid. LC-MS: $[M+H]^+$ = 336.1.

Step (3) Preparation of methyl 3,5-difluoro-4-(7-methyl-3-((3-oxo-4-morpholinyl)methyl)imidazo[1,2-*a*]pyridin-2-yl)benzoate



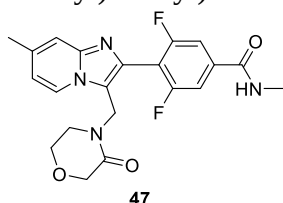
Morpholin-3-one (101 mg, 1.0 mmol) was dissolved in DMF (5 mL), and sodium hydride (80 mg, 2 mol) was added in an ice bath. The reaction system was reacted for 0.5 h, added with methyl **intermediate 47-4** (351 mg, 1 mmol), and reacted at room temperature for 16 h. Then the reaction system was poured into ice water (10 mL), extracted with dichloromethane (10 mL × 2), washed with saturated brine (10 mL), dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated by rotary evaporation to give **intermediate 47-5** (120 mg) in the form of a yellow solid. LC-MS: $[M+H]^+ = 416.4$.

Step (4) Preparation of 3,5-difluoro-4-(7-methyl-3-((3-oxo-4-morpholinyl)methyl)imidazo[1,2-a]pyridin-2-yl)benzoic acid



Intermediate 47-5 (120 mg, 0.29 mmol) was dissolved in THF (9 mL), and MeOH (3 mL), H₂O (3 mL), and lithium hydroxide (61 mg, 1.45 mmol) were added sequentially. The reaction system was stirred at room temperature overnight, then diluted with water (10 mL), and concentrated by rotary evaporation under reduced pressure to remove the organic solvent. The aqueous phase was adjusted to pH 5-6 with hydrochloric acid (1 N), and extracted with dichloromethane (10 mL × 2). The organic phase was dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated by rotary evaporation. The residue was purified by silica gel column chromatography (DCM/MeOH = 15:1) to give **intermediate 47-6** (85 mg, 72.4% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 402.3$.

Step (5) Preparation of 3,5-difluoro-N-methyl-4-(7-methyl-3-((3-oxo-4-morpholinyl)methyl)imidazo[1,2-a]pyridin-2-yl)benzamide

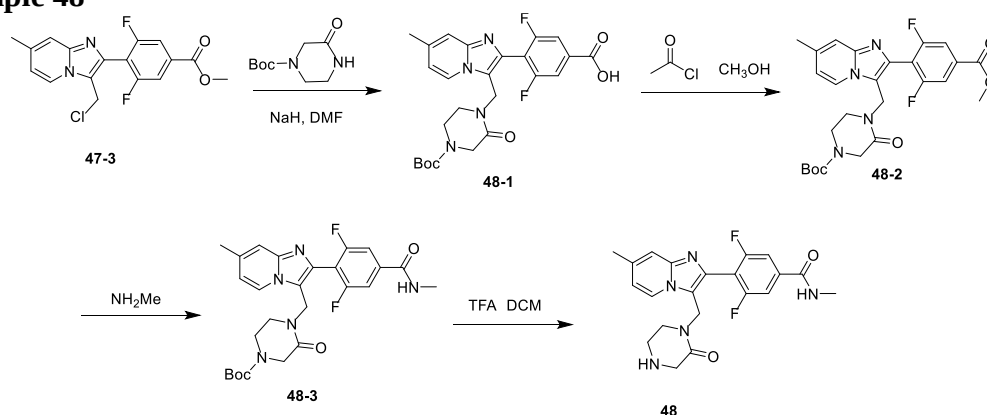


Intermediate 47-6 (80 mg, 0.2 mmol) was dissolved in DMF (5 mL), and methylamine hydrochloride (20 mg, 0.3 mmol), 2-(7-azabenzotriazol-1-yl)-N,N,N',N'-tetramethyluronium hexafluorophosphate (114 mg, 0.3 mmol) and triethylamine (44 mg, 0.4 mmol) were added

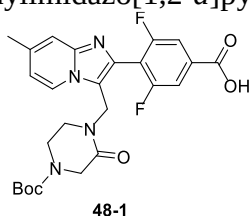
sequentially. The reaction system was stirred at room temperature overnight. Then the reaction system was diluted with water (15 mL), extracted with ethyl acetate (10 mL × 2), washed with saturated brine (10 mL), dried over anhydrous sodium sulfate and filtered, and the filtrate was concentrated by rotary evaporation. The residue was purified by silica gel column chromatography (DCM/MeOH = 30:1) to give **compound 47** (21 mg, 20% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 415.1$. ^1H NMR (400 MHz, MeOD) : δ 8.44 (d, $J = 6.9$ Hz, 1H), 7.64 (d, $J = 7.8$ Hz, 2H), 7.43 (s, 1H), 6.96 (d, $J = 6.9$ Hz, 1H), 5.00 (s, 2H), 4.14-4.12 (m, 2H), 3.75-3.71 (m, 2H), 3.04-3.01 (m, 2H), 2.98 (s, 3H), 2.47 (s, 3H).

Example 48

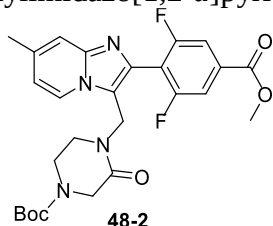


Step (1) Preparation of 4-(3-((4-(*tert*-butoxycarbonyl)-2-oxopiperazin-1-yl)methyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)-3,5-difluorobenzoic acid



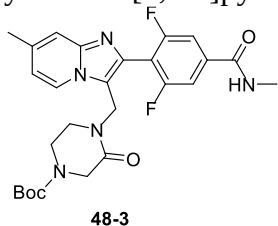
Tert-butyl 3-carboxypiperazine-carboxylate (400 mg, 2.0 mmol) was dissolved in *N,N*-dimethylformamide (10 mL). The reaction system was cooled in an ice water bath for 5 min, added with sodium hydride (120 mg, 3.0 mmol) slowly, and reacted for 30 min. Then the reaction system was added with **intermediate 47-3** (350 mg, 1.0 mmol) (60% purity shown on LC-MS) and stirred overnight. After the reaction was quenched with saturated ammonium chloride solution (5 mL), the reaction system was added dropwise with hydrochloric acid (1 N) to adjust the pH to 3-4, diluted with water (20 mL), and extracted with dichloromethane extraction (45 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by column chromatography (DCM/CH₃OH = 10/1) to give **intermediate 48-1** (300 mg, 50% purity shown on LC-MS) in the form of a light yellow solid. LC-MS: $[M+H]^+ = 501.2$.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-carbomethoxyphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)-2-carboxylpiperazine-carboxylate



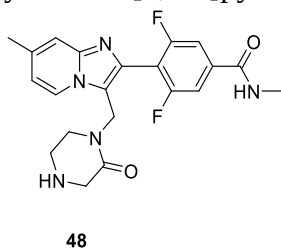
Intermediate 48-1 (300 mg, 0.69 mmol) was dissolved in methanol (5 mL). The reaction system was added dropwise with acetyl chloride (0.5 mL), and stirred overnight. Then the reaction system was concentrated, diluted with dichloromethane (25 mL), and washed with saturated sodium bicarbonate solution (15 mL). The aqueous phase was extracted with dichloromethane (20 mL × 2). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by column chromatography (PE/EA = 1/1) to give **intermediate 48-2** (70 mg, 60% purity shown on LC-MS) in the form of a colorless oily liquid, LC-MS: $[M+H]^+ = 515.1$.

Step (3) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)-2-carboxylpiperazine-carboxylate



Intermediate 48-2 (70 mg, 0.13 mmol) was dissolved in ethanol (5 mL), and aqueous methylamine solution (2 mL, 25-30% w/w) was added. The reaction system was stirred at room temperature overnight. Then the reaction system was added with H₂O (15 mL) to quench the reaction, and extracted with ethyl acetate (15 mL × 2). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give **intermediate 48-3** (60 mg, 60% purity shown on LC-MS) in the form of a colorless oily liquid. LC-MS: $[M+H]^+ = 514.1$.

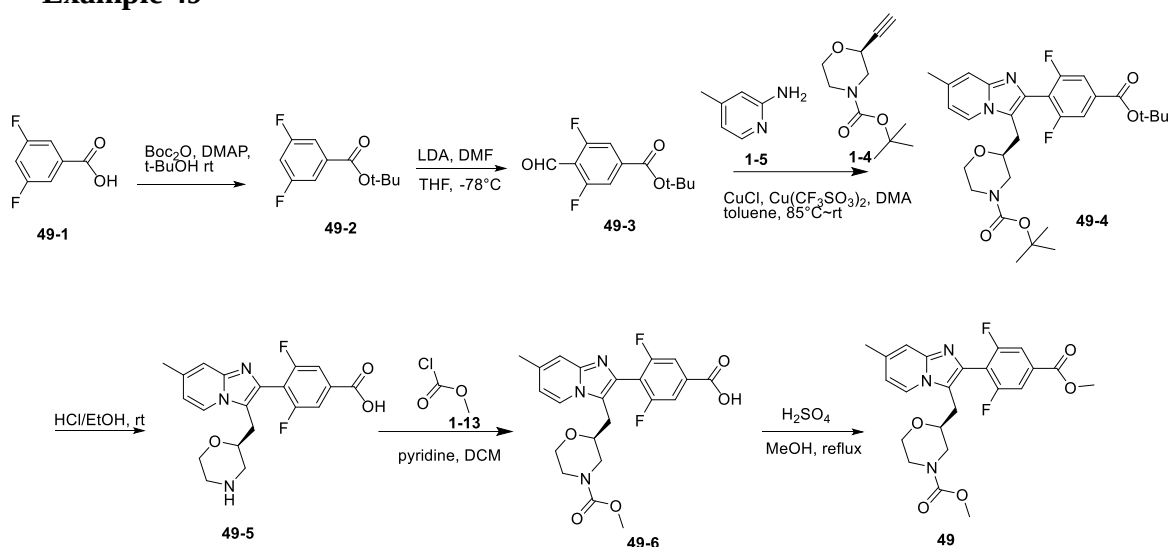
Step (4) Preparation of (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)-2-carboxyl-piperazine



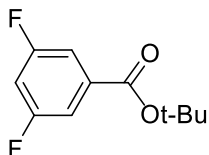
Intermediate 48-3 (60 mg, 0.12 mmol) was dissolved in dichloromethane (3 mL), and trifluoroacetic acid (3 mL) was added. The reaction system was stirred overnight at room temperature, concentrated by rotary evaporation, diluted with dichloromethane (40 mL), washed with saturated sodium bicarbonate solution (10 mL), washed with saturated brine (10 mL), and dried over anhydrous sodium sulfate. The residue was purified by preparative chromatography, and lyophilized to give **compound 48** (3 mg) in the form of a white solid.

LC-MS: $[M+H]^+ = 414.1$. ^1H NMR (400 MHz, MeOD) : δ 8.47 (d, $J = 7.2$ Hz, 1H), 7.64 (d, $J = 7.8$ Hz, 2H), 7.42 (s, 1H), 6.95 (d, $J = 7.2$ Hz, 2H), 4.99 (s, 2H), 3.42 (s, 2H), 3.00 – 2.95 (m, 5H), 2.87 (t, $J = 5.2$ Hz, 2H), 2.49 (s, 3H).

Example 49



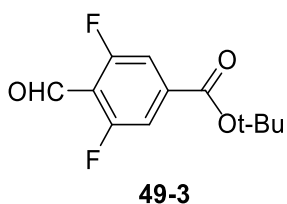
Step (1) Preparation of *tert*-butyl 3,5-difluorobenzoate



49-2

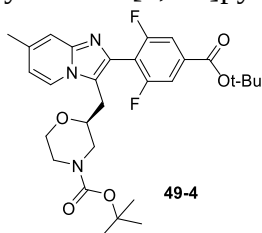
3,5-difluorobenzoic acid (10 g, 63.25 mmol) was dissolved in *tert*-butanol (100 mL), and DMAP (12.3 g, 18.975 mmol) and Boc-anhydride (27.6 g, 126.5 mmol) were added. The reaction system was reacted at room temperature for 15 h. Then the reaction system was concentrated, diluted with water (150 mL) and extracted with ethyl acetate (50 mL $\times$ 3). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by column chromatography (PE/EA = 20/1) to give **compound 49-2** (12 g, 88% yield) in the form of a colorless oily liquid.

Step (2) Preparation of *tert*-butyl 3,5-difluoro-4-formylbenzoate



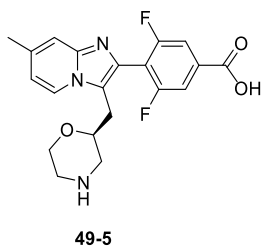
LDA (2 M, 8.4 mL) was dissolved in tetrahydrofuran (10 mL). The reaction system was cooled to -78 °C under nitrogen atmosphere. **Compound 49-2** (3 g, 14 mmol) was dissolved in tetrahydrofuran (20 mL), and the solution was added to the reaction system. The reaction system was stirred at -78 °C for 1.5 h, added with DMF (1.228 g, 16.8 mmol), and stirred for another 1 h with the temperature maintained not higher than -70 °C. Then the reaction system was added with saturated aqueous ammonium chloride solution (60 mL) to quench the reaction, and extracted with ethyl acetate (30 mL × 3). The ethyl acetate phases were combined, dried over anhydrous sodium sulfate, and concentrated. The residue was separated by column chromatography (PE/EA = 10/1) to give **compound 49-3** (1.5 g, 44% yield) in the form of a yellow solid.

Step (3) Preparation of *tert*-butyl (S)-2-((2-(4-(*tert*-butoxycarbonyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



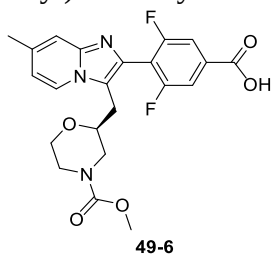
Compound 49-3 (480 mg, 1.98 mmol) was dissolved in anhydrous toluene (4 mL), and **compound 1-4** (419 mg, 1.98 mmol), **compound 1-5** (215 mg, 1.98 mmol), copper(I) chloride (62 mg, 0.63 mmol), and copper(II) trifluoromethanesulfonate (216 mg, 0.60 mmol) were added. Under nitrogen atmosphere, the reaction system was heated to 85 °C, added with two drops of *N,N*-dimethylacetamide, heated and stirred for 6 h. Then the reaction system was cooled to room temperature and stirred overnight. Then the reaction system was filtered, and the filtrate was concentrated. The residue was separated by column chromatography (PE/EA = 2/1) to give **intermediate 49-4** (340 mg, 31% yield) in the form of a brown solid. LC-MS: $[M+H]^+ = 544.7$.

Step (4) Preparation of (S)-3,5-difluoro-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)benzoic acid



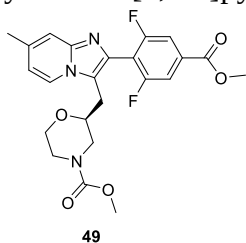
Intermediate 49-4 (340 mg, 0.62 mmol) was dissolved in ethanol (3 mL), and a solution of HCl in ethanol (1 mL, 33%) was added. The reaction system was stirred at room temperature for 1.5 h. Then the reaction system was concentrated to give **intermediate 49-5** (400 mg) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 388.7$.

Step (5) Preparation of (S)-3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)benzoic acid



Intermediate 49-5 (400 mg) was dissolved in dichloromethane (5 mL), and pyridine (244 mg, 3.09 mmol) was added, followed by the addition of methyl chloroformate (62 mg, 0.65 mmol). The reaction system was stirred at room temperature for 1.5 h, and concentrated to give **intermediate 49-6** (190 mg). LC-MS: $[M+H]^+ = 446.6$.

Step (6) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-carbomethoxyphenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate

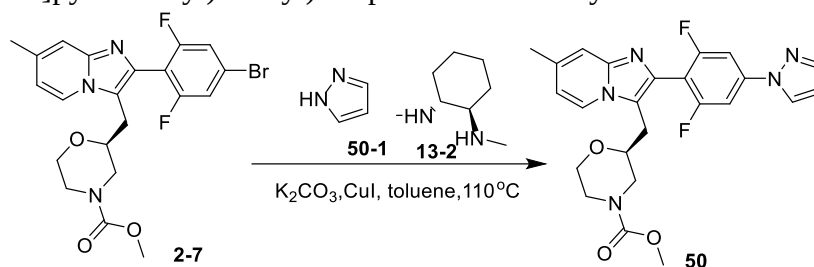


Intermediate 49-6 (50 mg, 0.1124 mmol) was dissolved in absolute methanol (5 mL), and concentrated sulfuric acid (0.1 mL) was added. The reaction system was stirred under reflux overnight. Then the reaction system was concentrated, diluted with water (10 mL), and extracted with ethyl acetate (10 mL $\times$ 2). The organic phase was dried over anhydrous sodium sulfate and concentrated. The residue was separated by thin-layer chromatography (PE/EA = 3/1) to give **compound 49** (20.8 mg, 40% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 460.7$. ^1H NMR (400 MHz, CDCl_3) : δ 8.27 (brs, 1H), 7.73 (d, $J = 7.1$ Hz, 2H), 7.54 (brs, 1H), 6.80 (brs, 1H), 4.00 (s, 3H), 3.99 -3.75 (m, 3H), 3.70 (s, 3H), 3.65 – 3.56 (m, 1H), 3.45 – 3.35 (m, 1H), 3.10 – 3.85 (m, 3H), 2.67 – 2.57 (m, 1H), 2.48 (s, 3H).

Example 50

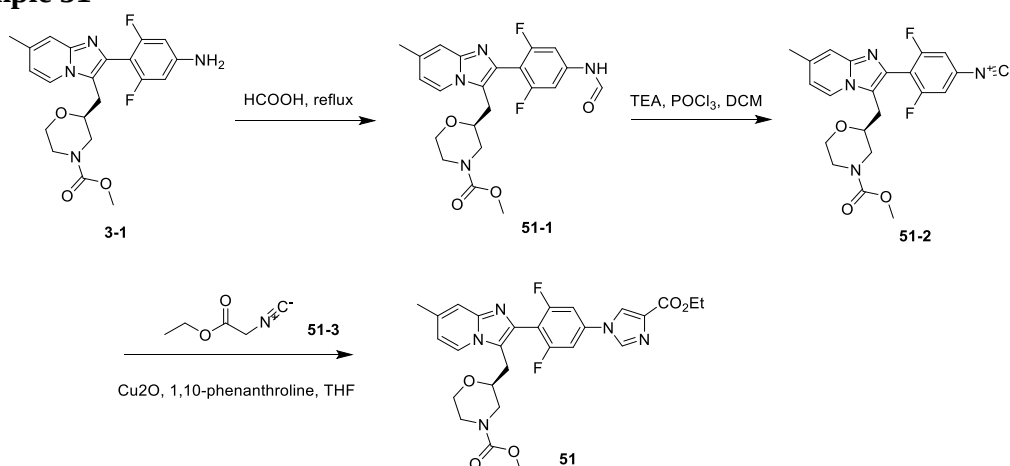
5 Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1*H*-pyrazole)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



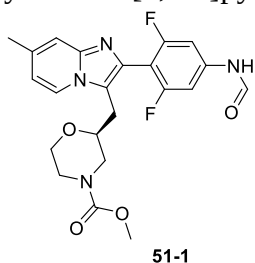
Intermediate 2-7 (120 mg, 0.25 mmol) was dissolved in anhydrous toluene (3 mL), and potassium carbonate (104 mg, 0.75 mmol), copper(I) iodide (4.8 mg, 0.025 mmol), 1*H*-pyrazole (34 mg, 0.50 mmol) and *trans*-(1*R*,2*R*)-*N,N'*-dimethyl-1,2-cyclohexanediamine (3.6 mg, 0.025 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 110 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (10 mL) and extracted with dichloromethane (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL $\times$ 1), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 50** (18.7 mg, 16% yield) in the form of a yellow solid.

LC-MS: $[M+H]^+ = 468.7$. ^1H NMR (400 MHz, MeOD): δ 8.44 – 8.40 (m, 2H), 7.80 (s, 1H), 7.67 (d, $J = 8.1$ Hz, 2H), 7.35 (s, 1H), 6.88 (d, $J = 7.0$ Hz, 1H), 6.61 (s, 1H), 3.89 – 3.76 (m, 3H), 3.66 (s, 3H), 3.65 – 3.56 (m, 1H), 3.42 – 3.36 (m, 1H), 3.15 – 3.06 (m, 2H), 2.96 – 2.80 (m, 1H), 2.72 – 2.60 (m, 1H), 2.47 (s, 3H).

Example 51



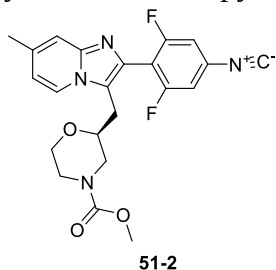
Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-formylaminophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



5

Intermediate 3-1 (360 mg, 0.86 mmol) was dissolved in formic acid (5 mL). The reaction system was reacted at 90 °C for 2 h. After the reaction was completed as monitored by TLC, the reaction system was concentrated under reduced pressure, added with saturated brine (10 mL), adjusted to pH 7-8 with saturated sodium bicarbonate solution, and extracted with DCM (10 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 51-1** (220 mg, 52% yield) in the form of a yellow solid. MS $[M+H]^+ = 445.1$.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-isocyanophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

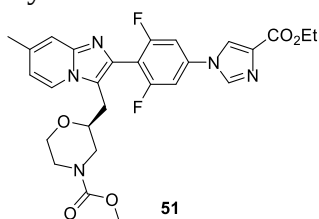


15

Intermediate 51-1 (220 mg, 0.49 mmol) and triethylamine (149 mg, 1.5 mmol) were dissolved in dichloromethane (25 mL) and phosphorus oxychloride (115 mg, 0.75 mmol) was added dropwise slowly at 0 °C. After the completion of the dropwise addition, the reaction

system was reacted at 0 °C for 2 h. After the reaction was completed as monitored by LCMS, the reaction system was added with water at 0 °C to quench the reaction, with the temperature maintained at 25 °C below. Then the reaction system was adjusted to pH 7-8 with saturated sodium bicarbonate solution, and washed with saturated saline (10 mL). The organic phase was dried, and concentrated to give **intermediate 51-2** (210 mg, 84% yield, crude product) in the form of a yellow oily liquid, which was used directly in the next step without purification. MS $[M+H]^+ = 427.0$.

Step (3) Preparation of methyl (S)-2-((2-(4-(4-(carbethoxy<ethoxycarbonyl>)-1H-imidazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



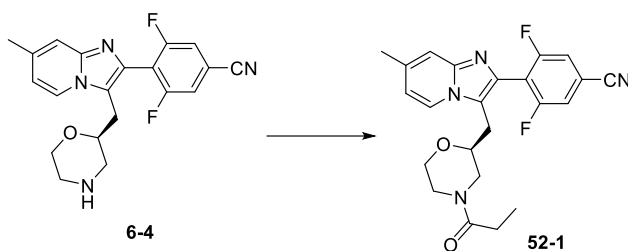
Intermediate 51-2 (170 mg, 0.4 mmol), ethyl isocyanoacetate (45 mg, 0.4 mmol), copper(I) chloride (8.6 mg, 0.06 mmol) and 1,10-phenanthroline (22 mg, 0.12 mmol) were dissolved in tetrahydrofuran (15 mL). The reaction system was reacted at 60 °C for 3 h under nitrogen atmosphere, and reacted at room temperature overnight. Then the reaction system was added with saturated brine (20 mL), and extracted with ethyl acetate (10 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by FCC to give **compound 51** (160 mg, 70% yield) in the form of a white solid. MS $[M+H]^+ = 540.1$.

¹H NMR (400 MHz, d6-DMSO) δ 8.70 (d, $J = 1.3$ Hz, 1H), 8.60 (d, $J = 1.3$ Hz, 1H), 8.44 (d, $J = 7.1$ Hz, 1H), 7.89 (d, $J = 8.6$ Hz, 2H), 7.37 (s, 1H), 6.85 (dd, $J = 7.1, 1.5$ Hz, 1H), 4.30 (q, $J = 7.1$ Hz, 2H), 3.77 (d, $J = 12.2$ Hz, 1H), 3.66 (t, $J = 12.1$ Hz, 2H), 3.56 (s, 3H), 3.53 - 3.47 (m, 1H), 3.29 - 3.21 (m, 1H), 3.10 - 3.00 (m, 2H), 2.90 - 2.74 (m, 1H), 2.60 - 2.45 (m, 1H), 2.40 (s, 3H), 1.32 (t, $J = 7.1$ Hz, 3H).

Example 52

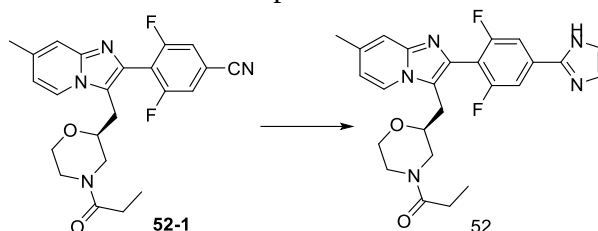
Step (1)

Compound 52-1 was prepared by replacing methyl chloroformate with acetyl chloride, according to the preparation method for **compound 6** in Example 6.



Step (2)

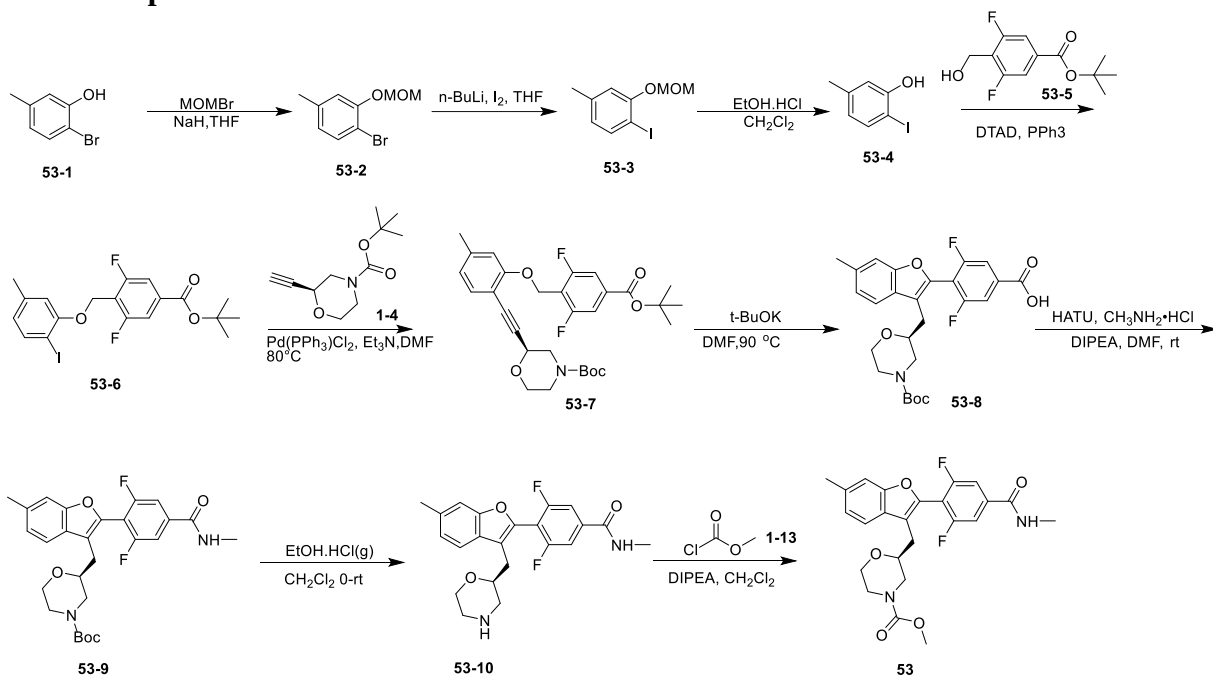
Compound 52 was prepared by using **compound 52-1** as the starting material, according to the preparation method as described in Example 7.



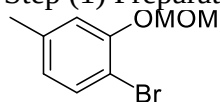
5

LC-MS: $[M + H]^+ = 466.1$.

Example 53



Step (1) Preparation of 1-bromo-2-(methoxymethoxy)-4-methylbenzene

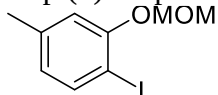


10

2-bromo-5-methylphenol (1.6 g, 8.55 mmol) was dissolved in tetrahydrofuran (20 mL), and sodium hydride (855 mg, 21.37 mmol) was added in portions under nitrogen atmosphere in an ice water bath. The reaction system was reacted at 0 °C for 30 min, added dropwise with bromomethyl methyl ether (1.4 g, 11.12 mmol), and stirred at room temperature overnight. Then

the reaction system was poured into ice water (20 mL), and extracted with ethyl acetate (20 mL $\times$ 2). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 53-2** (1.7 g, 68% yield) in the form of a colorless oil.

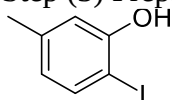
Step (2) Preparation of 1-iodo-2-(methoxymethoxy)-4-methylbenzene



53-3

Intermediate 53-2 (528 mg, 2.29 mmol) was dissolved in tetrahydrofuran (10 mL) under nitrogen atmosphere, and *n*-butyllithium (1.2 mL, 2.75 mmol) was added dropwise at -78 °C. The reaction system was reacted at this temperature for 20 min, added dropwise with a solution of iodine (874 mg, 3.44 mmol) in tetrahydrofuran (5 mL), and reacted for 2 h. Then the reaction system added with ice water (20 mL) to quench the reaction, added with saturated sodium thiosulfate solution (2 mL), stirred at room temperature for 5 min, and extracted with ethyl acetate (25 mL $\times$ 2). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give **intermediate 53-3** (570 mg, 80% purity, 72% yield, crude product) in the form of a brown oil.

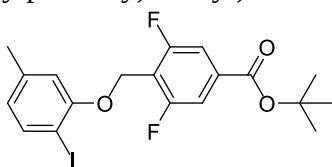
Step (3) Preparation of 2-iodo-5-methylphenol



53-4

Intermediate 53-3 (570 mg, 2.05 mmol) was dissolved in tetrahydrofuran (8 mL), and a solution of HCl in ethanol (1.03 mL, 4.1 mmol) was added dropwise at 0 °C. The reaction system was stirred at room temperature for 2 h and concentrated under reduced pressure. The residue was purified by silica gel column chromatography to give **intermediate 53-4** (400 mg, 70% purity, 58% yield) in the form of a colorless oil.

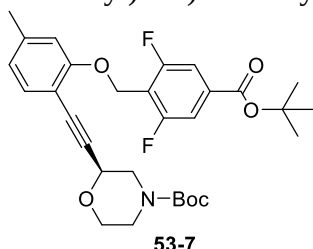
Step (4) Preparation of *tert*-butyl 3,5-difluoro-4-((2-iodo-5-methylphenoxy)methyl)benzoate



53-6

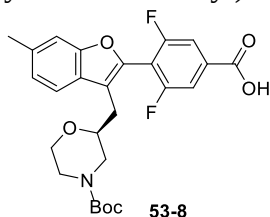
Intermediate 53-4 (400 mg, 1.71 mmol), triphenylphosphine (673 mg, 2.57 mmol) and *tert*-butyl 3,5-difluoro-4-(hydroxymethyl)benzoate (418 mg, 1.71 mmol) were dissolved in tetrahydrofuran (6 mL) under nitrogen atmosphere, and di-*tert*-butyl azodicarboxylate (591 mg, 2.57 mmol) was added. The reaction system was reacted at room temperature overnight. Then the reaction system was concentrated under reduced pressure, added with ethyl acetate (30 mL), washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 53-6** (375 mg, 75% purity, 37% yield) in the form of a yellow oil. ESI-MS [M+Na]⁺: 462.5.

Step (5) Preparation of *tert*-butyl (S)-2-((2-((4-(*tert*-butoxycarbonyl)-2,6-difluorobenzyl)oxo)-4-methylphenyl)ethynyl)morpholine-4-carboxylate



Intermediate 53-6 (290 mg, 0.63 mmol) and *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (146 mg, 0.69 mmol) were dissolved in *N,N*-dimethylformamide (1.5 mL) under nitrogen atmosphere, and copper(I) iodide (12 mg, 0.063 mmol), bis(triphenylphosphine)palladium(II) chloride (91 mg, 0.13 mmol) and triethylamine (4 mL) were added. The reaction system was reacted at 80 °C for 8 h, cooled to room temperature, added with water (40 mL), and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 53-7** (132 mg, 75% purity, 29% yield) in the form of a brown solid. ESI-MS [M+Na]⁺: 565.6.

Step (6) Preparation of (S)-4-(3-((4-(*tert*-butoxycarbonyl)morpholin-2-yl)methyl)-6-methylbenzofuran-2-yl)-3,5-difluorobenzoic acid

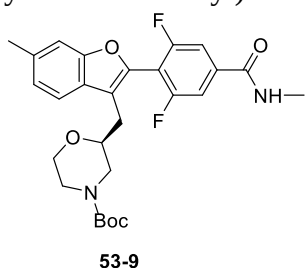


Intermediate 53-7 (132 mg, 0.24 mmol) was dissolved in *N,N*-dimethylformamide (3 mL) under nitrogen atmosphere, and potassium *tert*-butoxide (96 mg, 0.72 mmol) was added. The

reaction system was reacted at 80 °C for 3 h. Then the reaction system was cooled to room temperature, poured into water (10 mL), adjusted to pH 4 with diluted hydrochloric acid (1 N), and extracted with ethyl acetate (20 mL × 2). The organic phases were combined, washed with saturated brine (15 mL × 2), dried over anhydrous magnesium sulfate, filtered and concentrated.

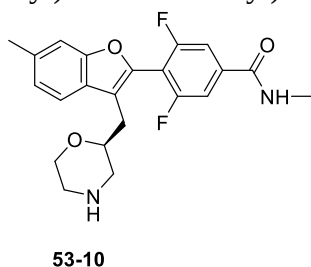
- 5 The crude product was purified by silica gel column chromatography to give **intermediate 53-8** (100 mg, 70% purity, 55% yield) in the form of a brown solid. ESI-MS $[M+Na]^+$: 510.1.

Step (7) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylbenzofuran-3-yl)methyl)morpholine-4-carboxylate



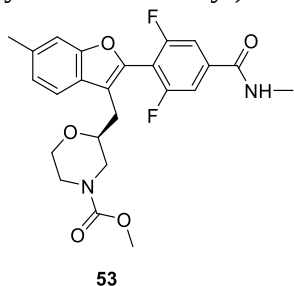
- 10 **Intermediate 53-8** (100 mg, 0.20 mmol) and 2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (114 mg, 0.30 mmol) were dissolved in *N,N*-dimethylformamide (3 mL) under nitrogen atmosphere. The reaction system was stirred at room temperature for 10 min, added with *N,N'*-diisopropylethylamine (1.79 mL, 1.40 mmol) and methylamine hydrochloride (68 mg, 1.00 mmol), and stirred at room temperature for another 4
- 15 h. Then the reaction system was added with water (20 mL) and extracted with ethyl acetate (20 mL × 2). The organic phases were combined and washed with saturated brine (15 mL × 2). The crude product was purified by silica gel column chromatography to give **intermediate 53-9** (65 mg, 63% yield) in the form of a light yellow solid. ESI-MS $[M+Na]^+$: 522.7.

- 20 Step (8) Preparation of (S)-3,5-difluoro-*N*-methyl-4-(6-methyl-3-(morpholin-2-ylmethyl)benzofuran-2-yl)benzamide hydrochloride



- Intermediate 53-9** (65 mg, 0.13 mmol) was dissolved in dichloromethane (2 mL) and a solution of HCl in ethanol (0.5 mL, 0.52 mmol) was added dropwise. The reaction system was stirred at room temperature for 4 h and concentrated under reduced pressure to give **intermediate**
- 25 **53-10** in the form of a light yellow solid (70 mg, crude product), ESI-MS $[M+H]^+$: 400.7.

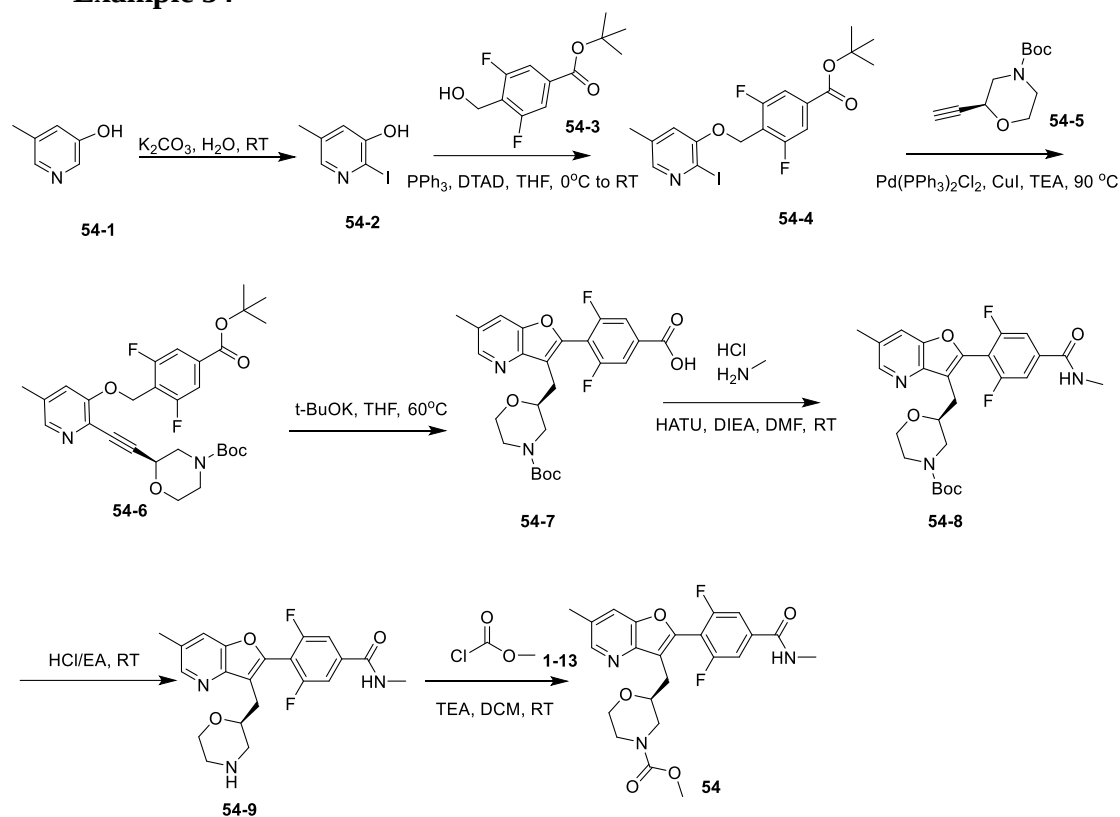
Step (9) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylbenzofuran-3-yl)methyl)morpholine-4-carboxylate



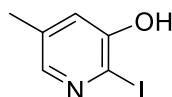
Intermediate 53-10 (70 mg, 0.18 mmol) and triethylamine (0.99 mL, 0.72 mmol) were dissolved in dichloromethane (2 mL), and methyl chloroformate (21 mg, 0.22 mmol) was added dropwise at 0 °C. The reaction system was reacted at this temperature for 4 h. Then the reaction system was poured into ice water (10 mL), and extracted with dichloromethane (15 mL × 2). The organic phases were combined, washed with saturated brine (10 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 53** (22 mg, 22% yield) in the form of a white solid.

ESI-MS $[M + H]^+$: 458.8. ^1H NMR (400 MHz, MeOD) δ 7.63 (d, $J = 8.0$ Hz, 1H), 7.62 (d, $J = 8.4$ Hz, 2H), 7.35 (s, 1H), 7.17 (d, $J = 8.0$ Hz, 1H), 3.81-3.78 (m, 3H), 3.65 (s, 3H), 3.63 – 3.58 (m, 1H), 3.39 – 3.38 (m, 1H), 2.97 (s, 3H), 2.95–2.86 (m, 2H), 2.83 – 2.76 (m, 1H), 2.62-2.58 (m, 1H), 2.51 (s, 3H).

Example 54



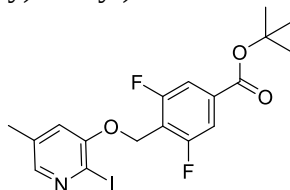
Step (1) Preparation of 2-iodo-5-methylpyridin-3-ol



54-2

- 5 Iodine (2.56 g, 10.08 mmol) was added to a solution of 5-methylpyridin-3-ol (1.10 g, 10.08 mmol) and potassium carbonate (4.18 g, 30.24 mmol) in water (100 mL) while stirring at room temperature. The reaction mixture was stirred at room temperature for 3 h. Then the reaction system was adjusted to pH 6 with diluted hydrochloric acid and extracted with DCM (100 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, and evaporated to dryness. The crude product was purified by silica gel column chromatography to give **intermediate 54-2** (1.9 g, 79.4% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 235.9$.

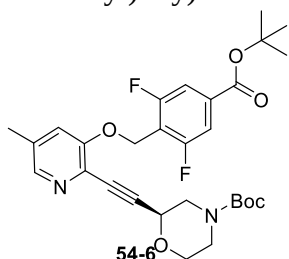
Step (2) Preparation of *tert*-butyl 3,5-difluoro-4-(((2-iodo-5-methylpyridin-3-yl)oxy)methyl)benzoate



54-4

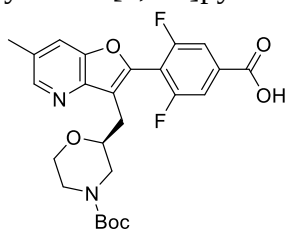
Intermediate 54-2 (773 mg, 3.29 mmol), *tert*-butyl 3,5-difluoro-4-(hydroxymethyl)benzoate (884 mg, 3.62 mmol) and triphenylphosphine (949 mg, 3.62 mmol) were dissolved in THF (15 mL) at 0 °C under nitrogen atmosphere, and a solution of DTAD (833 mg, 3.619 mmol) in THF (5 mL) was added dropwise. After the completion of the dropwise addition, the reaction system was slowly warmed to room temperature and stirred for 4 h. Then the reaction system was added with water (50 mL) to quench the reaction, and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (40 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by column chromatography to give **intermediate 54-4** (1.5 g, 97.9% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 462.0$.

Step (3) Preparation of *tert*-butyl (S)-2-((3-((4-(*tert*-butoxycarbonyl)-2,6-difluorobenzyl)oxy)-5-methylpyridin-2-yl)ethynyl)morpholine-4-carboxylate



To a solution of **intermediate 54-4** (500 mg, 1.08 mmol) and *tert*-butyl (2S)-2-ethynylmorpholin-4-yl formate (344 mg, 1.62 mmol) in TEA (30 mL) were added copper(I) iodide (21 mg, 0.11 mmol) and bis(triphenylphosphine)palladium(II) chloride (38 mg, 0.054 mmol) under nitrogen atmosphere. The reaction system was stirred at 90 °C overnight. Then the reaction system was evaporated to dryness to remove the solvent. The crude product was purified by silica gel column chromatography (petroleum ether/ethyl acetate = 10/1) to give **intermediate 54-6** (452 mg, 73.2% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 545.2$.

Step (4) Preparation of (S)-4-(3-((4-(*tert*-butoxycarbonyl)morpholin-2-yl)methyl)-6-methylfurano[3,2-*b*]pyridin-2-yl)-3,5-difluorobenzoic acid

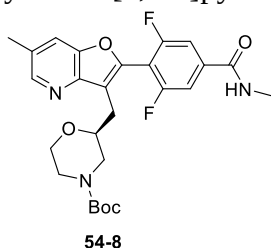


To a solution of **intermediate 54-6** (453 mg, 0.83 mmol) in THF (20 mL) was added *t*-BuOK (232 mg, 2.08 mmol) at room temperature. The reaction system was reacted at 60 °C for 2 h. Then the reaction system was cooled to room temperature, added with water (50 mL),

adjusted to pH 6 with formic acid, and extracted with EA (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 54-7** (230 mg, 56.7% yield) in the form of a yellow solid.

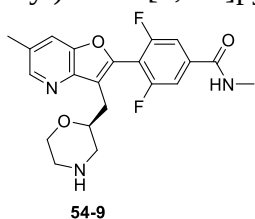
5 LC-MS: $[M+H]^+ = 489.2$.

Step (5) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[3,2-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate



To a solution of **intermediate 54-7** (230 mg, 0.47 mmol) and HATU (268 mg, 0.71 mmol) in DMF (10 mL) was added DIEA (243 mg, 1.88 mmol) at room temperature under nitrogen atmosphere. The reaction system was stirred for 10 min, added with methylamine hydrochloride (63 mg, 0.94 mmol), and stirred at room temperature overnight. Then the reaction system was added with water (50 mL) to quench the reaction and extracted with EA (30 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 54-8** (162 mg, 68.1% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 502.2$.

Step (6) Preparation of (S)-3,5-difluoro-N-methyl-4-(6-methyl-3-(morpholin-2-ylmethyl)furano[3,2-*b*]pyridin-2-yl)benzamide

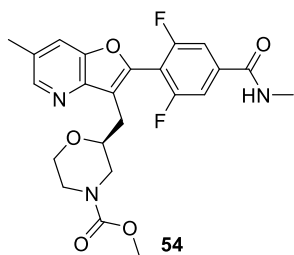


20

A solution of HCl in EA (3 M, 2 mL) was added dropwise to a solution of **intermediate 54-8** (162 mg, 0.32 mmol) in EA (1 mL). The reaction system was stirred at room temperature for 2 h, and concentrated by rotary evaporation to give (S)-3,5-difluoro-N-methyl-4-(6-methyl-3-(morpholin-2-ylmethyl)furano[3,2-*b*]pyridin-2-yl)benzamide hydrochloride in the form of a yellow solid (130 mg, crude product), LC-MS: $[M+H]^+ = 402.1$.

25

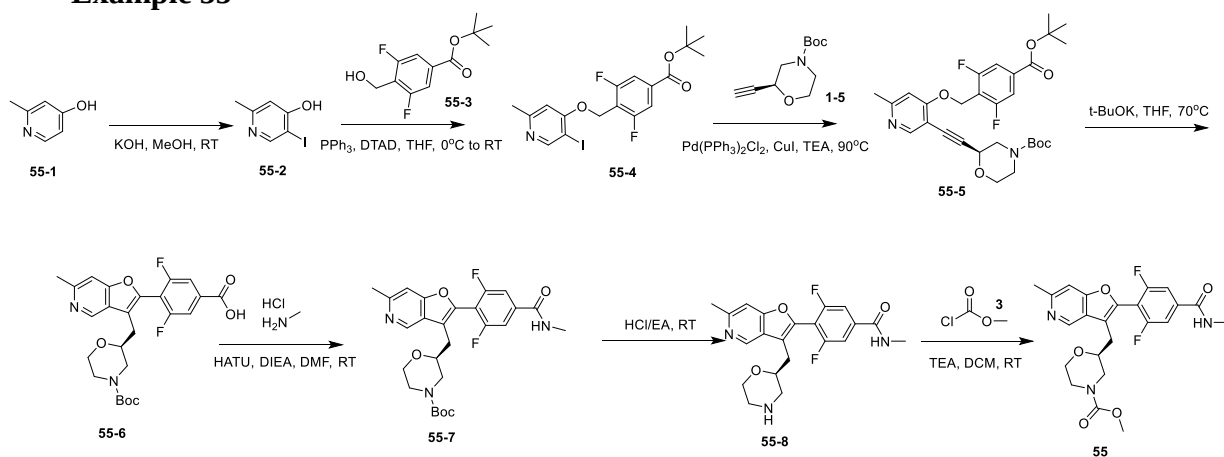
Step (7) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[3,2-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate



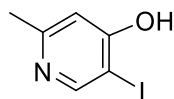
To a solution of **intermediate 54-8** (130 mg, 0.32 mmol) and TEA (100 mg, 0.96 mmol) in DCM (10 mL) was added methyl chloroformate (61 mg, 0.64 mmol) dropwise while stirring at room temperature. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h, and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography, and lyophilized to give **compound 54** (110 mg, 74.3% yield over two steps) in the form of a white solid.

LC-MS: $[M + H]^+ = 460.1$. ^1H NMR (400 MHz, MeOD): δ 8.45 (s, 1H), 7.84 (s, 1H), 7.64 (d, $J = 8.3$ Hz, 2H), 3.93 (d, $J = 13.6$ Hz, 1H), 3.77 (d, $J = 13.2$ Hz, 2H), 3.71-3.63 (m, 4H), 3.38 - 3.29 (m, 1H), 3.03-2.92 (m, 5H), 2.90-2.80 (m, 1H), 2.67-2.56 (m, 1H), 2.55 (s, 3H).

Example 55



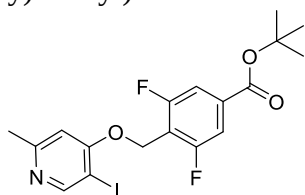
Step (1) Preparation of 5-iodo-2-methylpyridin-4-ol



55-2

Iodine (7.8 g, 31 mmol) was added to a solution of 6-methylpyridin-2-ol (3.0 g, 28 mmol) and potassium hydroxide (6.3 g, 112 mmol) in methanol (200 mL) in portions while stirring at room temperature. The reaction system was stirred at room temperature overnight. Then the reaction system was adjusted to pH 6 with concentrated hydrochloric acid, and concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **intermediate 55-2** (1.2 g, 18.6% yield) in the form of an off-white solid. LC-MS: $[M + H]^+ = 235.8$.

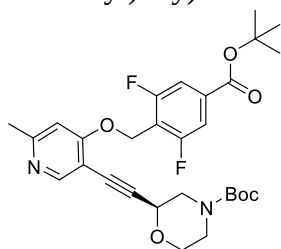
Step (2) Preparation of *tert*-butyl 3,5-difluoro-4-(((5-iodo-2-methylpyridin-4-yl)oxy)methyl)benzoate



55-4

Intermediate 55-2 (470 mg, 2.0 mmol), *tert*-butyl 3,5-difluoro-4-(hydroxymethyl)benzoate (733 mg, 3.0 mmol) and triphenylphosphine (1049 mg, 4.0 mmol) were added to THF (25 mL) at 0 °C under nitrogen atmosphere, and a solution of DTAD (1382 mg, 6.0 mmol) in THF (5 mL) was added dropwise. After the completion of the dropwise addition, the reaction system was slowly warmed to room temperature and stirred for 2 h. Then the reaction system was added with water (50 mL) to quench the reaction, and extracted with dichloromethane (30 mL × 3). The organic phases were combined, and dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 55-4** (620 mg, 66.5% yield) in the form of a white solid. LC-MS $[M+H]^+ = 461.6$.

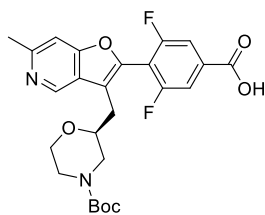
Step (3) Preparation of *tert*-butyl (S)-2-((4-((4-(*tert*-butoxycarbonyl)-2,6-difluorobenzyl)oxy)-6-methylpyridin-3-yl)ethynyl)morpholine-4-carboxylate



55-5

To a solution of **intermediate 55-4** (620 mg, 1.35 mmol) and *tert*-butyl (2S)-2-ethynylmorpholin-4-yl formate (436 mg, 2.06 mmol) in TEA (20 mL) were added copper(I) iodide (26 mg, 0.14 mmol) and bis(triphenylphosphine)palladium(II) chloride (48 mg, 0.07 mmol) under nitrogen atmosphere. The reaction system was heated to 90 °C and reacted overnight. Then the reaction system was concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **intermediate 55-5** (520 mg, 68.6% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 544.8$.

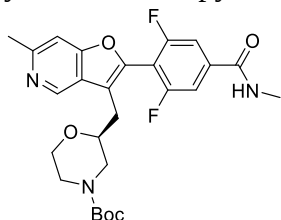
Step (4) Preparation of (S)-4-(3-((4-(*tert*-butoxycarbonyl)morpholin-2-yl)methyl)-6-methylfurano[3,2-*c*]pyridin-2-yl)-3,5-difluorobenzoic acid



55-6

To a solution of **intermediate 55-5** (500 mg, 0.92 mmol) in THF (15 mL) was added t-BuOK (258 mg, 2.30 mmol) under nitrogen atmosphere. The reaction system was stirred at 70 °C for 2 h. Then the reaction system was cooled to room temperature, adjusted to pH 6 with formic acid, and concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **intermediate 55-6** (250 mg, 53.3% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 488.7$.

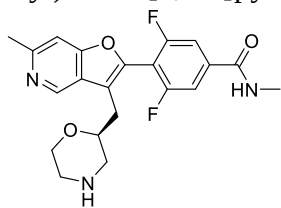
Step (5) Preparation of *tert*-butyl (*S*)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[3,2-*c*]pyridin-3-yl)methyl)morpholine-4-carboxylate



55-7

To a solution of **intermediate 55-6** (250 mg, 0.51 mmol) and HATU (233 mg, 0.61 mmol) in DMF (4 mL) was added DIEA (264 mg, 2.04 mmol) under nitrogen atmosphere. The reaction system was stirred at room temperature for 10 min, added with methylamine hydrochloride (52 mg, 0.77 mmol), and stirred at room temperature overnight. Then the reaction system was added with water (30 mL), and extracted with ethyl acetate (10 mL $\times$ 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 55-7** (250 mg, 94.12% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 502.2$.

Step (6) Preparation of (*S*)-3,5-difluoro-*N*-methyl-4-(6-methyl-3-(morpholin-2-ylmethyl)furano[3,2-*c*]pyridin-2-yl)benzamide

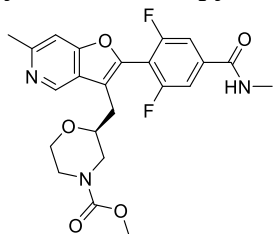


55-8

To a solution of **intermediate 55-7** (250 mg, 0.50 mmol) in EA (5 mL) was added a solution of HCl (g) in EA (3 M, 10 mL) at room temperature. The reaction system was stirred at room

temperature for 2 h, and concentrated under reduced pressure to give **intermediate 55-8** (250 mg, crude product). LC-MS: $[M+H]^+ = 402.1$.

Step (7) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[3,2-c]pyridin-3-yl)methyl)morpholine-4-carboxylate

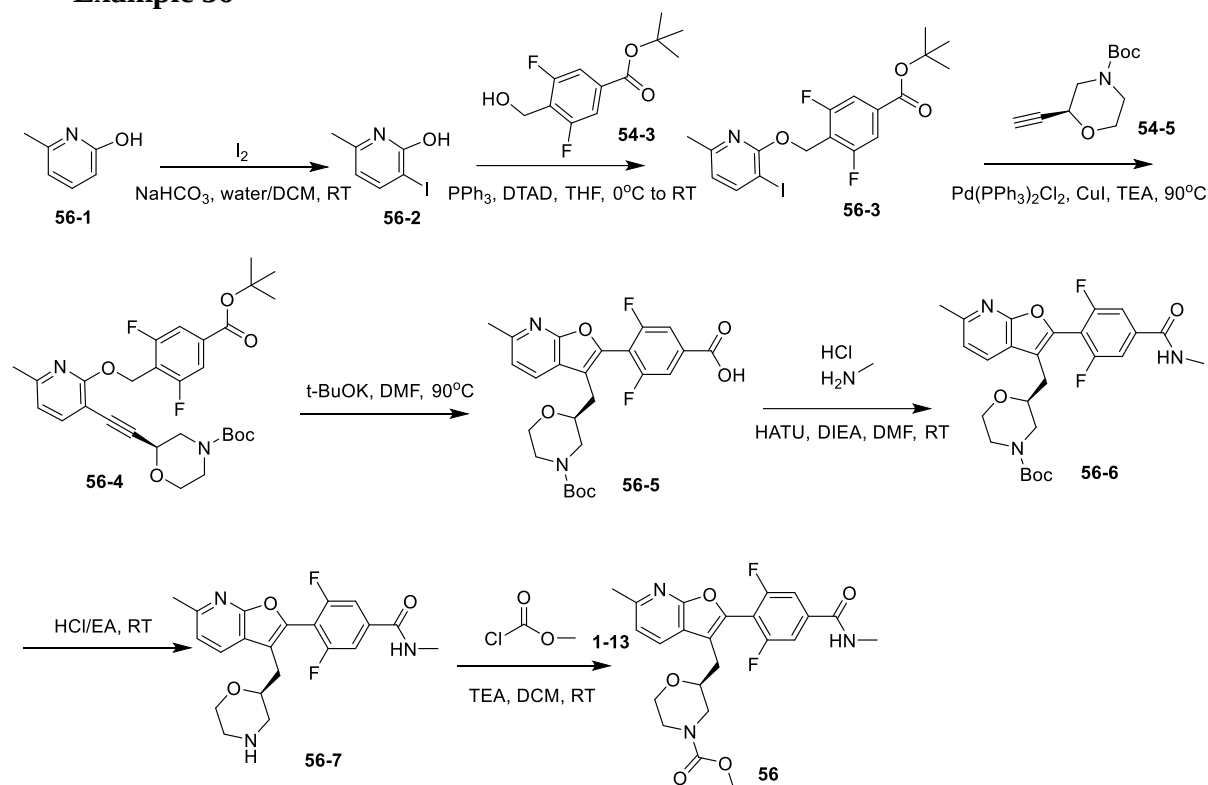


55

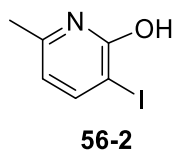
To a solution of **intermediate 55-8** (300 mg, 0.75 mmol) and TEA (228 mg, 2.25 mmol) in DCM (10 mL) was added methyl chloroformate (142 mg, 1.50 mmol) dropwise under nitrogen atmosphere in an ice water bath. After the completion of the dropwise addition, the reaction system was warmed to room temperature and stirred for 2 h. Then the reaction system was concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **compound 55** (70 mg, 20.0% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 460.2$.

^1H NMR (400 MHz, MeOD) δ 8.93 (s, 1H), 7.63 (d, $J = 8.4$ Hz, 2H), 7.48 (s, 1H), 3.90 (d, $J = 13.0$ Hz, 1H), 3.83 – 3.77 (m, 2H), 3.65 (s, 3H), 3.64 – 3.58 (m, 1H), 3.39 (dt, $J = 11.8, 2.8$ Hz, 1H), 2.95 (s, 3H), 2.94 – 2.81 (m, 3H), 2.70 – 2.63 (m, 1H), 2.69 (s, 3H).

Example 56

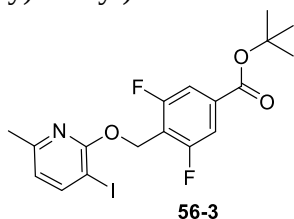


Step (1) Preparation of 3-iodo-6-methylpyridin-2-ol



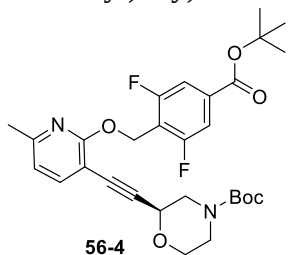
- 5 6-methylpyridin-2-ol (5 g, 46 mmol) and sodium hydrogen carbonate (7.73 g, 92 mmol) were added to a mixture of water and dichloromethane (500 mL, 3/2) while stirring at room temperature, and iodine (11.68 g, 46 mmol) was added. The reaction mixture was stirred at room temperature overnight. Then the reaction system was extracted with dichloromethane (200 mL $\times$ 2). The organic phases were combined, washed with saturated brine (100 mL), dried over
- 10 anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 56-2** (2.5 g, 21.74% yield) in the form of an off-white solid. LC-MS: $[M+H]^+ = 235.0$.

Step (2) Preparation of *tert*-butyl 3,5-difluoro-4-(((3-iodo-6-methylpyridin-2-yl)oxy)methyl)benzoate



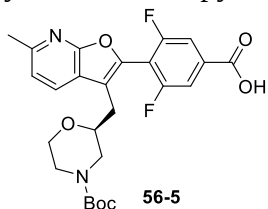
Intermediate 56-2 (510 mg, 2.17 mmol), *tert*-butyl 3,5-difluoro-4-(hydroxymethyl)benzoate (583 mg, 2.39 mmol) and triphenylphosphine (626 mg, 2.39 mmol) were dissolved in THF (15 mL) at 0 °C under nitrogen atmosphere, and a solution of DTAD (550 mg, 2.39 mmol) in THF (5 mL) was added dropwise. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. Then the reaction system was added with water (50 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate and filtered, and the filtrate was concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 56-3** (860 mg, 94.84% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 462.1$.

Step (3) Preparation of *tert*-butyl (S)-2-((2-((4-(*tert*-butoxycarbonyl)-2,6-difluorobenzyl)oxy)-6-methylpyridin-3-yl)ethynyl)morpholine-4-carboxylate



To a solution of **intermediate 56-3** (500 mg, 1.08 mmol) and *tert*-butyl (2S)-2-ethynylmorpholin-4-yl formate (344 mg, 1.62 mmol) in triethylamine (20 mL) were added copper(I) iodide (21 mg, 0.11 mmol) and bis(triphenylphosphine)palladium(II) chloride (38 mg, 0.05 mmol) under nitrogen atmosphere. The reaction mixture was stirred at 90 °C for 12 h. Then the reaction system was concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **intermediate 56-4** (570 mg, 91.67% yield) in the form of a brownish-yellow solid. LC-MS: $[M+H]^+ = 567.1$.

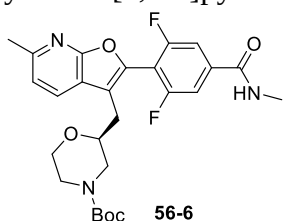
Step (4) Preparation of (S)-4-(3-((4-(*tert*-butoxycarbonyl)morpholin-2-yl)methyl)-6-methylfurano[2,3-*b*]pyridin-2-yl)-3,5-difluorobenzoic acid



To a solution of **intermediate 56-4** (502 mg, 0.92 mmol) in DMF (20 mL) was added potassium *tert*-butoxide (310 mg, 2.76 mmol) under nitrogen atmosphere. The reaction system was stirred at 90 °C for 2 h. After the reaction was completed, the reaction system was cooled to

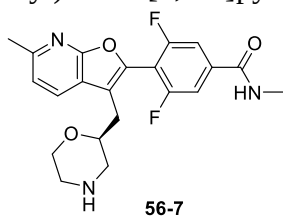
room temperature, adjusted to pH 6 with formic acid, and concentrated by rotary evaporation (oil pump) to remove the solvent. The crude product was purified by silica gel column chromatography to give **intermediate 56-5** (108 mg, 39.13% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 489.8$.

- 5 Step (5) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[2,3-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate



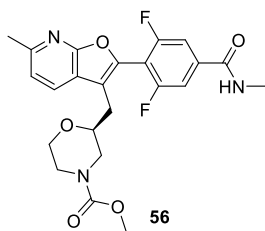
- To a solution of **intermediate 56-5** (180 mg, 0.37 mmol) in DMF (5 mL) were added HATU (211.06 mg, 0.56 mmol) and DIEA (143 mg, 1.11 mmol) under nitrogen atmosphere.
- 10 The reaction system was stirred at room temperature for 10 min, added with methylamine hydrochloride (50 mg, 0.74 mmol), and stirred at room temperature overnight. Then the reaction system was added with water (20 mL), and extracted with ethyl acetate (10 mL $\times$ 3). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column
- 15 chromatography to give **intermediate 56-6** (90 mg, 48.65% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 502.8$.

Step (6) Preparation of (S)-3,5-difluoro-N-methyl-4-(6-methyl-3-(morpholin-2-ylmethyl)furano[2,3-*b*]pyridin-2-yl)benzamide



- 20 To a solution of **intermediate 56-6** (100 mg, 0.20 mmol) in EA (1 mL) was added a solution of HCl (g) in EA (3 M, 2 mL) at room temperature. The reaction system was stirred at room temperature for 2 h. Then the reaction system was concentrated to give **intermediate 56-7** hydrochloride (100 mg, crude product), LC-MS: $[M+H]^+ = 401.9$.

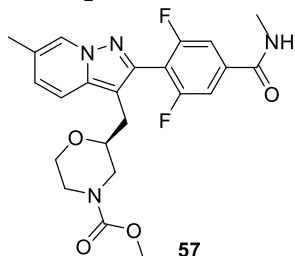
- Step (7) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methylfurano[2,3-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate
- 25



To a solution of **intermediate 56-7** hydrochloride (100 mg, 0.20 mmol) and TEA (101 mg, 1.0 mmol) in DCM (10 mL) was added a solution of methyl chloroformate (47.25 mg, 0.50 mmol) in dichloromethane (1 mL) dropwise in an ice water bath. The reaction system was stirred at room temperature for 2 h. Then the reaction system was added with water (10 mL) to quench the reaction, and extracted with dichloromethane (10 mL × 2). The organic phases were combined, washed with saturated brine (10 mL), filtered and concentrated. The crude product was purified by silica gel column chromatography, and lyophilized to give **compound 56** (45 mg, 49.01% yield) in the form of an off-white solid.

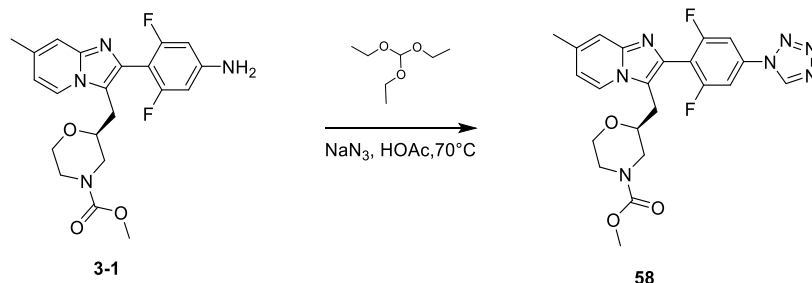
LC-MS: $[M+H]^+ = 460.8$. ^1H NMR (400 MHz, MeOD) : δ 8.19 (d, $J = 7.9$ Hz, 1H), 7.65 (d, $J = 8.4$ Hz, 2H), 7.31 (d, $J = 7.9$ Hz, 1H), 3.89 (d, $J = 12.9$ Hz, 1H), 3.85 – 3.75 (m, 2H), 3.67 (s, 3H), 3.66 – 3.58 (m, 1H), 3.44 – 3.35 (m, 1H), 2.97 (s, 3H), 2.92 – 2.82 (m, 3H), 2.73 – 2.58 (m, 1H), 2.65(s, 3H).

Example 57



The compound was prepared according to the preparation method as described in Example 39. LC-MS: $[M+H]^+ = 459.4$.

Example 58

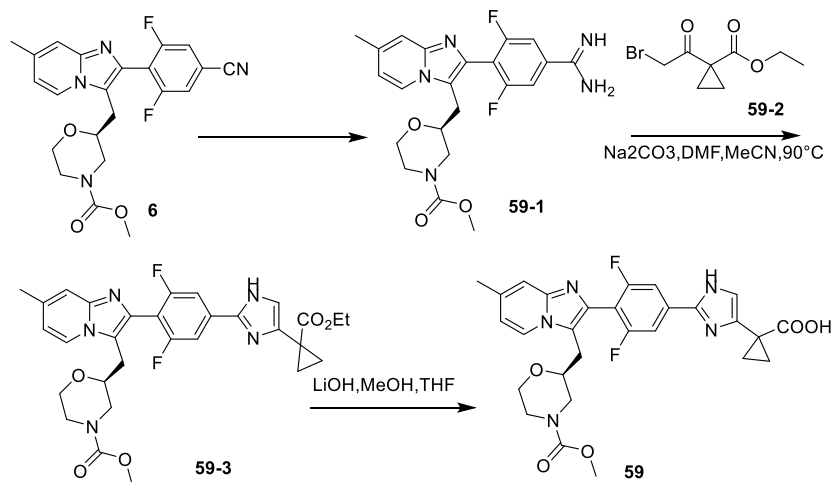


Intermediate 3-1 (400 mg, 0.96 mmol), triethyl orthoformate (456 mg, 3.07 mmol), sodium azide (78 mg, 1.20 mmol) and acetic acid (923 mg, 15.37 mmol) were sequentially added. The reaction system was stirred at room temperature for 10 min, and then stirred at 70 °C for 3 days. Then the reaction system was adjusted to pH 9, and extracted with ethyl acetate three times. The

organic phases were combined, dried and concentrated by rotary evaporation. The crude product was purified by column chromatography to give a product (270 mg) in the form of a white solid, which was then purified by preparative HPLC to give **compound 58** (109 mg, 99% purity, 24% yield) in the form of a white solid.

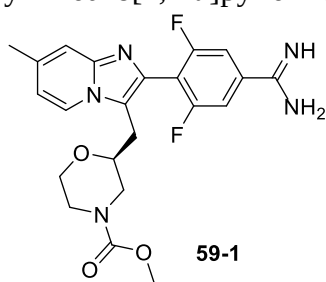
- 5 LCMS $[M+H]^+ = 470$. ^1H NMR (400 MHz, DMSO- d_6) δ 10.18 (s, 1H), 8.39 (dd, $J = 24.5$, 7.3 Hz, 1H), 8.01 – 7.83 (m, 2H), 7.41 – 7.28 (m, 1H), 6.82 (dd, $J = 7.1$, 1.6 Hz, 1H), 3.79 – 3.71 (m, 1H), 3.61 (t, $J = 9.8$ Hz, 2H), 3.52 (s, 3H), 3.46 (dd, $J = 7.5$, 2.8 Hz, 1H), 3.20 (m, 2.8 Hz, 1H), 3.10 – 2.93 (m, 2H), 2.77 (s, 1H), 2.51 (s, 1H), 2.35 (s, 3H).

Example 59



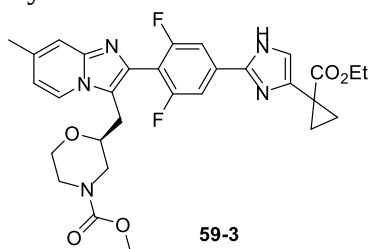
10

Step (1) Preparation of methyl (S)-2-((2-(4-carbamimidoyl-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



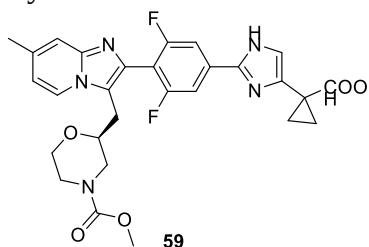
- 15 **Compound 6** (800 mg, 1.88 mmol) and sodium methoxide (51 mg, 0.94 mmol) were dissolved in methanol (8 mL) under nitrogen atmosphere. The reaction system was stirred at room temperature overnight, added with NH_4Cl solid (152 mg, 2.80 mmol), and stirred at room temperature overnight. Then the reaction system was adjusted to pH 8 with saturated sodium bicarbonate solution (10 mL), concentrated under reduced pressure to 1/2 volume, and extracted with dichloromethane (20 mL $\times$ 3). The organic phases were combined, washed with saturated
- 20 brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated to give **intermediate 59-1** (760 mg, crude product) in the form of a brown solid. LC-MS $[M+H]^+ : 444$.

Step (2) Preparation of methyl (S)-2-((2-(4-(4-(1-(ethoxycarbonyl)cyclopropyl)-1H-imidazol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 59-1 (800 mg, 1.80 mmol) and ethyl 1-(2-bromoacetyl)cyclopropane-1-carboxylate (329 mg, 1.4 mmol) were dissolved in *N,N*-dimethylformamide (8 mL) under nitrogen atmosphere, and potassium bicarbonate (360 mg, 3.6 mmol) was added. The reaction system was stirred at 70 °C overnight. Then the reaction system was added with water (30 mL) to quench the reaction, and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by silica gel column chromatography to give **intermediate 59-3** (600 mg, 57% yield) in the form of a brown solid. LC-MS $[M+H]^+$: 580.

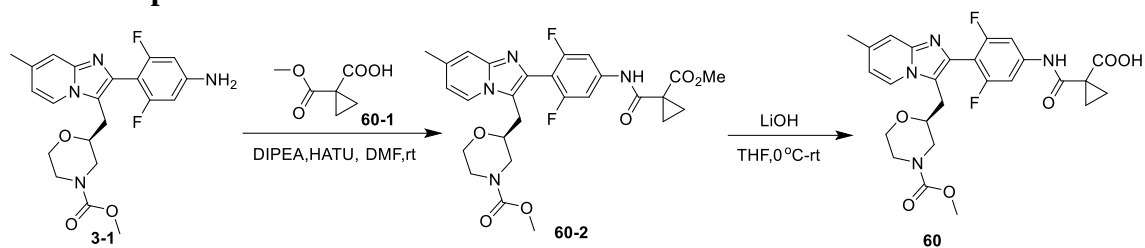
Step (3) Preparation of (S)-1-(2-(3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)-1H-imidazol-4-yl)cyclopropane-1-carboxylic acid



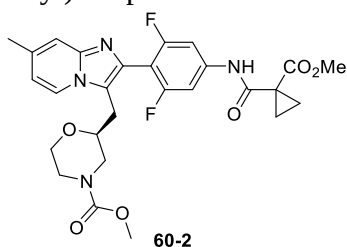
Intermediate 59-3 (600 mg, 1.1 mmol) was dissolved in tetrahydrofuran (4 mL) and absolute methanol (2 mL), and lithium hydroxide solution (1.5 mL, 3 M) was added dropwise at 0 °C. The reaction system was warmed to room temperature and stirred for 3 h. Then the reaction system was adjusted to pH 4 with diluted hydrochloric acid (1 N), and concentrated to give a crude product (450 mg) in the form of a yellow solid. The crude product (50 mg) was purified by preparative chromatography to give **compound 59** (4.6 mg) in the form of a white solid.

LC-MS $[M+H]^+$: 552. ^1H NMR (400 MHz, $CDCl_3$) δ 8.28 (d, J = 8.0 Hz, 1H), 7.47 (s, 1H), 7.30 (d, J = 8.3 Hz, 2H), 6.81 (d, J = 7.7 Hz, 1H), 6.75 (s, 1H), 4.00 – 3.81 (m, 3H), 3.67 (s, 3H), 3.63 – 3.57 (m, 1H), 3.45 – 3.35 (m, 1H), 3.05 – 2.85 (m, 3H), 2.66 – 2.57 (m, 1H), 2.52 (s, 3H), 1.92 – 1.86 (m, 2H), 1.25 – 1.19 (m, 2H).

Example 60

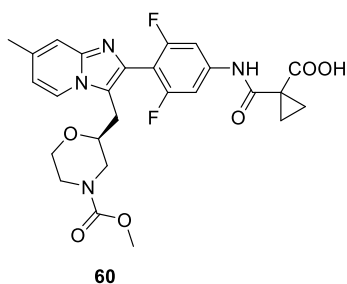


Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1-(methoxycarbonyl)cyclopropane-1-carboxamido)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



1-(methoxycarbonyl)cyclopropane-1-carboxylic acid (104 mg, 0.72 mmol) and 2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (275 mg, 0.72 mmol) were dissolved in anhydrous *N,N*-dimethylformamide (2 mL) under nitrogen atmosphere. The reaction system was reacted at room temperature for 10 min, added with *N,N*-diisopropylethylamine (252 mg, 1.95 mmol) and **intermediate 3-1** (200 mg, 0.48 mmol), and reacted at room temperature overnight. Then the reaction system was added with water (20 mL), and extracted with ethyl acetate (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **intermediate 60-2** (100 mg, 51.3% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 543.7$.

Step (2) Preparation of (S)-2-((2-(2,6-difluoro-4-(1-(methoxycarbonyl)cyclopropane-1-carboxamido)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylic acid

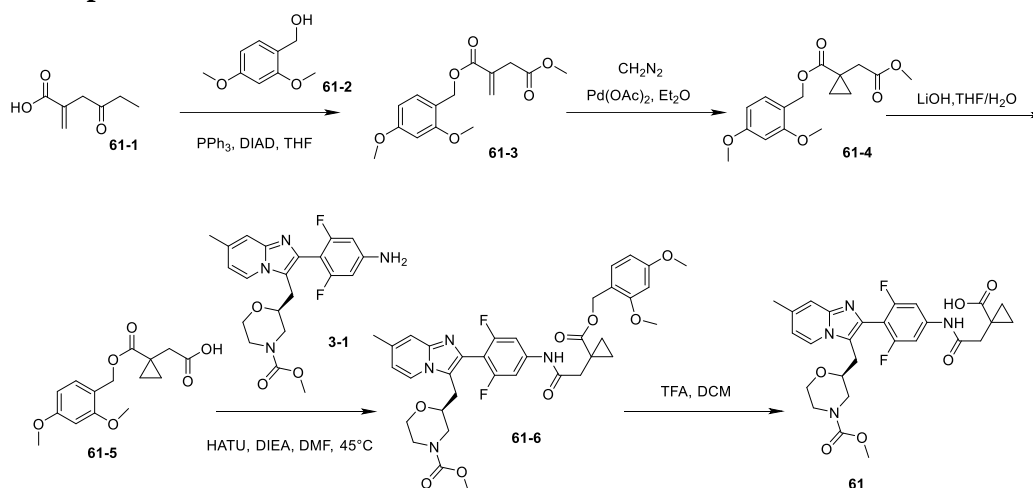


Intermediate 60-2 (100 mg, 0.18 mmol) was dissolved in tetrahydrofuran (1.5 mL), and lithium hydroxide solution (0.18 mL, 3 M) was added dropwise. The reaction system was reacted

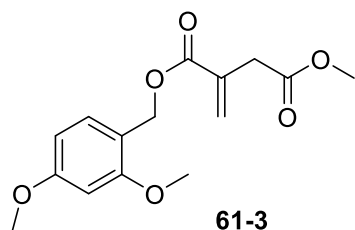
at room temperature for 1 h. After the reaction was completed, the reaction system was added dropwise with diluted hydrochloric acid (1 M) to adjust the pH to 6, added with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 60** (65.9 mg, 67.6% yield) in the form of a white solid.

LC-MS: $[M + H]^+ = 529.6$. ^1H NMR (400 MHz, MeOD) : δ 8.84 (d, $J = 7.1$ Hz, 1H), 7.70 (s, 1H), 7.64 (d, $J = 7.4$ Hz, 2H), 7.41 (dd, $J = 7.1, 1.5$ Hz, 1H), 4.01-3.97(m, 1H), 3.86-3.74 (m, 2H), 3.70 (s, 3H), 3.69 – 3.65 (m, 1H), 3.39-3.36 (m, 1H), 3.25-3.24 (m, $J = 5.2$ Hz, 2H), 2.99-2.85 (m, 1H), 2.81-2.67 (m, 1H), 2.65 (s, 3H), 1.75-1.72 (m, 2H), 1.71-1.68 (m, 2H).

Example 61



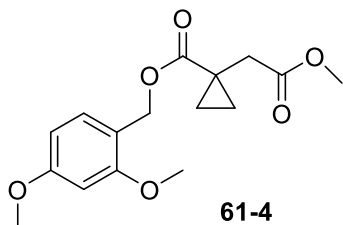
Step (1) Preparation of 1-(2,4-dimethoxybenzyl)-4-methyl 2-methylenesuccinate



Monomethyl itaconate (3.00 g, 21.10 mmol), (2,4-dimethoxy)benzyl alcohol (3.90 g, 23.21 mmol) and triphenylphosphine (6.65 g, 25.33 mmol) were dissolved in THF (50 mL) at 0 °C under nitrogen atmosphere, and a solution of DIAD (5.12 g, 25.33 mmol) in THF (10 mL) was added dropwise. After the completion of the dropwise addition, the reaction system was warmed to room temperature and stirred overnight. Then the reaction system was added with water (60 mL) to quench the reaction, and extracted with ethyl acetate (40 mL × 3). The organic phases were combined, washed with saturated brine (40 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by silica gel

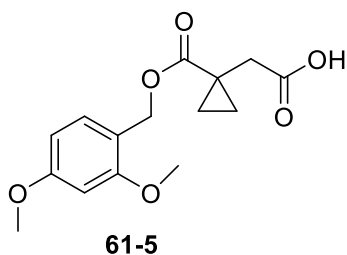
column chromatography to give **intermediate 61-3** (4.5 g, 72.2% yield) in the form of a white solid. MS $[M + Na]^+ = 316.8$.

Step (2) Preparation of 2,4-dimethoxybenzyl 1-(2-methoxy-2-carbonylethyl)cyclopropanecarboxylate



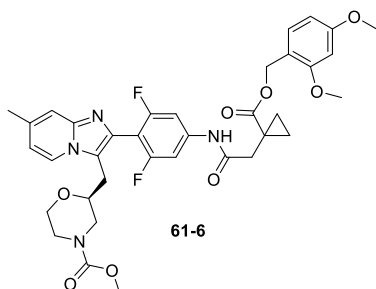
To a mixture of potassium hydroxide (30 mL, 40% aq) and diethyl ether (20 mL) was added 1-methyl-3-nitro-1-nitrosoguanidine (4.0 g, 13.6 mmol, 50% water content) at 0 °C under nitrogen atmosphere. The reaction system was stirred at 0 °C for 15 min and left to stand for liquid separation. The upper organic phase was collected, and dried over anhydrous sodium sulfate. **Intermediate 61-3** (600 mg, 2.04 mmol) and palladium acetate (92 mg, 0.41 mmol) were dissolved in diethyl ether (15 mL) at 0 °C under nitrogen atmosphere, and a fresh solution of diazomethane in diethyl ether was added dropwise. After the completion of the dropwise addition, the reaction system was slowly warmed to room temperature and stirred for 16 h. The reaction system was added with absolute methanol (10 mL) to quench the reaction and filtered, and the filtrate was concentrated by rotary evaporation to give a crude product. The crude product was purified by column chromatography to give **intermediate 61-4** (520 mg, 82.7% yield) in the form of a colorless oil. MS $[M + Na]^+ = 330.8$.

Step (3) Preparation of 2-(1-(((2,4-dimethoxybenzyl)oxo)carbonyl)cyclopropyl)acetic acid



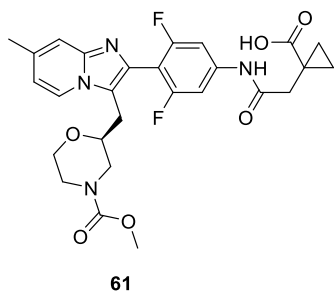
To a mixture of **intermediate 61-4** (500 mg, 1.62 mmol) in tetrahydrofuran (40 mL) and water (10 mL) was added lithium hydroxide monohydrate (11.9 mL, 11.9 mmol) at 0 °C. The reaction was stirred at 0 °C for 8 h. Then the reaction system was warmed to room temperature, adjusted to pH 4 with diluted hydrochloric acid, and extracted with DCM (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 61-5** (320 mg, 73.2% yield, crude product) in the form of a yellow oil. MS $[M - H]^- = 292.4$.

Step (4) Preparation of methyl (S)-2-((2-(4-(2-(1-(((2,4-dimethoxybenzyl)oxo)carbonyl)cyclopropyl)acetamido)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



5 To a solution of **intermediate 61-5** (300 mg, 1.02 mmol) in DMF (5 mL) were added DIEA (527 mg, 4.08 mmol) and HATU (466 mg, 1.22 mmol) at room temperature under nitrogen atmosphere. The reaction system was stirred at 45 °C for 0.5 h, added with **intermediate 3-1** (213 mg, 0.51 mmol), and stirred at 45 °C for another 18 h. Then the reaction system was cooled to room temperature, added with water (10 mL), and extracted with EA (10 mL × 3). The organic
10 phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 61-6** (160 mg, 22.6% yield) in the form of a yellow solid. MS [M+H]⁺ = 692.7.

Step (5) Preparation of (S)-1-(2-((3,5-difluoro-4-(3-((4-(carbomethoxy<methoxycarbonyl>)morpholin-2-yl)methyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)phenyl)amino)-2-carbonylethyl)cyclopropanecarboxylic acid

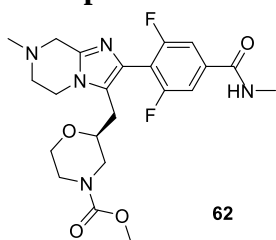


To a solution of **intermediate 61-6** (100 mg, 0.14 mmol) in dichloromethane (2 mL) was added trifluoroacetic acid (1 mL) at room temperature. The reaction system was stirred at room
20 temperature for 3 h and concentrated. The residue was separated by preparative high performance liquid chromatography, and lyophilized to give **compound 61** (29 mg, 36.9% yield) in the form of a white solid. MS [M+H]⁺ = 542.7.

¹H NMR (400 MHz, CDCl₃) δ 10.44 (brs, 1H), 8.51 (d, J = 7.0 Hz, 1H), 7.96 (s, 1H), 7.43 (d, J = 11.1 Hz, 2H), 7.09 (d, J = 6.7 Hz, 1H). 4.10 – 3.75 (m, 3H), 3.69 (s, 3H), 3.65-3.61(m,

1H), 3.45 - 3.35 (m, 1H), 3.05 – 2.85 (m, 3H), 2.76-2.68 (m, 1H), 2.58 (s, 3H), 1.39 (s, 2H), 0.94 (s, 2H).

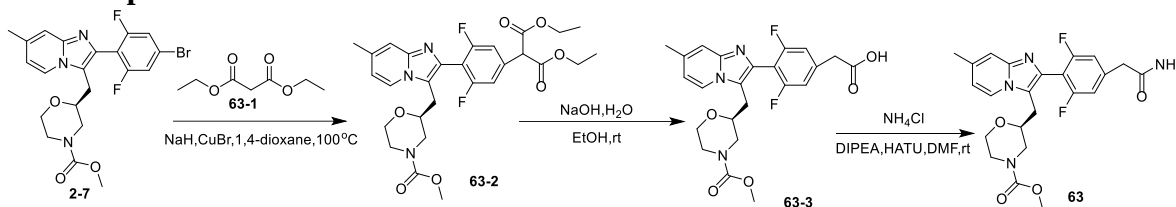
Example 62



62

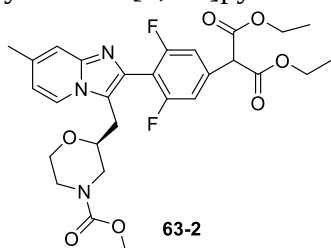
5 The compound was prepared according to the preparation method as described in Example 44. LC-MS: $[M+H]^+ = 464.2$.

Example 63



Step (1) Preparation of methyl (S)-2-((2-(4-(diethylmalonate)-2,6-difluorophenyl)-7-

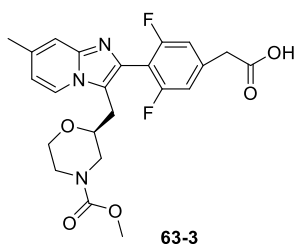
10 methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



63-2

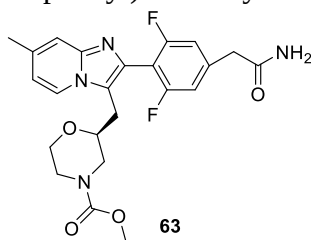
Intermediate 2-7 (300 mg, 0.62 mmol), 1,3-diethyl malonate (200 mg, 1.24 mmol) and bromoidenone (180 mg, 1.24 mmol) were dissolved in 1,4-dioxane (3 mL), and sodium hydride (60%, 55 mg, 1.36 mmol) was added in portions under nitrogen atmosphere. The reaction system was reacted at 100 °C for 10 h. After the reaction was completed, the reaction system was cooled to room temperature, added with water (20 mL) to quench the reaction, and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by column chromatography to give **intermediate 63-2** (230 mg, 66% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 560.2$.

Step (2) (S)-2-(3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)acetic acid



Intermediate 63-2 (210 mg, 0.38 mmol) was dissolved in absolute ethanol (1.5 mL), and sodium hydroxide solution (3 M, 45 mg, 1.14 mmol) was added. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was adjusted to pH 6 with hydrochloric acid solution (1 M), added with water (5 mL), and extracted with dichloromethane (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 63-3** (140 mg, 77% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ = 460.2$.

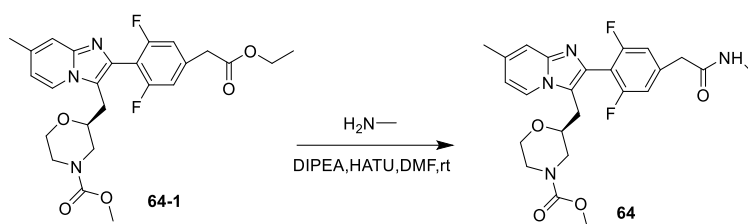
Step (3) Preparation of methyl (S)-2-((2-(4-(2-amino-2-carbonylethyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



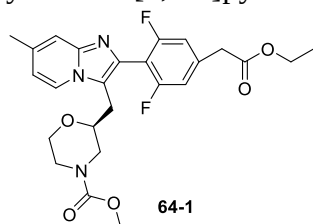
Intermediate 63-3 (70 mg, 0.15 mmol) and 2-(7-azobenzotriazol)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (87 mg, 0.23 mmol) were dissolved in *N,N*-dimethylformamide (2 mL) under nitrogen atmosphere. The reaction system was reacted at room temperature for 10 min, added with *N,N*-diisopropylethylamine (60 mg, 0.46 mmol) and ammonium chloride (10 mg, 0.19 mmol), and reacted at room temperature for another 2 h. After the reaction was completed, the reaction system was added with water (20 mL), and extracted with ethyl acetate (30 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by preparative chromatography to give **compound 63** (10.5 mg, 15% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 459.2$. ^1H NMR (400 MHz, CDCl_3): δ 8.60 (brs, 1H), 7.97 (brs, 1H), 7.67-7.62 (brs, 1H), 7.27-7.08 (m, 3H), 6.50 (brs, 1H), 4.08-3.78 (m, 4H), 3.73 (s, 3H), 3.71-3.58 (m, 2H), 3.45-3.37 (m, 1H), 3.07-3.03 (m, 2H), 3.00-2.92 (m, 1H), 2.77-2.69 (m, 1H), 2.62 (s, 3H).

Example 64



Step (1) Preparation of methyl (S)-2-((2-(4-(2-ethoxy-2-oxoethyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

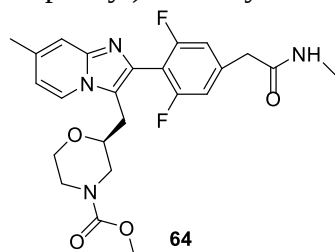


5

Compound 63-3 (459 mg, 1 mmol) was dissolved in absolute ethanol (10 mL), and H₂SO₄ (0.1 mmol) was added. The reaction system was heated to reflux. After the reaction was completed as monitored by TLC, the reaction system was diluted with water with the pH adjusted to be neutral, and concentrated under reduced pressure. The aqueous phase was extracted with EA. The organic phase was dried over anhydrous MgSO₄, filtered, and concentrated under reduced pressure to give **intermediate 64-1** (0.36 g, 74% yield), LC-MS: [M+H]⁺ = 488.2.

10

Step (2) Preparation of methyl (S)-2-((2-(4-(2-(methylamino)-2-oxoethyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



15

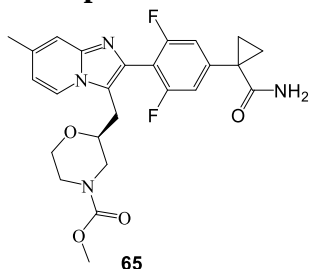
Intermediate 64-1 (70 mg, 0.15 mmol) and 2-(7-azobenzotriazol)-N,N,N',N'-tetramethyluronium hexafluorophosphate (87 mg, 0.23 mmol) were dissolved in N,N-dimethylformamide (2 mL) under nitrogen atmosphere. The reaction system was reacted at room temperature for 10 min, added with N,N-diisopropylethylamine (60 mg, 0.46 mmol) and a solution of methylamine in tetrahydrofuran (1 M, 0.1 mL, 0.15 mmol), and reacted at room temperature for another 2 h. After the reaction was completed, the reaction system was added with water (20 mL), and extracted with ethyl acetate (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 3), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was

20

purified by high performance liquid chromatography to give **compound 64** (17.2 mg, 24% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 473.2$. ^1H NMR (400 MHz, $CDCl_3$): δ 8.59 (brs, 1H), 7.92 (s, 1H), 7.55 (brs, 1H), 7.19 (d, $J = 7.4$ Hz 1H), 7.08 (d, $J = 8.1$ Hz, 2H), 4.10- 3.90 (m, 3H), 3.86-3.78 (m, 1H), 3.73 (s, 3H), 3.73-3.65 (m, 2H), 3.46 -3.35 (m, 1H), 3.08-3.02 (m, 2H), 3.00-2.90 (m, 1H), 2.85 (s, 3H), 2.76-2.70 (m, 1H), 2.62 (s, 3H).

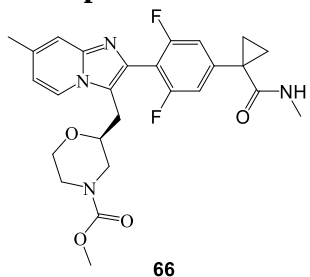
Example 65



The compound was prepared according to the preparation method as described in Example

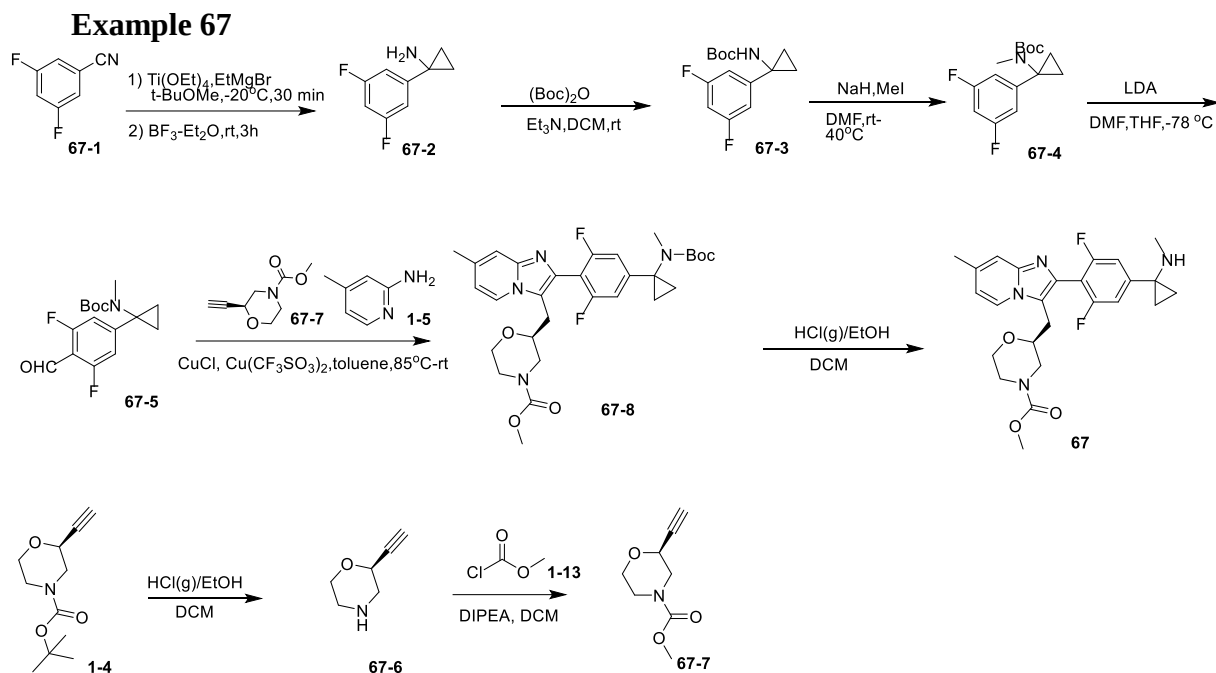
63. LC-MS: $[M+H]^+ = 485.2$.

Example 66

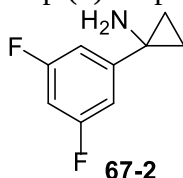


The compound was prepared according to the preparation method as described in Example

63. LC-MS: $[M+H]^+ = 499.3$.

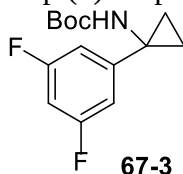


Step (1) Preparation of 1-(3,5-difluorophenyl)cyclopropane-1-amine



- 5 3,5-difluorobenzonitrile (10.0 g, 71.89 mmol) and tetraisopropyl titanate (27.0 g, 95.00 mmol) were dissolved in methyl *tert*-butyl ether (120 mL), and ethyl magnesium bromide (3 M in diethyl ether, 55 mL, 165.35 mmol) was added at -20°C under nitrogen atmosphere. The reaction system was reacted at this temperature for 30 min, added with boron trifluoride diethyl ether (46.5%, 44 g, 143.78 mmol), and stirred at room temperature for another 3 h. Then the
- 10 reaction system was added with sodium hydroxide solution (3 M, 72 mL, 215.67 mmol) and filtered, and the filtrate was added with water (100 mL) and extracted with dichloromethane (150 mL $\times$ 3). The organic phases were combined, washed with saturated brine (100 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give
- 15 **intermediate 67-2** (2 g, 16.6% yield) in the form of a yellow oil, $\text{MS } [\text{M}+\text{H}]^+ = 170.0$.

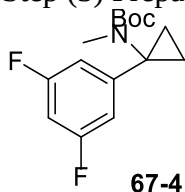
Step (2) Preparation of *tert*-butyl (1-(3,5-difluorophenyl)cyclopropyl)carbamate



Intermediate 67-2 (2.0 g, 11.83 mmol) was dissolved in anhydrous dichloromethane (20 mL), and di-*tert*-butyl dicarbonate (3.9 g, 17.75 mmol) and triethylamine (2.4 g, 23.67 mmol)

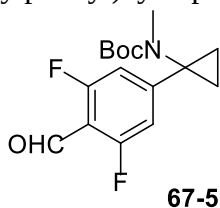
were added under nitrogen atmosphere. The reaction system was reacted at room temperature overnight. Then the reaction system was added with water (50 mL), and extracted with dichloromethane (50 mL). The organic phase was washed with saturated brine (50 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 67-3** (640 mg, 18.08% yield) in the form of a white solid. MS $[M+H]^+ = 269.9$.

Step (3) Preparation of *tert*-butyl (1-(3,5-difluorophenyl)cyclopropyl)(methyl)carbamate



Intermediate 67-3 (620 mg, 2.30 mmol) was dissolved in anhydrous *N,N*-dimethylformamide (5 mL) under nitrogen atmosphere, and sodium hydride (60%, 138 mg, 3.45 mmol) was added. The reaction system was reacted at room temperature for 30 min, added with methyl iodide (720 mg, 5.07 mmol) and reacted at 40 °C for 1 h. Then the reaction system was added with ice water (20 mL) to quench the reaction, and extracted with ethyl acetate (20 mL $\times$ 2). The organic phases were combined, washed with saturated brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 67-4** (260 mg, 40% yield) in the form of a white solid. MS $[M+H]^+ = 283.9$.

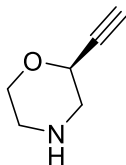
Step (4) Preparation of *tert*-butyl (1-(3,5-difluoro-4-formylphenyl)cyclopropyl)(methyl)carbamate



Intermediate 67-4 (260 mg, 0.92 mmol) was dissolved in anhydrous tetrahydrofuran (5 mL) under nitrogen atmosphere, and lithium diisopropylamide (2 M in tetrahydrofuran, 0.6 mL, 1.10 mmol) was added dropwise at -78 °C. After the completion of the dropwise addition, the reaction system was reacted at -78 °C for 1 h, added with anhydrous *N,N*-dimethylformamide (67 mg, 0.92 mmol), and reacted for another 30 min. After the reaction was completed, the reaction system was added with saturated ammonium chloride solution to quench the reaction, diluted with water (20 mL), and extracted with ethyl acetate (20 mL $\times$ 2). The organic phases were combined, washed with saturated brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel

column chromatography to give **intermediate 67-5** (250 mg, 85% yield) in the form of a white solid. MS $[M+H]^+ = 311.9$.

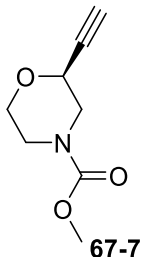
Step (5) Preparation of (S)-2-ethynylmorpholine



67-6

Intermediate 1-4 (200 mg, 0.95 mmol) was dissolved in anhydrous dichloromethane (3 mL), and a solution of HCl in ethanol (33%, 0.5 mL) was added at 0 °C. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give **intermediate 67-6** (230 mg, crude product) in the form of a yellow oil, MS $[M+H]^+ = 120.0$.

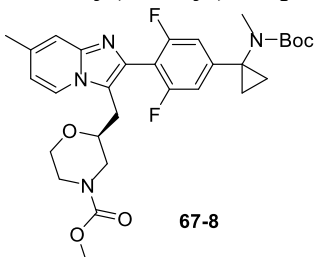
Step (6) Preparation of methyl (S)-2-ethynylmorpholine-4-carboxylate



67-7

Intermediate 67-6 (230 mg, crude product, 0.95 mmol) and *N,N*-diisopropylethylamine (1.1 g, 8.51 mmol) were dissolved in anhydrous dichloromethane (3 mL), and methyl chloroformate (180 mg, 1.90 mmol) was added dropwise slowly in an ice water bath. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 1 h. After the reaction was completed, the reaction system was added with water (15 mL), and extracted with dichloromethane (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 67-7** (160 mg, crude product) in the form of a yellow oil. MS $[M+H]^+ = 169.7$.

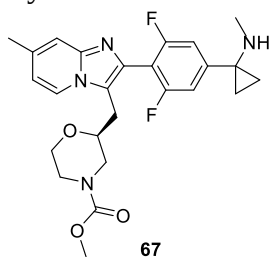
Step (7) Preparation of methyl (S)-2-((2-(4-(1-((*tert*-butoxycarbonyl)(methyl)amino)cyclopropyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



67-8

Intermediate 67-5 (250 mg, 0.80 mmol), 4-methylpyridin-2-amine (87 mg, 0.80 mmol), methyl (S)-2-ethynylmorpholine-4-carboxylate (136 mg, 0.80 mmol), copper(I) chloride (24 mg, 0.24 mmol) and copper(II) trifluoromethanesulfonate (87 mg, 0.24 mmol) were dissolved in toluene solution (3 mL) under nitrogen atmosphere. The reaction system was reacted at 85 °C for 10 min, added with *N,N*-dimethylacetamide (0.1 mL), and stirred at 85 °C for another 5 h. Then the reaction system was reacted at room temperature overnight. The resulting reaction system was filtered, and the filtrate was added with water (20 mL) and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 67-8** (150 mg, 32.6% yield) in the form of a yellow oil. MS $[M+H]^+ = 570.8$.

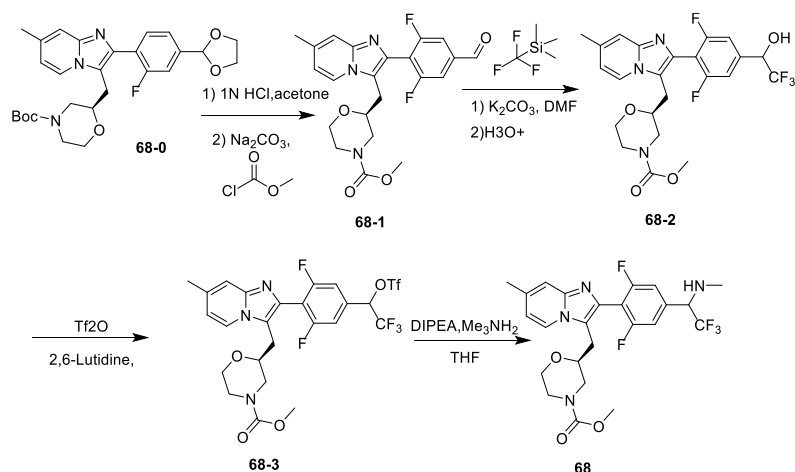
Step (8) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1-(methylamino)cyclopropyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



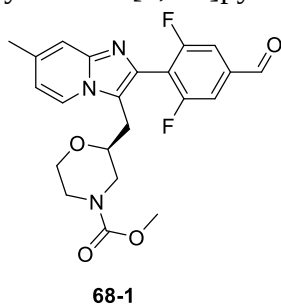
Intermediate 67-8 (150 mg, 0.26 mmol) was dissolved in anhydrous dichloromethane (3 mL), and a solution of HCl in ethanol (33%, 0.5 mL) was added at 0 °C. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation, diluted with dichloromethane (30 mL), dissolved with saturated sodium bicarbonate solution (10 mL), and stirred for 5 min. Then the reaction system was extracted. The organic phase was dried over anhydrous sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 67** (55.4 mg, 43.5% yield) in the form of a white solid. MS $[M+H]^+ = 470.8$.

¹H NMR (400 MHz, MeOD) δ 8.85 (d, *J* = 7.1 Hz, 1H), 7.73 (s, 1H), 7.59 (d, *J* = 8.4 Hz, 2H), 7.41 (d, *J* = 7.0 Hz, 1H), 4.02 – 3.93(m, 1H), 3.87 – 3.73 (m, 2H), 3.70 (s, 3H), 3.69 – 3.62 (m, 1H), 3.39 – 3.35 (m, 1H), 3.28 – 3.23(m, 2H), 2.99 – 2.85 (m, 1H), 2.75 (s, 3H), 2.74 – 2.66 (m, 1H), 2.65 (s, 3H), 1.61 – 1.54(m, 2H), 1.52 – 1.46 (m, 2H).

Example 68



Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-formylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



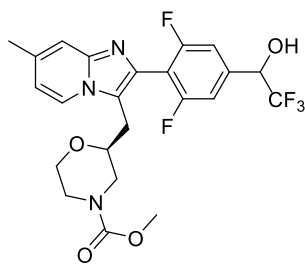
5

Tert-butyl (*R*)-2-((2-(4-(1,3-dioxan-2-yl)-2-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate (2.0 g, 4.0 mmol) was dissolved in acetone (15 mL). The reaction system was cooled to 0 °C, added dropwise with diluted hydrochloric acid (1 N, 5 mL), and warmed to room temperature and stirred for 4 h. Then the reaction system was adjusted to pH 8 with saturated sodium carbonate solution (8 mL), added dropwise with methyl chloroformate (0.45 g, 4.8 mmol) at 0 °C, and stirred at 0 °C for 1 h. The resulting reaction system was added with water (50 mL), and extracted with dichloromethane (60 mL × 3). The organic phases were combined, washed with saturated brine (100 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by silica gel column chromatography to give **intermediate 68-1** (770 mg, 46% yield) in the form of a yellow solid. LC-MS [M+H]⁺ = 430.

10

15

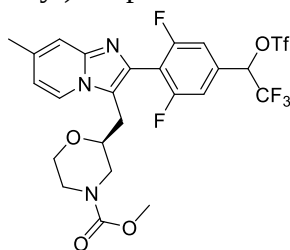
Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(2,2,2-trifluoro-1-hydroxyethyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



68-2

Intermediate 68-1 (100 mg, 0.23 mmol) was dissolved in *N,N*-dimethylformamide (2 mL) under nitrogen atmosphere, and potassium carbonate (16 mg, 0.11 mmol) was added, followed by the dropwise addition of trifluoromethyl trimethylsilane (50 mg, 0.30 mmol). The reaction system was stirred at room temperature for 4 h, added with HCl (2 mL, 2 N), and stirred at room temperature overnight. Then the reaction system was added with water (20 mL), and extracted with dichloromethane (15 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by silica gel column chromatography to give **intermediate 68-2** (80 mg, 69% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 500$.

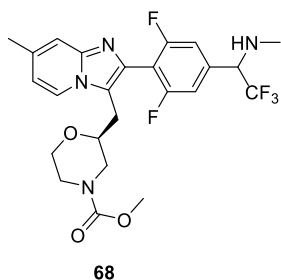
Step (3) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(2,2,2-trifluoro-1-((trifluoromethyl)sulfonyl)oxy)ethyl))-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



68-3

Intermediate 68-2 (80 mg, 0.16 mmol) and 2,6-dimethylpyridine (34 mg, 0.32 mmol) were dissolved in dichloromethane (3 mL) under nitrogen atmosphere. The reaction system was cooled to 0 °C, added dropwise with trifluoromethanesulfonic anhydride (81 mg, 0.29 mmol), and stirred for 1 h. Then the reaction system was concentrated to give **intermediate 68-3** (100 mg) in the form of a brown solid, which was used directly in the next step. LC-MS: $[M+H]^+ = 632$.

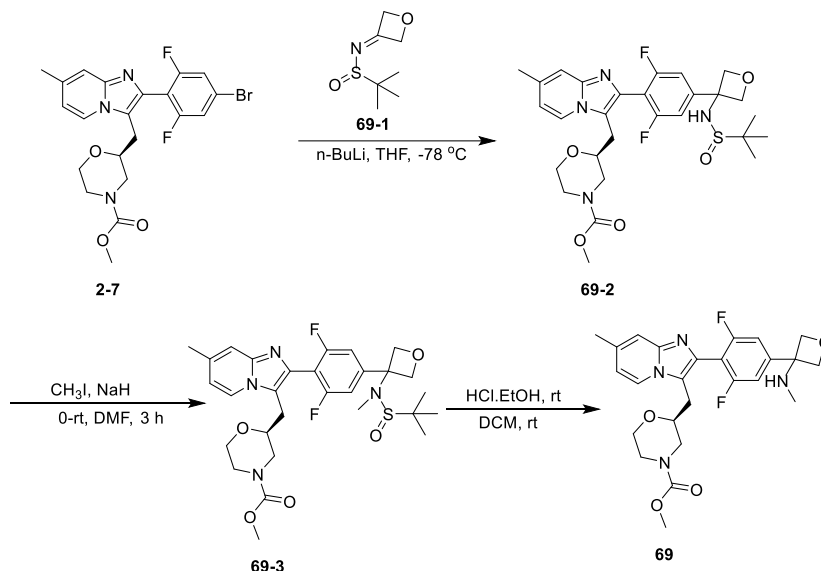
Step (4) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(2,2,2-trifluoro-1-(methylamino)ethyl)phenyl))-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



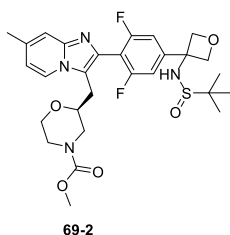
Intermediate 68-3 (100 mg, 0.16 mmol) and *N,N*-diisopropylethylamine (62 mg, 0.48 mmol) were dissolved in tetrahydrofuran (2 mL), and methylamine in tetrahydrofuran (1 mL, 2 N) was added. The reaction system was stirred at room temperature for 2 h under nitrogen atmosphere. Then the reaction system was added with water (20 mL) to quench the reaction, and extracted with dichloromethane (10 mL $\times$ 3). The organic phases were combined, washed with saturated brine (10 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by preparative chromatography to give **compound 68** (trifluoroacetate salt, 18.3 mg, 22% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 513$.

^1H NMR (400 MHz, CDCl_3) δ 8.58 (d, $J = 6.9$ Hz, 1H), 7.95 (s, 1H), 7.37 (d, $J = 8.1$ Hz, 2H), 7.16 (d, $J = 7.0$ Hz, 1H), 6.56 (brs, 2H), 4.62 – 4.52 (m, 1H), 4.14 – 3.77 (m, 3H), 3.72 (s, 3H), 3.71 – 3.65 (m, 1H), 3.46 – 3.36 (m, 1H), 3.12 – 3.11 (m, 2H), 2.01 – 2.88 (m, 1H), 2.78 – 2.68 (m, 1H), 2.61 (s, 6H).

Example 69

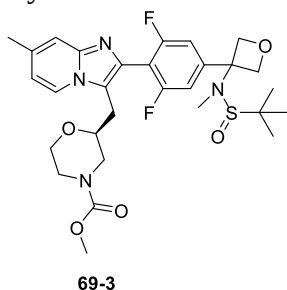


Step (1) Preparation of methyl (2S)-2-((2-(4-(3-((*tert*-butylsulfinyl)amino)oxetan-3-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



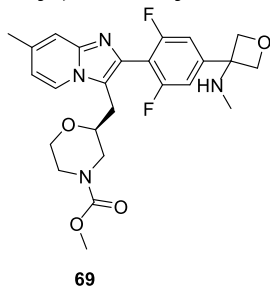
Intermediate 2-7 (300 mg, 0.63 mmol) was dissolved in tetrahydrofuran (5 mL) under nitrogen atmosphere, and *n*-butyllithium (0.32 mL, 0.76 mmol) was added dropwise at -78 °C in a dry ice-acetone bath. After the completion of the dropwise addition, the reaction system was reacted for 15 min, added dropwise with a solution of 2-methyl-*N*-(oxetan-3-ylidene)propane-2-sulfinamide (121 mg, 0.69 mmol) in tetrahydrofuran (1 mL), and reacted at -78 °C for 3 h. Then the reaction system was added with saturated aqueous ammonium chloride solution (2 mL) to quench the reaction, diluted with water (20 mL), and extracted with ethyl acetate (20 mL × 2). The organic phases were combined, washed with saturated brine (15 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 69-2** (60 mg, 70% purity, 12% yield) in the form of a brown solid. ESI-MS [M+H]⁺: 576.7.

Step (2) Preparation of methyl (2*S*)-2-((2-(4-(3-((*tert*-butylsulfinyl)(methyl)amino)oxetan-3-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 69-2 (60 mg, 0.10 mmol) was dissolved in anhydrous *N,N*-dimethylformamide (1 mL) under nitrogen atmosphere, and sodium hydride (10 mg, 0.25 mmol) was added in an ice water bath. The reaction system was stirred for 0.5 h, added dropwise with a solution of iodomethane (21 mg, 0.15 mmol) in *N,N*-dimethylformamide (0.5 mL), and stirred at 0 °C for 3 h. Then the reaction system was poured into ice water (15 mL), and extracted with ethyl acetate (15 mL × 2). The organic phases were combined, washed with saturated brine (10 mL × 3), dried over sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 69-3** (50 mg, 70% purity, 57% yield) in the form of a brown solid. ESI-MS [M +H]⁺: 590.8.

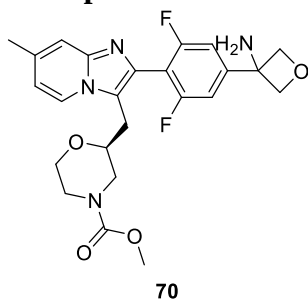
Step (3) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(3-(methylamino)oxetan-3-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 69-3 (50 mg, 0.085 mmol) was dissolved in dichloromethane (1.5 mL), and a solution of HCl in ethanol (0.1 mL, 0.40 mmol) was added dropwise at 0 °C. The reaction system was stirred for 15 min, poured into ice water (10 mL), adjusted to pH 8 with saturated sodium bicarbonate solution, and extracted with dichloromethane (10 mL × 2). The organic phases were combined, washed with saturated brine (10 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 69** (6.4 mg, 15.6% yield) in the form of a white solid. ESI-MS $[M+H]^+$: 486.8.

^1H NMR (400 MHz, CDCl_3) δ 8.26 (d, $J = 7.0$ Hz, 1H), 7.50 (s, 1H), δ 7.20 (d, $J = 8.5$ Hz, 2H), 6.74 (d, $J = 6.2$ Hz, 1H), 4.93 (d, $J = 6.7$ Hz, 2H), 4.82 (d, $J = 6.7$ Hz, 2H), 3.98 – 3.80 (m, 3H), 3.70 (s, 3H), 3.64 – 3.60 (m, 1H), 3.47 – 3.36 (m, 1H), 3.10 – 2.97 (m, 2H), 2.97 – 2.86 (m, 1H), 2.65 (dd, $J = 13.1, 10.7$ Hz, 1H), 2.47 (s, 3H), 2.35 (s, 3H).

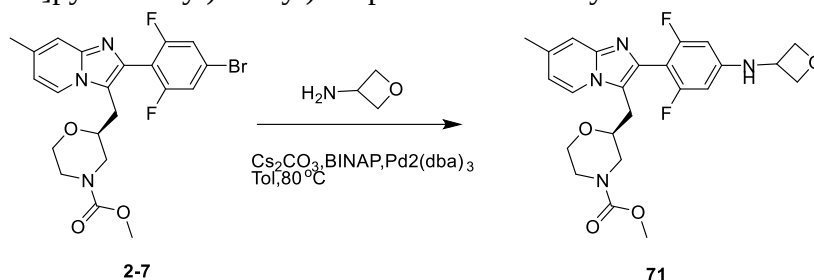
Example 70



The compound was prepared according to the preparation method as described in Example 63. LC-MS: $[M+H]^+ = 473.3$.

Example 71

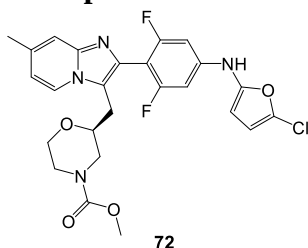
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(oxetan-3-ylamino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



5 **Intermediate 2-7** (40 mg, 0.08 mmol), 3-oxetane (7 mg, 0.096 mmol), cesium carbonate (78 mg, 0.24 mmol), tris(dibenzylideneacetone)dipalladium (7 mg, 0.008 mmol) and (±)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (10 mg, 0.016 mmol) were dissolved in anhydrous toluene (5 mL). The reaction system was reacted at 80 °C for 16 h under nitrogen atmosphere, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified
10 by preparative high performance liquid chromatography to give **compound 71** (25 mg, 62.5% yield) in the form of a white solid. MS $[M+H]^+ = 473.1$.

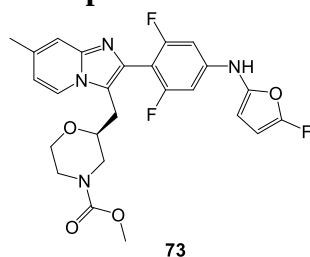
1H NMR (400 MHz, d6-DMSO) δ 8.35 (d, $J = 7.0$ Hz, 1H), 7.30 (s, 1H), 7.15 (d, $J = 6.5$ Hz, 1H), 6.78 (dd, $J = 7.1, 1.6$ Hz, 1H), 6.27 (d, $J = 10.6$ Hz, 2H), 4.88 (t, $J = 6.5$ Hz, 2H), 4.61 (dt, $J = 13.1, 6.7$ Hz, 1H), 4.44 (td, $J = 6.1, 2.1$ Hz, 2H), 3.73 - 3.64 (m, 3H), 3.56 (s, 3H), 3.51 -
15 3.45 (m, 1H), 3.25 (dt, $J = 11.7, 2.6$ Hz, 1H), 3.04 - 2.92 (m, 2H), 2.90 - 2.77 (m, 1H), 2.60 - 2.46 (m, 1H), 2.36 (s, 3H).

Example 72



The compound was prepared according to the preparation method as described in Example
20 25. LC-MS: $[M+H]^+ = 517.1$.

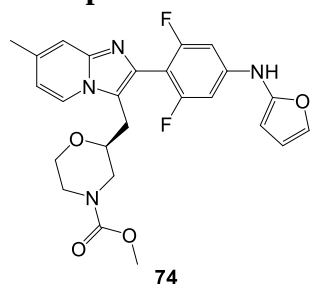
Example 73



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 501.2$.

5

Example 74

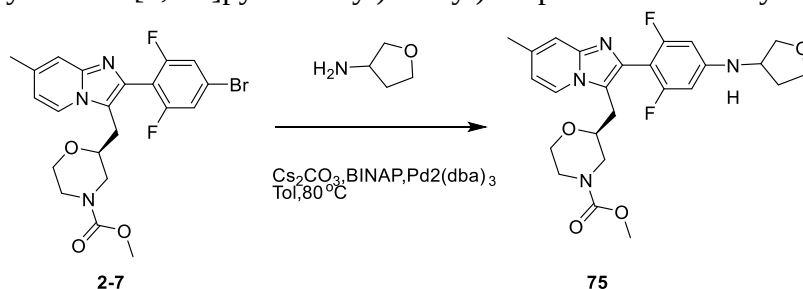


The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 483.3$.

Example 75

10

Preparation of methyl 2-((2-(2,6-difluoro-4-((tetrahydrofuran-3-yl)amino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 2-7 (50 mg, 0.10 mmol) was dissolved in toluene (5 mL) under nitrogen atmosphere, and oxolane-3-amine (11 mg, 0.12 mmol), cesium carbonate (98 mg, 0.30 mmol), tris(dibenzylideneacetone)dipalladium (9 mg, 0.01 mmol), and ($\pm$)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (13 mg, 0.02 mmol) were added. The reaction system was stirred at 80 °C overnight. Then the reaction system was filtered and concentrated. The residue was separated by preparative high performance liquid chromatography to give **compound 75** (14.7 mg, 30.0% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 487.1$.

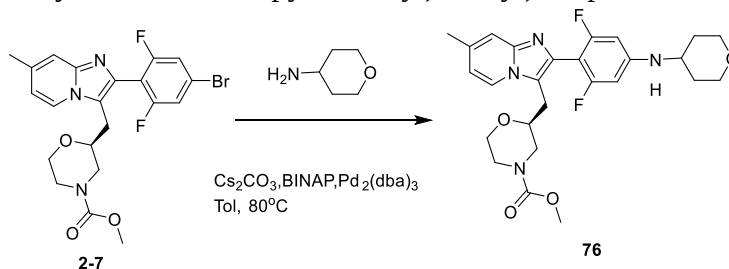
20

^1H NMR (400 MHz, MeOD) δ 8.47 (d, $J = 6.8$ Hz, 1H), 7.38 (s, 1H), 6.95 (d, $J = 6.5$ Hz, 1H), 6.37 (d, $J = 10.9$ Hz, 2H). 4.19 – 4.08 (m, 1H), 4.99 – 3.95 (m, 2H), 3.90 – 3.78 (m, 4H),

3.77 – 3.73 (m, 1H), 3.68 (s, 3H), 3.62 – 3.57 (m, 1H), 3.43 – 3.36 (m, 1H), 3.14 – 3.03 (m, 2H), 2.97 – 2.85 (m, 1H), 2.72 – 2.62 (m, 1H), 2.49 (s, 3H), 2.38 – 2.28 (m, 1H), 1.98 – 1.88 (m, 1H).

Example 76

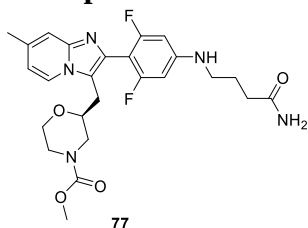
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-((tetrahydro-2H-pyran-4-yl)amino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 2-7 (80 mg, 0.17 mmol) was dissolved in toluene (5 mL) under nitrogen atmosphere, and tetrahydro-2H-pyran-4-amine (21 mg, 0.20 mmol), cesium carbonate (166 mg, 0.51 mmol), tris(dibenzylideneacetone)dipalladium (16 mg, 0.017 mmol), and (±)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (21 mg, 0.034 mmol) were added sequentially. The reaction system was stirred at 80 °C overnight. Then the reaction system was filtered and concentrated. The residue was separated by preparative high performance liquid chromatography to give **compound 76** (51.9 mg, 58.8% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 501.2$.

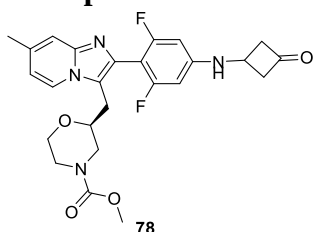
¹H NMR (400 MHz, DMSO) δ 8.36 (d, $J = 7.0$ Hz, 1H), 7.31 (s, 1H), 6.80 (d, $J = 6.6$ Hz, 1H), 6.41 – 6.34 (m, 1H), 6.38 (d, $J = 11.0$ Hz, 2H), 3.91 – 3.85 (m, 2H), 3.75 – 3.63 (m, 3H), 3.56 (s, 3H), 3.55 – 3.41 (m, 4H), 3.32 – 3.22 (m, 1H), 3.05 – 2.92 (m, 2H), 2.90 – 2.76 (m, 1H), 2.62 – 2.50 (m, 1H), 2.38 (s, 3H), 1.94 – 1.86 (m, 2H), 1.45 – 1.34 (m, 2H).

Example 77



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 502.2$.

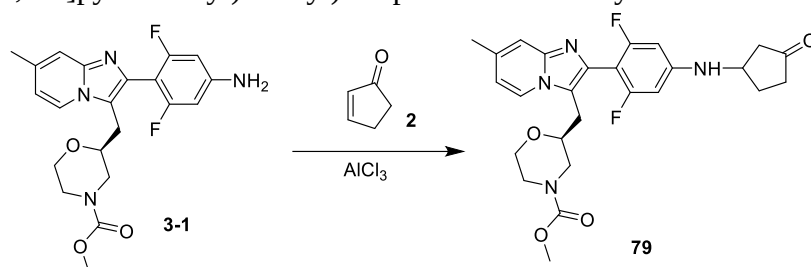
Example 78



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 485.2$.

5 Example 79

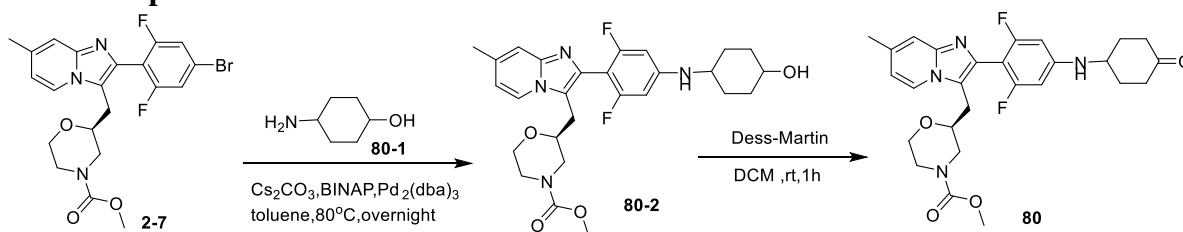
Preparation of methyl (2*S*)-2-((2-(2,6-difluoro-4-((3-carbonylcyclopentyl)amino)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



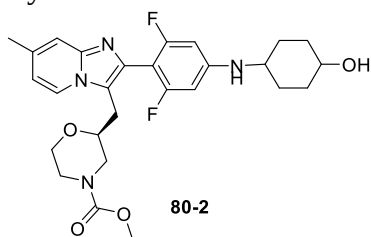
Intermediate 3-1 (60.0 mg, 0.14 mmol) was dissolved in cyclopent-2-enone (3 mL) under nitrogen atmosphere, and anhydrous aluminium trichloride (117.5 mg, 1.15 mmol) was added. The reaction system was reacted at room temperature for 18 h. Then the reaction system was filtered and concentrated. The crude product was separated by preparative high performance liquid chromatography, and lyophilized to give **compound 79** (29.7 mg, 42% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 499.3$.

¹H NMR (400 MHz, CDCl₃) δ 8.21 (brs, 1H), 7.46 (brs, 1H), 6.70 (brs, 1H), 6.27 (d, *J* = 9.9 Hz, 2H), 4.17 – 4.08 (m, 1H), 4.00 – 3.76 (m, 3H), 3.70 (s, 3H), 3.64 – 3.56 (m, 1H), 3.47 – 3.36 (m, 1H), 3.02 – 2.97 (m, 2H), 2.94 – 2.85 (m, 1H), 2.75 (dd, *J* = 18.2, 6.5 Hz, 1H), 2.64 (dd, *J* = 13.1, 10.7 Hz, 1H), 2.56 – 2.48 (m, 1H), 2.44 (s, 3H), 2.39 – 2.32 (m, 1H), 2.30 – 2.22 (m, 1H), 2.12 – 2.00 (m, 2H).

20 Example 80

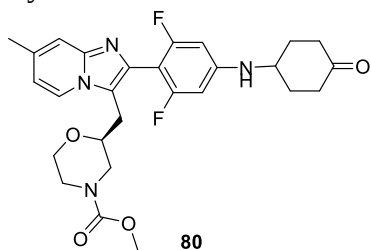


Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-((4-hydroxycyclohexyl)amino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



5 **Intermediate 2-7** (200 mg, 0.42 mmol) was dissolved in anhydrous toluene (3 mL) under nitrogen atmosphere, and 4-aminocyclohexanol (58 mg, 0.50 mmol), cesium carbonate (407 mg, 1.25 mmol), ($\pm$)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (52 mg, 0.084 mmol) and tris(dibenzylideneacetone)dipalladium (40 mg, 0.043 mmol) were added. The reaction system was reacted at 80 °C overnight. After the reaction was completed, the reaction system was cooled to
10 room temperature and filtered, and the filtrate was added with water (20 mL) and extracted with dichloromethane (30 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 80-2** (150 mg, 70% yield) in the form of a yellow oil. LC-MS: $[M+H]^+ =$
15 514.8.

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-((4-carbonylcyclohexyl)amino)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

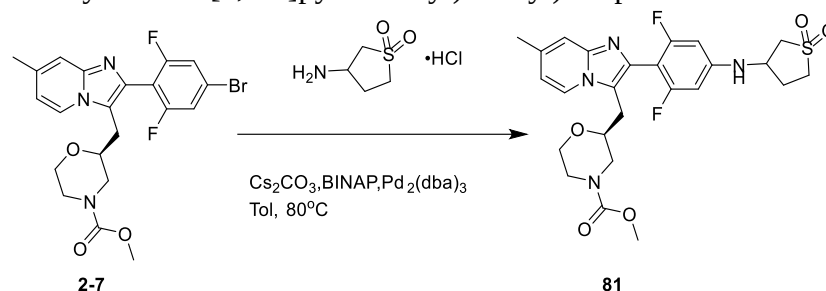


20 **Intermediate 80-2** (150 mg, 0.29 mmol) was dissolved in anhydrous dichloromethane (2 mL) under nitrogen atmosphere, and Dess-Martin periodinane (185 mg, 0.44 mmol) was added. The reaction system was reacted at room temperature for 1 h. Then the reaction system was added with water (20 mL), and extracted with dichloromethane (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL), dried over sodium sulfate, filtered and
25 concentrated by rotary evaporation to give a crude product. The crude product was purified by preparative high performance liquid chromatography to give **compound 80** (20.2 mg, 20.3% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 512.8$.

^1H NMR (400 MHz, MeOD) δ 8.82 – 8.78 (m, 1H), 7.65 – 7.64 (m, 1H), 7.38 – 7.36 (m, 1H), 6.53 (d, J = 11.7 Hz, 1H), 6.46 – 6.41 (m, 1H), 3.98 – 3.95 (m, 1H), 3.92 – 3.75 (m, 3H), 3.70 (s, 3H), 3.39 – 3.35 (m, 1H), 3.25 – 3.18 (m, 2H), 2.99 – 2.90 (m, 1H), 2.80 – 2.72 (m, 1H), 2.63 (s, 3H), 2.59 – 2.55 (m, 1H), 2.48 – 2.44 (m, 1H), 2.34 – 2.29 (m, 1H), 1.97 – 1.95 (m, 2H), 1.86 – 1.49 (m, 4H).

Example 81

Preparation of methyl (2S)-2-((2-(4-((1,1-dihydroxytetrahydrothiophen-3-yl)amino)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 2-7 (80 mg, 0.16 mmol), 3-aminotetrahydrothiophene 1,1-disulfone hydrochloride (43 mg, 0.24 mmol), cesium carbonate (217 mg, 0.64 mmol), ($\pm$)-2,2'-bis-(diphenylphosphino)-1,1'-binaphthalene (21 mg, 0.03 mmol), and tris(dibenzylideneacetone)dipalladium (15 mg, 0.02 mmol) were dissolved in toluene (3 mL) under nitrogen atmosphere. The reaction system was heated to 80 °C, and reacted overnight. Then the reaction system was cooled to room temperature, added with water (20 mL) to quench the reaction, and extracted with dichloromethane (15 mL $\times$ 3). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by preparative high performance liquid chromatography to give **compound 81** (32.4 mg, 36% yield) in the form of a yellow solid, LC-MS: $[\text{M}+\text{H}]^+ = 535.1$.

^1H NMR (400 MHz, CDCl_3) δ 8.26 (s, 1H), 7.55 (d, J = 7.4 Hz, 1H), 6.77 (d, J = 6.9 Hz, 1H), 6.27 (d, J = 9.8 Hz, 2H), 4.37 – 4.30 (m, 1H), 4.01 – 3.79 (m, 3H), 3.70 (s, 3H), 3.63 – 3.56 (m, 1H), 3.55–3.48 (m, 1H), 3.45 – 3.32 (m, 2H), 3.25 – 3.16 (m, 1H), 3.13 – 3.05 (m, 1H), 3.03–2.86 (m, 3H), 2.67 – 2.62 (m, 2H), 2.46 (s, 3H), 2.45 – 2.36 (m, 1H).

Cc1ccc2nc(C(F)(F)c3cc(NC(=O)C4CCCCC4C(=O)O)cc(F)c3)[nH]2c1[C@H]3CCOC(=O)N3

82

5

Cc1ccc2nc3c(cc12)C(c4cc(F)c(NC(=O)C5=CCCCC5=O)c(F)c4)[C@H]6CCN(C6)C(=O)OC

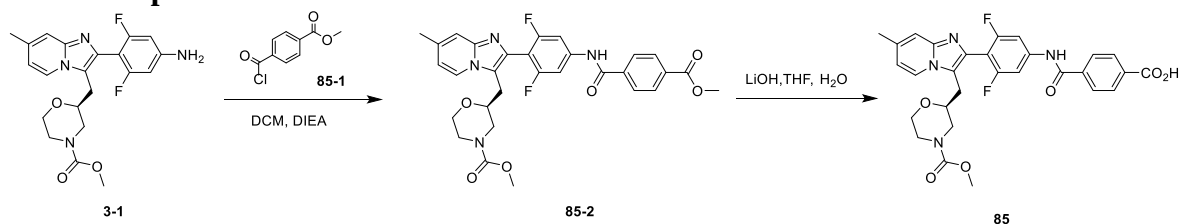
83

10

COC(=O)N1CC[C@H](C2=CN3C(=N2)C(=C3C=C4C(=C(C=C4)N(C)C)C(F)=C(F)C(=C4)NC(=O)C5=CC=CC=C5)C=C4)O1

84

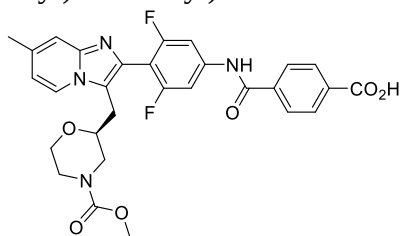
Example 85

COC(=O)Nc1cc(F)c(cc1F)C2=CN3C=CC=C(C)N3C[C@H]2C4CCN(C4)C(=O)OC

85-2

Intermediate 3-1 (80 mg, 0.19 mmol) and diisopropylethylamine (74 mg, 0.57 mmol) were dissolved in dichloromethane (10 mL). The reaction system was cooled to 0 °C, added dropwise with a solution of methyl 4-(chloroformyl)benzoate (75 mg, 0.38 mmol) in dichloromethane (0.2 mL), and stirred at room temperature for 12 h. Then the reaction system was diluted with dichloromethane (10 mL) and washed with saturated brine (10 mL). The organic phase was dried over anhydrous sodium sulfate, filtered, and concentrated to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 85-2** (100 mg, 49% yield) in the form of a white solid. MS $[M+H]^+ = 579.2$.

Step (2) Preparation of (S)-4-((3,5-difluoro-4-(3-((4-(carbomethoxy<methoxycarbonyl>morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)carbamoyl)benzoic acid

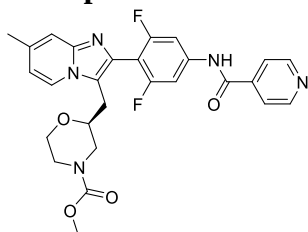


85

Intermediate 85-2 (50 mg, 0.09 mmol) was dissolved in tetrahydrofuran (5 mL) and water (1 mL), and lithium hydroxide monohydrate (22 mg, 0.55 mmol) was added. The reaction system was stirred at room temperature for 1 h. After the reaction was completed, the reaction system was adjusted to pH 4-5 with diluted hydrochloric acid (1 N), and concentrated by rotary evaporation to give a crude product. The crude product was purified by preparative high performance liquid chromatography to give **compound 85** (35 mg, 59% yield) in the form of a white solid. MS $[M+H]^+ = 565.2$.

¹H NMR (400 MHz, d₆-DMSO) δ 13.24 (s, 1H), 10.84 (s, 1H), 8.42 (d, $J = 6.9$ Hz, 1H), 8.10 (q, $J = 7.0$ Hz, 4H), 7.69 (d, $J = 9.9$ Hz, 2H), 7.35 (s, 1H), 6.83 (d, $J = 7.1$ Hz, 1H), 3.77 – 3.59 (m, 3H), 3.56 (s, 3H), 3.52 – 3.46 (m, 1H), 3.30 – 3.22 (m, 1H), 3.08 – 3.00 (m, 2H), 2.90 – 7.74 (m, 1H), 2.60 – 2.48 (m, 1H), 2.39 (s, 3H).

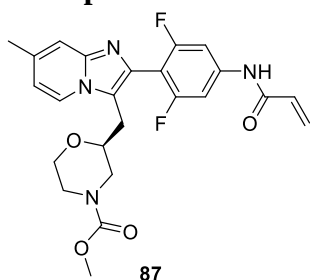
Example 86



86

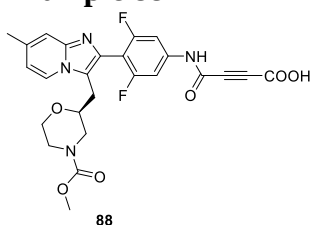
The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 522.2$.

Example 87



5 The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 471.2$.

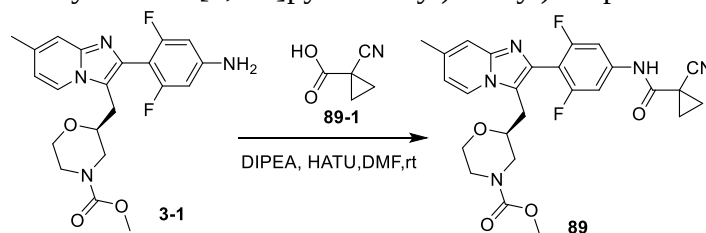
Example 88



10 The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 513.3$.

Example 89

Preparation of methyl (S)-2-((2-(4-(1-cyanocyclopropane-1-carboxamido<oxalylamino>)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

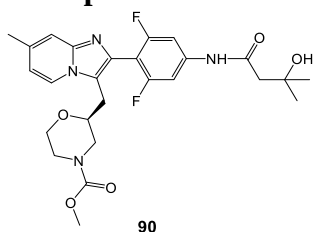


15 1-cyanocyclopropane-1-carboxylic acid (52 mg, 0.47 mmol) and 2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (180 mg, 0.47 mmol) were dissolved in anhydrous *N,N*-dimethylformamide (2 mL) under nitrogen atmosphere. The reaction system was reacted at room temperature for 10 min, added with *N,N*-diisopropylethylamine (162 mg, 1.25 mmol) and
20 **intermediate 3-1** (130 mg, 0.31 mmol), and reacted at room temperature overnight. Then the reaction system was added with water (20 mL), and extracted with ethyl acetate (30 mL $\times$ 3). The organic phases were combined, washed with saturated brine (30 mL $\times$ 3), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude

product. The crude product was purified by high performance liquid chromatography to give **compound 89** (35.8 mg, 23% yield) in the form of a white solid.

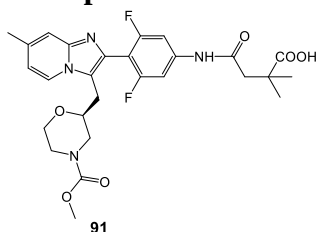
LC-MS: $[M+H]^+ = 510.6$. ^1H NMR (400 MHz, MeOD): δ 8.83 (d, $J = 7.1$ Hz, 1H), 7.69 (s, 1H), 7.67 (d, $J = 10.1$ Hz, 2H), 7.40 (dd, $J = 7.1, 1.5$ Hz, 1H), 3.98 (d, $J = 12.8$ Hz, 1H), 3.83 (d, $J = 13.6$ Hz, 1H), 3.76 (d, $J = 9.2$ Hz, 1H), 3.70 (s, 3H), 3.70 – 3.65 (m, 1H), 3.37 – 3.34 (m, 1H), 3.24 (d, $J = 5.3$ Hz, 2H), 3.00 – 2.85 (m, 1H), 2.80 – 2.67 (m, 1H), 2.64 (s, 3H), 1.81 – 1.76 (m, 2H), 1.75 – 1.70 (m, 2H).

Example 90



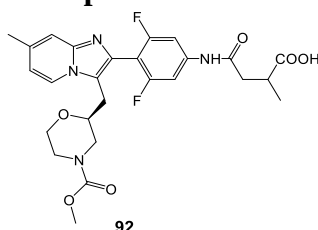
The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 517.2$.

Example 91



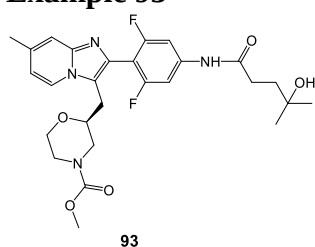
The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 545.4$.

Example 92



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 531.2$.

Example 93

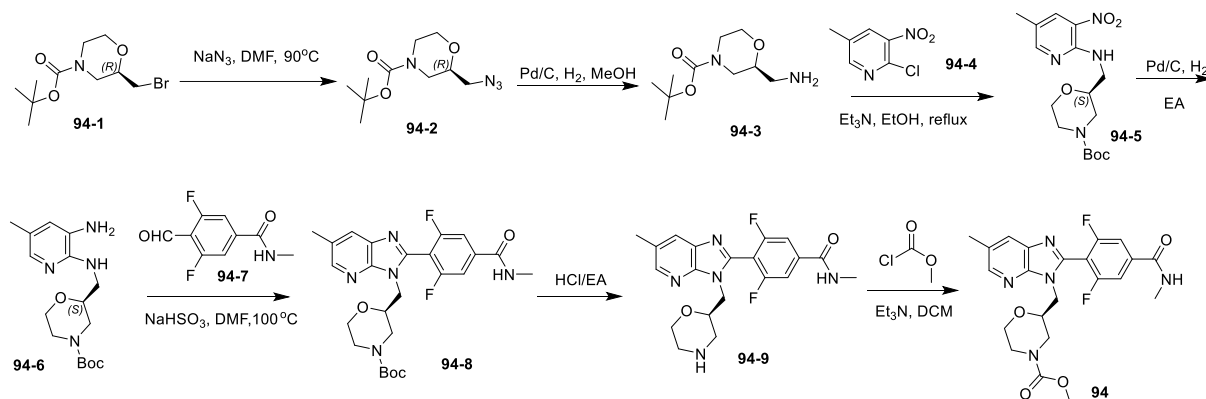


93

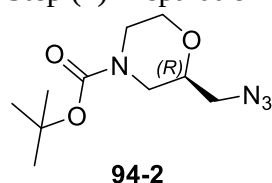
The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 531.4$.

5

Example 94



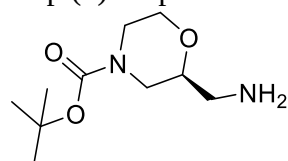
Step (1) Preparation of *tert*-butyl (*R*)-2-(azidomethyl)morpholine-4-carboxylate



94-2

Tert-butyl (*R*)-2-(bromomethyl)morpholine-4-carboxylate (1.0 g, 3.58 mmol) was dissolved in *N,N*-dimethylformamide (8 mL), and sodium azide (700 mg, 10.75 mmol) was added. The reaction system was heated to 90 °C for 8 h. Then the reaction system was cooled to room temperature, added with water (40 mL) to quench the reaction, and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (15 mL × 2), dried over anhydrous sodium sulfate, and concentrated. The residue was separated by column chromatography to give **intermediate 94-2** (738 mg, 92% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 243.2$.

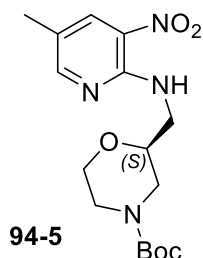
Step (2) Preparation of *tert*-butyl (*S*)-2-(aminomethyl)morpholine-4-carboxylate



94-3

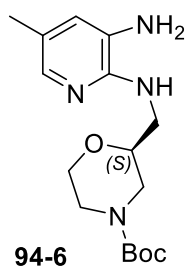
Intermediate 94-2 (738 mg, 3.05 mmol) was dissolved in methanol (10 mL), and wet palladium on carbon catalyst (200 mg, 10% w/w) was added. The reaction system was purged with hydrogen and reacted at room temperature for 2.5 h. Then the reaction system was filtered and concentrated to give **intermediate 94-3** (580 mg, 82% yield) in the form of a light yellow oil. LC-MS: $[M+H]^+ = 217.2$.

Step (3) Preparation of *tert*-butyl (S)-2-(((5-methyl-3-nitropyridin-2-yl)amino)methyl)morpholine-4-carboxylate



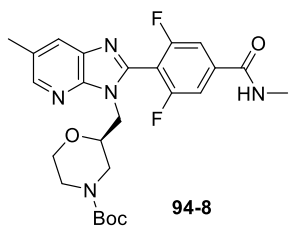
Intermediate 94-3 (580 mg, 2.68 mmol) was dissolved in ethanol (10 mL), and 2-chloro-5-methyl-3-nitropyridine (509 mg, 2.95 mmol) and triethylamine (895 mg, 8.84 mmol) were added. The reaction system was heated to reflux for 24 h. Then the reaction system was concentrated by rotary evaporation. The residue was separated by column chromatography (PE/EA = 4/1) to give **intermediate 94-5** (710 mg, 75% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 353.1$.

Step (4) Preparation of *tert*-butyl (S)-2-(((3-amino-5-methylpyridin-2-yl)amino)methyl)morpholine-4-carboxylate



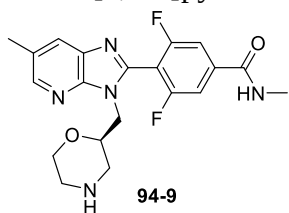
Intermediate 94-5 (326 mg, 0.96 mmol) was dissolved in ethyl acetate (20 mL), and wet palladium on carbon catalyst (50 mg, 10% w/w) was added. The reaction system was purged with hydrogen and stirred at room temperature for 2.5 h. Then the reaction system was filtered and concentrated to give **intermediate 94-6** (266 mg, 81% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 323.2$.

Step (5) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methyl-3H-imidazo[4,5-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate



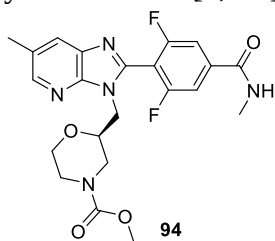
Intermediate 94-6 (266 mg, 0.82 mmol), sodium bisulfite (512 mg, 4.92 mmol), and 3,5-difluoro-4-formyl-*N*-methylbenzamide (213 mg, 1.07 mmol) were dissolved in *N,N*-dimethylformamide (8 mL). The reaction system was heated to 100 °C and reacted for 15 h. After the reaction was completed, the reaction system was added with water (30 mL) to quench the reaction, and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The reaction system was separated by column chromatography (EA) to give **intermediate 94-8** (210 mg, 46% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 502.1$.

Step (6) Preparation of (*R*)-3,5-difluoro-*N*-methyl-4-(6-methyl-3-(morpholin-2-yl-methyl)-3*H*-imidazo[4,5-*b*]pyridin-2-yl)benzamide



Intermediate 94-8 (210 mg, 0.52 mmol) was dissolved in ethyl acetate (4 mL), and a solution of HCl in ethyl acetate (1 mL, 4 N) was added dropwise. The reaction system was stirred at room temperature for 2.5 h. Then the reaction system was concentrated to give **intermediate 94-9** (hydrochloride, 222 mg, crude product) in the form of a yellow solid, which was used directly in the next step. LC-MS: $[M+H]^+ = 402.1$.

Step (7) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(methylcarbamoyl)phenyl)-6-methyl-3*H*-imidazo[4,5-*b*]pyridin-3-yl)methyl)morpholine-4-carboxylate

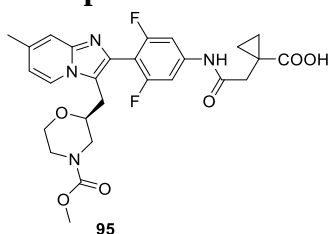


Intermediate 94-9 (222 mg, 0.52 mmol) was dissolved in anhydrous dichloromethane (10 mL), and triethylamine (157 mg, 1.56 mmol) was added. Under nitrogen atmosphere, the reaction system was cooled to 0 °C, added dropwise with methyl chloroformate (59 mg, 0.62 mmol), and reacted for 3 h. Then the reaction system was diluted with dichloromethane (20 mL)

and washed with water (20 mL). The aqueous phase was extracted with dichloromethane (15 mL $\times$ 2). The organic phases were combined, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by preparative high performance liquid chromatography to give **compound 94** (64 mg, 27% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 460.8$.

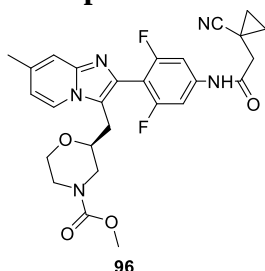
¹H NMR (400 MHz, CDCl₃): δ 8.39 (s, 1H), 8.03 (s, 1H), 7.56 (brs, 2H), 7.02 (brs, 1H), 4.50-4.41 (m, 1H), 4.25-4.15 (m, 1H), 4.12-3.96 (m, 1H), 3.85-3.65 (m, 5H), 3.65-3.55 (m, 1H), 3.30-3.20 (m, 1H), 3.07 (s, 3H), 2.85-2.70 (m, 1H), 2.57 (s, 3H), 2.56-2.46 (m, 1H).

Example 95



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 543.3$.

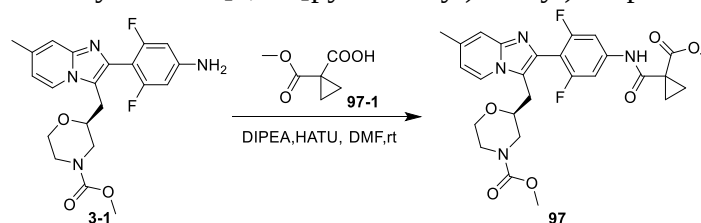
Example 96



The compound was prepared according to the preparation method as described in Example 25. LC-MS: $[M+H]^+ = 524.3$.

Example 97

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methoxycarbonyl)cyclopropane-1-carboxamido)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

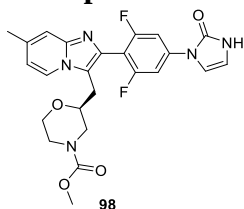


Intermediate 97-1 (31 mg, 0.22 mmol) and 2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (82 mg, 0.22 mmol) were dissolved in anhydrous *N,N*-dimethylformamide (1.0 mL) under nitrogen atmosphere. The reaction system was reacted at room temperature for 10 min, added with *N,N*-diisopropylethylamine (75 mg, 0.58 mmol) and

intermediate 3-1 (60 mg, 0.14 mmol), and reacted at room temperature overnight. Then the reaction system was added with water (20 mL) to quench the reaction, and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 97** (11.7 mg, 15% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 543.7$.

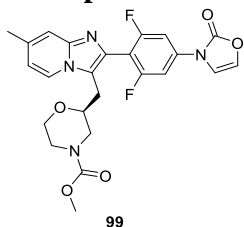
^1H NMR (400 MHz, MeOD) δ 8.84 (d, $J = 7.1$ Hz, 1H), 7.70 (s, 1H), 7.64 (d, $J = 10.1$ Hz, 2H), 7.41 (dd, $J = 7.1, 1.4$ Hz, 1H), 4.01 – 3.97 (m, 1H), 3.85 – 3.82 (m, 1H), 3.80 (s, 3H), 3.78 – 3.71 (m, 2H), 3.70 (s, 3H), 3.39 – 3.35 (m, 1H), 3.28 – 3.21 (m, 2H), 2.98 – 2.90 (m, 1H), 2.80 – 2.70 (m, 1H), 2.65 (s, 3H), 1.71 – 1.64 (m, 4H).

Example 98



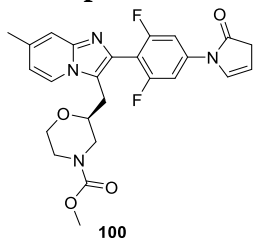
The compound was prepared according to the preparation method as described in Example 5. LC-MS: $[M+H]^+ = 484.2$.

Example 99



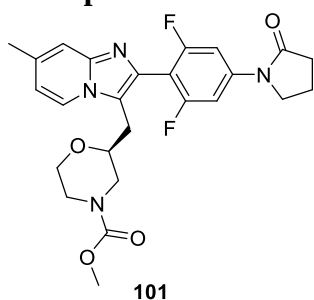
The compound was prepared according to the preparation method as described in Example 5. LC-MS: $[M+H]^+ = 485.3$.

Example 100



The compound was prepared according to the preparation method as described in Example 5. LC-MS: $[M+H]^+ = 483.4$.

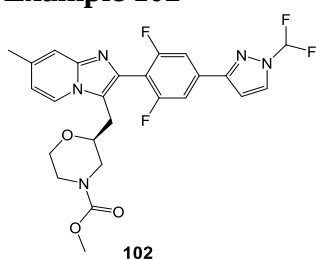
Example 101



The compound was prepared according to the preparation method as described in Example 5. LC-MS: $[M+H]^+ = 485.2$.

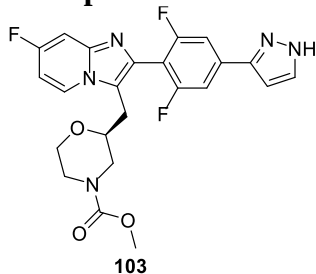
5

Example 102



The compound was prepared according to the preparation method as described in Example 4. LC-MS: $[M+H]^+ = 518.2$.

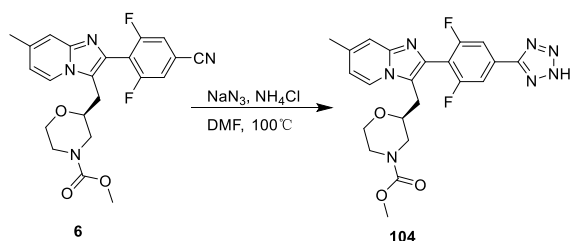
Example 103



10

The compound was prepared according to the preparation method as described in Example 15. LC-MS: $[M+H]^+ = 472.5$.

Example 104



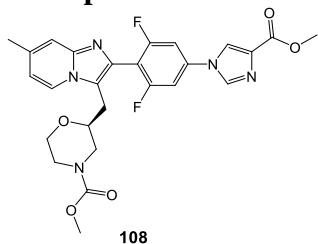
15

Compound 6 (540 mg, 1.27 mmol) was dissolved in DMF (15 mL), and a magnetic stirrer was added, followed by the addition of NaN_3 (247.7 mg, 3.81 mmol) and NH_4Cl (271 mg, 5.08 mmol). The reaction system was slowly heated to 100 °C and reacted overnight. After the reaction was completed as detected by LCMS, the reaction system was dissolved with water, and

was purified by preparative high performance liquid chromatography to give **compound 107** (10 mg, 33% yield) in the form of a white solid. MS $[M+H]^+ = 512.1$.

^1H NMR (400 MHz, d_6 -DMSO) δ 8.56 (brs, 1H), 8.44 (d, $J = 5.0$ Hz, 1H), 7.89 (brs, 1H), 7.37 (brs, 1H), 6.84 (d, $J = 6.3$ Hz, 1H), 3.74 - 3.60 (m, 3H), 3.56 (s, 3H), 3.51 - 3.45 (m, 1H), 3.26 - 3.21 (m, 1H), 3.10 - 2.98 (m, 2H), 2.88 - 2.75 (m, 1H), 2.60 - 2.48 (m, 1H), 2.38 (s, 3H).

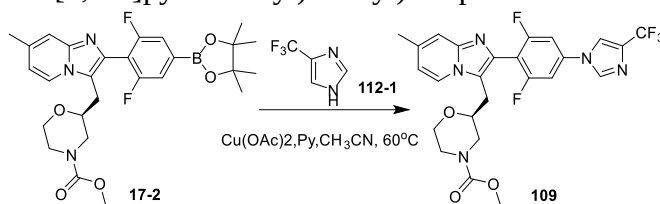
Example 108



The compound was prepared according to the preparation method as described in Example 50. LC-MS: $[M+H]^+ = 526.2$.

Example 109

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-(trifluoromethyl)-1H-imidazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

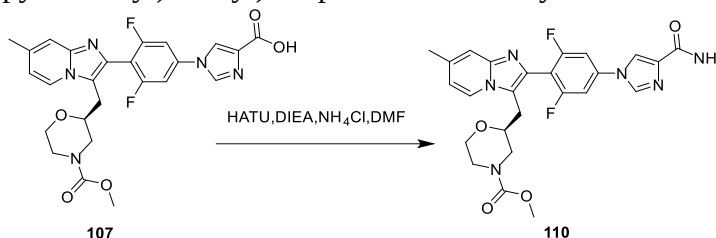


Intermediate 17-2 (130 mg, 0.25 mmol), 4-(trifluoromethyl)-1H-imidazole (40 mg, 0.3 mmol), copper(II) acetate (67 mg, 0.37 mmol), and pyridine (39 mg, 0.5 mmol) were dissolved in acetonitrile (2 mL). The reaction system was heated to 60 °C and reacted overnight. Then the reaction system was cooled to room temperature, diluted with water (10 mL), and extracted with dichloromethane (10 mL $\times$ 3). The organic phases were combined, washed with saturated brine (10 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 109** (13 mg, 9% yield) in the form of a white solid.

LC-MS: $[M+H]^+ = 536.4$. ^1H NMR (400 MHz, CDCl_3) : δ 8.29 (d, $J = 7.3$ Hz, 1H), 7.99 (s, 1H), 7.70 (s, 1H), 7.61 (s, 1H), 7.19 (d, $J = 7.2$ Hz, 1H), 6.83 (d, $J = 7.0$ Hz, 1H), 4.10 - 3.80 (m, 3H), 3.71 (s, 3H), 3.65 - 3.57 (m, 1H), 3.46 - 3.35 (m, 1H), 3.09 - 2.95 (m, 3H), 2.70 - 2.62 (m, 1H), 2.49 (s, 3H).

Example 110

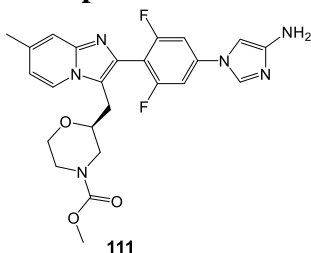
Preparation of methyl (S)-2-((2-(4-(4-carbamoyl-1*H*-imidazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



5 **Compound 107** (30 mg, 0.06 mmol), HATU (34 mg, 0.09 mmol) and *N,N*-diisopropylethylamine (23 mg, 0.18 mmol) were dissolved in *N,N*-dimethylformamide (2 mL). The reaction system was stirred at room temperature for 5 min, added with anhydrous ammonium chloride (10 mg, 0.18 mmol), and stirred at room temperature for another 1 h. Then the reaction system was added with saturated brine (15 mL) to quench the reaction, and extracted with ethyl acetate (10 mL × 2). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by silica gel column chromatography to give **compound 110** (28 mg, 83% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 511.1$.

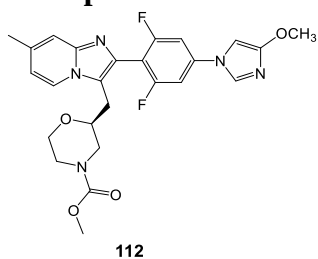
15 ¹H NMR (400 MHz, MEOD) δ 8.43 (d, *J* = 7.2 Hz, 1H), 8.36 (d, *J* = 1.3 Hz, 1H), 8.27 (d, *J* = 1.3 Hz, 1H), 7.62 (d, *J* = 8.1 Hz, 2H), 7.37 (s, 1H), 6.90 (dd, *J* = 7.1, 1.6 Hz, 1H), 3.89 – 3.76 (m, 3H), 3.67 (s, 3H), 3.63 – 3.58 (m, 1H), 3.41 – 3.35 (m, 1H), 3.16 – 3.10 (m, 2H), 2.98 – 2.82 (m, 1H), 2.72 – 2.58 (m, 1H), 2.48 (s, 3H).

Example 111



20 The compound was prepared according to the preparation method for compound 5. MS $[M+H]^+ = 483.3$.

Example 112

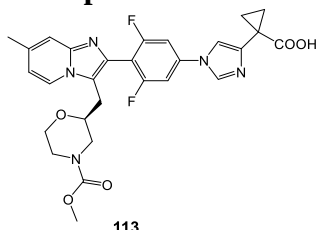


112

The compound was prepared according to the preparation method for compound 5. MS $[M+H]^+ = 498.2$.

5

Example 113



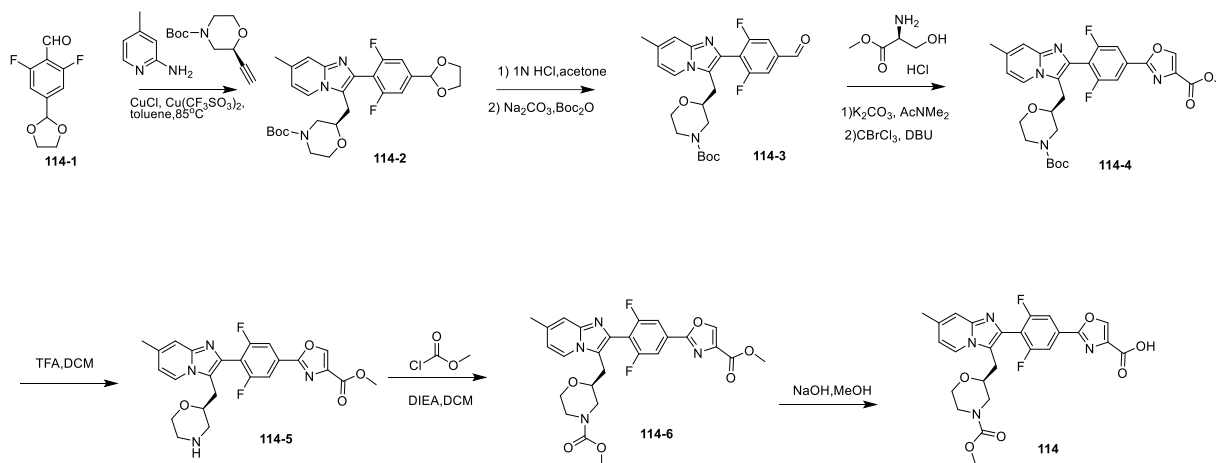
113

The compound was prepared according to the preparation method for compound 5. MS $[M+H]^+ = 552.2$.

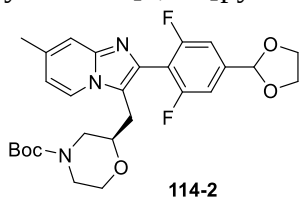
Example 114

10

Preparation of (S)-2-((3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)oxazole-4-carboxylic acid



Step (1) Preparation of methyl (R)-2-((2-(4-(1,3-dioxolan-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

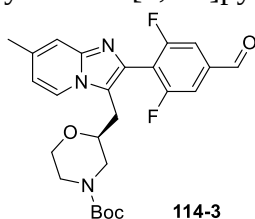


114-2

15

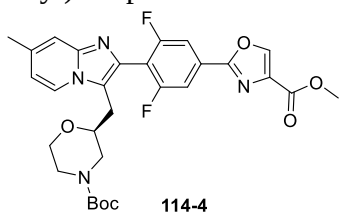
4-(1,3-dioxolan-2-yl)-2,6-difluorobenzaldehyde (600 mg, 2.8 mmol), 4-methylpyridin-2-amine (302.8 mg, 2.8 mmol), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (595 mg, 2.8 mmol), copper(I) chloride (84 mg, 0.84 mmol), and copper(II) trifluoromethanesulfonate (304 mg, 0.84 mmol) were dissolved in toluene (10 mL). The reaction system was stirred at 85 °C for 30 min, added dropwise with *N,N*-dimethylacetamide (0.1 mL), and stirred at 85 °C overnight. Then the reaction system was concentrated under reduced pressure. The crude product was purified by silica gel column chromatography to give **intermediate 114-2** (1.2 g, 55.36% yield) in the form of a yellow solid. MS $[M+H]^+ = 516.2$.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-formylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 114-2 (1.2 g, 2.33 mmol) was dissolved in acetone (10 mL) at room temperature, and diluted hydrochloric acid (10 mL, 1 N) was added. The reaction system was stirred overnight. After the reaction was completed, the reaction system was adjusted to pH 8-9 with saturated sodium carbonate solution, added with BOC-anhydride (766 mg, 3.5 mmol), and stirred at room temperature for 5 h. Then the reaction system was extracted with ethyl acetate (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 114-3** (563 mg, 51.3% yield) in the form of a yellow solid. MS $[M+H]^+ = 471.2$.

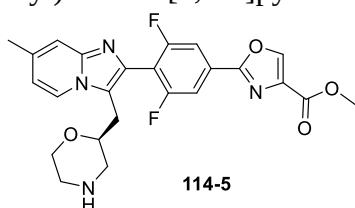
Step (3) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(4-(methoxycarbonyl)oxazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 114-3 (495 mg, 1.05 mmol) was dissolved in *N,N*-dimethylacetamide (4 mL) under nitrogen atmosphere, and methyl (2S)-2-amino-3-hydroxypropionate acid (163 mg, 1.05 mmol) and potassium carbonate (290 mg, 2.10 mmol) were added. The reaction system was stirred at room temperature for 18 h. Then the reaction system was cooled to 0 °C, added

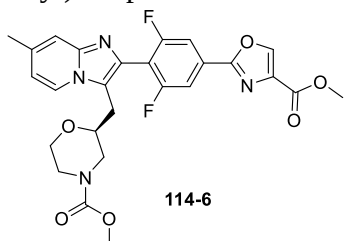
with bromotrichloromethane (625 mg, 3.15 mmol) and 1,8-diaza-cyclo[5.4.0]undec-7-ene (480 mg, 3.15 mmol), stirred at 0 °C for 2 h, and stirred at room temperature for 24 h. The resulting reaction system was added with water (20 mL) to quench the reaction and extracted with diethyl ether (20 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 114-4** (231 mg, 37.2% yield) in the form of a yellow solid. MS $[M+H]^+ = 569.2$.

Step (4) Preparation of methyl (S)-2-(3,5-difluoro-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-a]pyridin-2-yl)phenyl)oxazole-4-carboxylate



Intermediate 114-4 (231, 0.41 mmol) was dissolved in dichloromethane (4 mL), and trifluoroacetic acid (1 mL) was added. The reaction system was stirred at room temperature for 2 h. Then the reaction system was concentrated by rotary evaporation under reduced pressure, diluted with dichloromethane (30 mL), and slowly added with saturated sodium bicarbonate solution (20 mL) while stirring, followed by liquid separation. The aqueous phase was extracted with dichloromethane (20 mL). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 114-5** (195 mg, 97.6% yield) in the form of a brown solid. MS $[M+H]^+ = 469.1$.

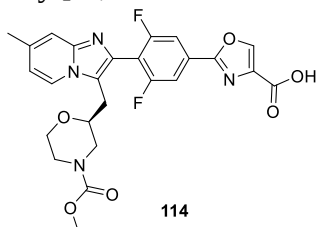
Step (5) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-(carbomethoxy<methoxycarbonyl>)oxazol-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 114-5 (150 mg, 0.30 mmol) and diisopropylethylamine (160 mg, 1.20 mmol) were dissolved in DCM (10 mL) at 0 °C under nitrogen atmosphere. The reaction system was stirred for 10 min, and added dropwise with methyl chloroformate (34 mg, 0.36 mmol). After the completion of the dropwise addition, the reaction was reacted for 2 h. Then the reaction system was concentrated by rotary evaporation. The crude product was purified by

silica gel column chromatography to give **intermediate 114-6** (110 mg, 66.7% yield) in the form of a white solid. MS $[M+H]^+ = 527.1$.

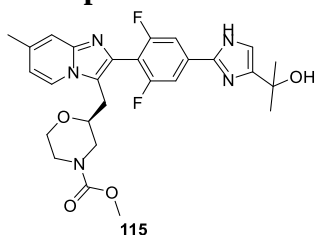
Step (6) Preparation of 2-[3,5-difluoro-4-(3-{[(2*S*)-4-(carbomethoxy<methoxycarbonyl>morpholin-2-yl)methyl}-7-methylimidazo[1,2-*a*]pyridin-2-yl)phenyl]-1,3-oxazole-4-carboxylic acid



Intermediate 114-6 (80 mg, 0.15 mmol) was dissolved in methanol (2 mL), and sodium hydroxide solution (1.5 mL, 1 N) was added. The reaction system was reacted at room temperature for 2 h. Then the reaction system was adjusted to pH 5 with diluted hydrochloric acid and concentrated. The residue was separated by preparative high performance liquid chromatography to give **compound 114** (33 mg, 40% yield) in the form of a white solid. LC-MS $[M+H]^+ = 513.1$.

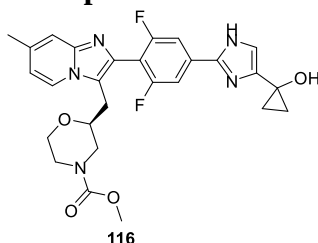
¹H NMR (400 MHz, DMSO) δ 13.24 (brs, 1H), 8.99 (s, 1H), 8.47 (d, *J* = 7.1 Hz, 1H), 7.81 (d, *J* = 7.5 Hz, 2H), 7.39 (s, 1H), 6.88 (d, *J* = 7.0 Hz, 1H), 3.82 – 3.73 (m, 1H), 3.69 – 3.60 (m, 2H), 3.56 (s, 3H), 3.49 (s, 1H), 3.27 – 3.18 (m, 1H), 3.14 – 3.00 (m, 2H), 2.87 – 2.75 (m, 1H), 2.55 – 2.45 (m, 1H), 2.40 (s, 3H).

Example 115



The compound was prepared according to the preparation method as described in Example 114. LC-MS: $[M+H]^+ = 493.2$.

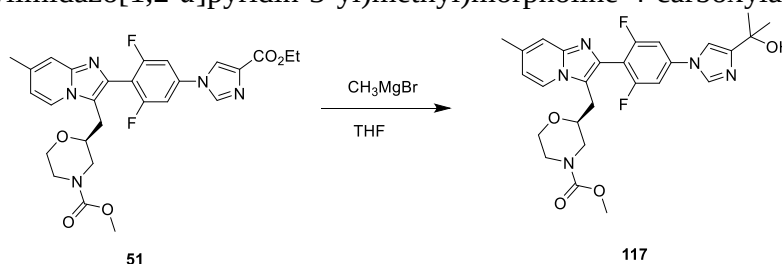
Example 116



The compound was prepared according to the preparation method as described in Example 114. LC-MS: $[M+H]^+ = 493.2$.

Example 117

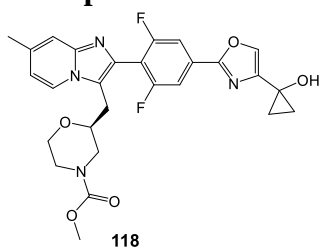
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-(2-hydroxypropan-2-yl)-1H-imidazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



5 **Compound 51** (30 mg, 0.06 mmol) was dissolved in tetrahydrofuran (2 mL), and a solution of methyl magnesium bromide in *n*-hexane (3 M, 0.06 mL, 0.18 mmol) was added dropwise under nitrogen atmosphere in an ice water bath. After the completion of the dropwise addition, the reaction system was stirred at room temperature for 1 h. Then the reaction system was added with saturated ammonium chloride solution to quench the reaction, and concentrated by rotary
10 evaporation to give a crude product. The crude product was purified by preparative high performance liquid chromatography to give **compound 117** (5 mg, 17% yield) in the form of a white solid. MS $[M+H]^+ = 526.1$.

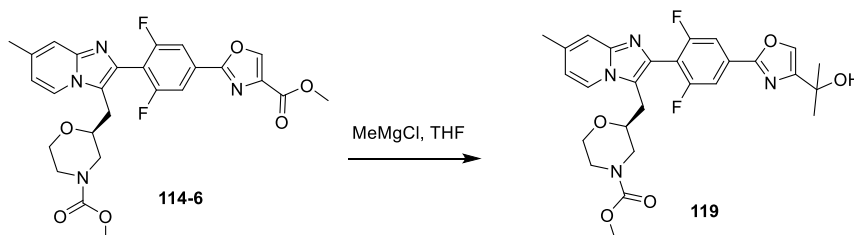
1H NMR (400 MHz, d6-DMSO) δ 8.43 (d, $J = 7.0$ Hz, 1H), 8.38 (d, $J = 1.2$ Hz, 1H), 7.73 (d, $J = 8.8$ Hz, 2H), 7.71 (d, $J = 1.4$ Hz, 1H), 7.36 (s, 1H), 6.85 (dd, $J = 7.1, 1.5$ Hz, 1H), 4.84 (s,
15 1H), 3.78 - 3.60 (m, 3H), 3.56 (s, 3H), 3.52 - 3.46 (m, 1H), 3.29 - 3.21 (m, 1H), 3.08 - 3.02 (m, 2H), 2.90 - 2.75 (m, 1H), 2.60 - 2.46 (m, 1H), 2.39 (s, 3H), 1.45 (s, 6H).

Example 118



The compound was prepared according to the preparation method as described in Example
20 114. MS $[M+H]^+ = 525.2$.

Example 119

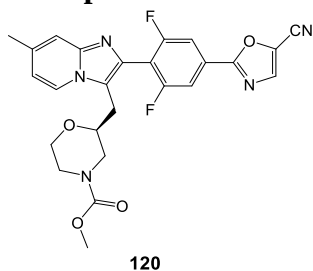


Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-(2-hydroxypropyl-2-yl)oxazol-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

Intermediate 114-6 (150 mg, 0.29 mmol) was dissolved in tetrahydrofuran (8 mL), and methylmagnesium chloride (0.3 mL, 3.0 M) was added dropwise with the temperature maintained at -20 °C. The reaction system was stirred at this temperature for 4 h. Then the reaction system was added with a small amount of water to quench the reaction, and concentrated. The residue was separated by preparative chromatography to give **compound 119** (10.8 mg, 7.1% yield) in the form of a white solid. LC-MS [M+H]⁺: 527.0.

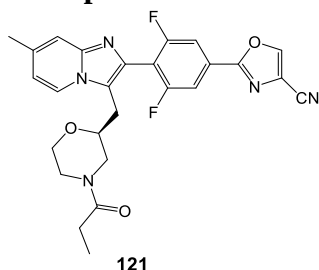
¹H NMR (400 MHz, CDCl₃) δ 8.43 (d, J = 7.2 Hz, 1H), 7.89 (s, 1H), 7.80 (d, J = 7.9 Hz, 1H), 7.36 (s, 1H), 6.89 (dd, J = 7.1, 1.5 Hz, 1H), 3.89 – 3.74 (m, 3H), 3.70 (s, 3H), 3.64 – 3.57 (m, 1H), 3.41 – 3.33 (m, 1H), 3.16 – 3.09 (m, 2H), 2.96 – 2.80 (m, 1H), 2.70 – 2.55 (m, 1H), 2.49 (s, 3H), 1.61 (s, 6H).

Example 120

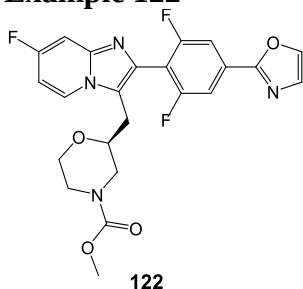


The compound was prepared according to the preparation method as described in Example 10. LC-MS: [M+H]⁺ = 494.3.

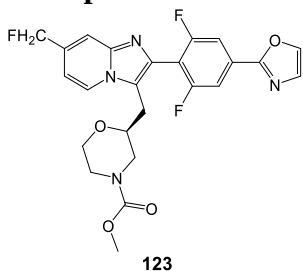
Example 121



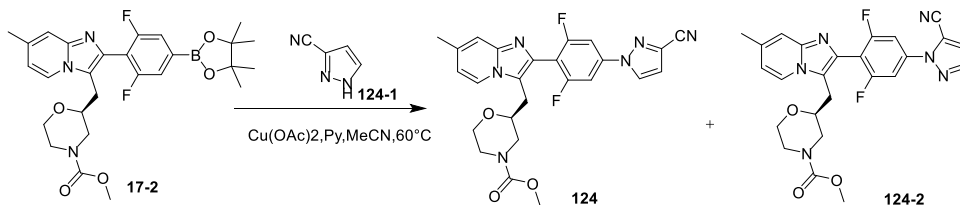
The compound was prepared according to the preparation method as described in Example 10. LC-MS: [M+H]⁺ = 492.4.

Example 122

The compound was prepared according to the preparation method as described in Example 10. LC-MS: $[M+H]^+ = 473.5$.

Example 123

The compound was prepared according to the preparation method as described in Example 10. LC-MS: $[M+H]^+ = 487.1$.

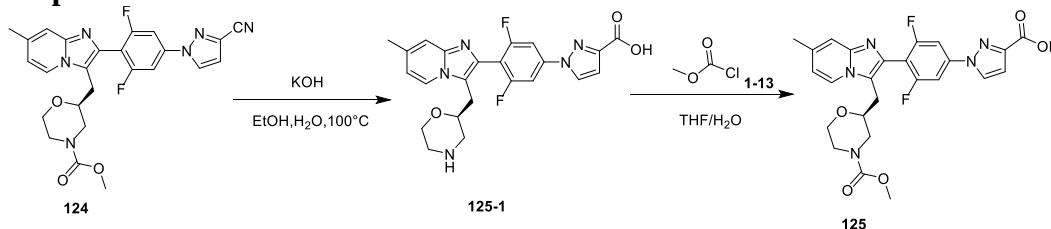
Example 124

Intermediate 17-2 (190 mg, 0.36 mmol), *1H*-pyrazole-3-carbonitrile (40 mg, 0.43 mmol), anhydrous copper(II) acetate (98 mg, 0.54 mmol) and pyridine (57 mg, 0.72 mmol) were dissolved in acetonitrile (3 mL). The reaction system was heated to 60 °C and reacted overnight. Then the reaction system was cooled to room temperature, diluted with water (20 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 124** (70 mg, white solid) and **compound 124-2** (5 mg, 42% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 493.3$.

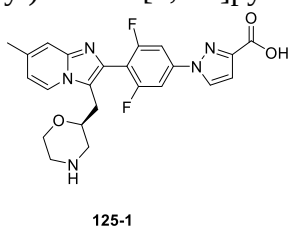
Compound 124: ^1H NMR (400 MHz, CDCl_3) δ 8.33 (d, $J = 6.0$ Hz, 1H), 8.08 (d, $J = 2.4$ Hz, 1H), 7.70 (brs, 1H), 7.52 (d, $J = 8.0$ Hz, 2H), 6.95 (d, $J = 2.5$ Hz, 1H), 6.88 (d, $J = 6.0$ Hz, 1H), 4.05-3.76 (m, 3H), 3.71 (s, 3H), 3.66-3.58 (m, 1H), 3.46-3.36 (m, 1H), 3.10-2.83 (m, 3H), 2.70-2.62 (m, 1H), 2.51 (s, 3H).

Compound 124-2: ^1H NMR (400 MHz, CDCl_3) δ 8.29 (brs, 1H), 7.88 (s, 1H), 7.70-7.50 (m, 3H), 7.14 (s, 1H), 6.83 (brs, 1H), 4.08-3.75 (m, 3H), 3.70 (s, 3H), 3.66-3.58 (m, 1H), 3.46-3.36 (m, 1H), 3.10-2.85 (m, 3H), 2.72-2.62 (m, 1H), 2.49 (s, 3H).

Example 125

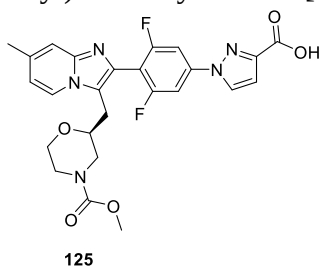


Step (1) Preparation of (S)-3,5-difluoro-4-(7-methyl-3-(morpholinyl-2-methyl)imidazo[1,2-a]pyridin-2-yl)-1H-pyrazole-3-carboxylic acid



Compound 124 (40 mg, 0.08 mmol) was dissolved in ethanol (2 mL) and water (1 mL), and potassium hydroxide (13 mg, 0.24 mmol) was added. The reaction system was reacted at 100 °C for 12 h under nitrogen atmosphere. The the reaction system was cooled to room temperature, and concentrated to give **intermediate 125-1** (40 mg, crude product), which was used directly in the next step. LC-MS: $[\text{M}+\text{H}]^+ = 454.2$.

Step (2) Preparation of (S)-1-(3,5-difluoro-4-(3-((4-(methoxycarbonyl)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)-1H-pyrazole-3-carboxylic acid

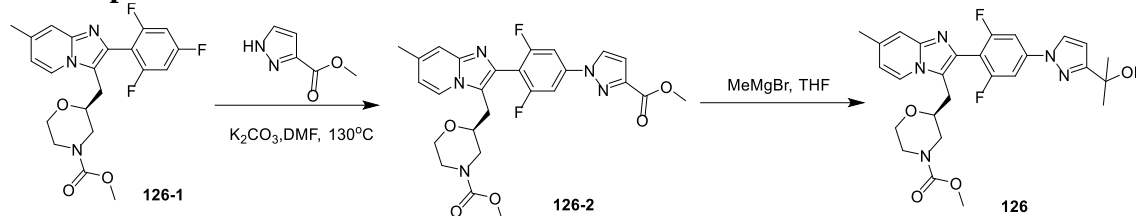


The **intermediate 125-1** (40 mg, 0.08 mmol, crude product) was dissolved in tetrahydrofuran/water (2 mL, 1/1), and methyl chloroformate (10 mg, 0.108 mmol) was added dropwise in an ice water bath. The reaction system was reacted at this temperature for 1 h. Then the reaction system was adjusted to pH 4 with diluted hydrochloric acid, and concentrated. The residual aqueous phase was lyophilized. The crude product was purified by preparative high performance liquid chromatography to give **compound 125** (7.6 mg, 18% yield), LC-MS: $[\text{M}+\text{H}]^+ = 512.3$.

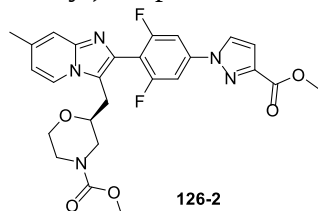
^1H NMR (400 MHz, CDCl_3) δ 8.28 (d, J = 7.0 Hz, 1H), 7.98 (d, J = 2.1 Hz, 1H), 7.70 (s, 1H), 7.49 (d, J = 8.2 Hz, 2H), 7.07 (d, J = 2.4 Hz, 1H), 6.81 (d, J = 7.0 Hz, 1H), 4.06 – 3.78 (m, 3H), 3.70 (s, 3H), 3.66 – 3.58 (m, 1H), 3.46 – 3.35 (m, 1H), 3.05 – 2.85 (m, 3H), 2.70 – 2.62 (m, 1H), 2.49 (s, 3H).

5

Example 126



Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(3-(carbomethoxy<methoxycarbonyl>)-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

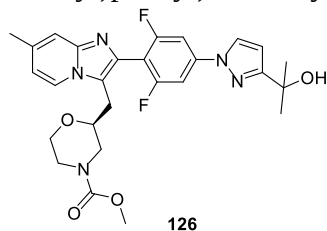


10

Methyl (S)-2-((7-methyl-2-(2,4,6-trifluorophenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate (700 mg, 1.67 mmol), methyl 3-pyrazolecarboxylate (210 mg, 1.67 mmol) and anhydrous potassium carbonate (692 mg, 5.00 mmol) were dissolved in *N,N*-dimethylformamide (15 mL) under nitrogen atmosphere. The reaction system was stirred at 130°C for 4 h. Then the reaction system was cooled, added with ice water (15 mL) to quench the reaction, and extracted with ethyl acetate (10 mL $\times$ 3). The organic phases were combined, washed with saturated brine (10 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 126-2** (300 mg, 80% purity, 27% yield) in the form of a yellow solid. MS $[\text{M}+\text{H}]^+ = 526.2$.

20

Step (2) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(3-(2-hydroxypropan-2-yl)-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



126

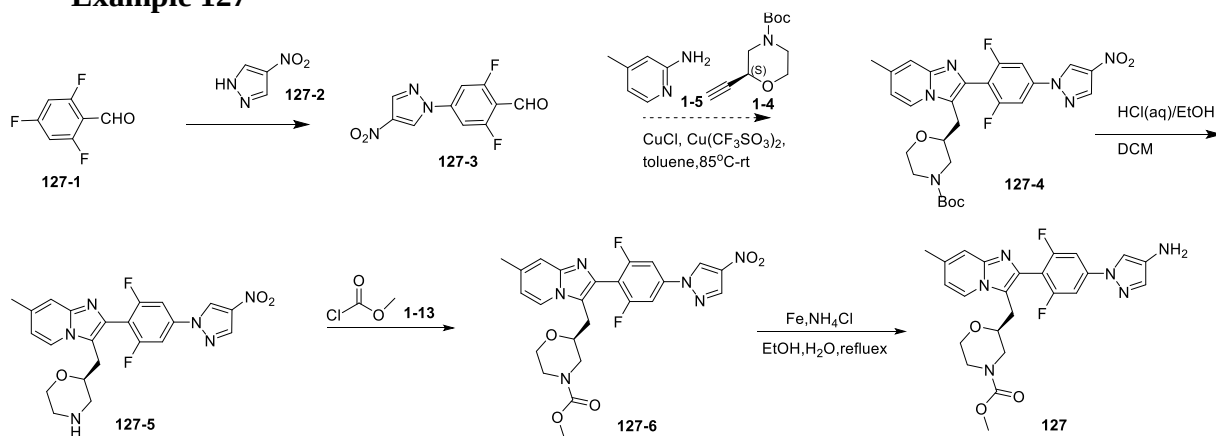
Intermediate 126-2 (150 mg, 0.28 mmol) was dissolved in tetrahydrofuran (5 mL) under nitrogen atmosphere. The reaction system was cooled to 0°C in an ice water bath, added

25

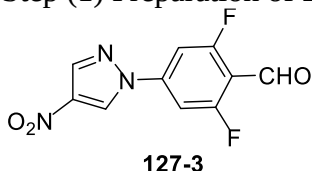
dropwise with methyl magnesium bromide solution (3 M, 0.3 mL, 0.9 mmol) with the temperature maintained at 0 °C, and stirred for 1 h. Then the reaction system was added with saturated ammonium chloride solution (10 mL) at 0 °C to quench the reaction, and extracted with ethyl acetate (10 mL × 3). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 126** (30 mg, 17% yield) in the form of a white solid. MS [M+H]⁺ = 525.8.

¹H NMR (400 MHz, MEOD): δ 8.45 (d, J = 7.0 Hz, 1H), 8.31 (d, J = 2.6 Hz, 1H), 7.67 (d, J = 8.9 Hz, 2H), 7.39 (s, 1H), 6.93 (d, J = 7.1 Hz, 1H), 6.60 (d, J = 2.6 Hz, 1H), 3.89 - 3.76 (m, 3H), 3.66 (s, 3H), 3.64 - 3.60 (m, 1H), 3.42 - 3.35 (m, 1H), 3.16 - 3.11 (m, 2H), 3.00 - 2.85 (m, 1H), 2.71 - 2.60 (m, 1H), 2.49 (s, 3H), 1.64 (s, 6H).

Example 127

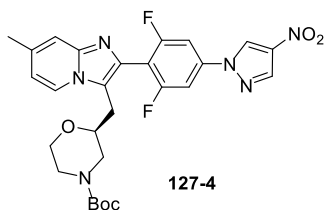


Step (1) Preparation of 2,6-difluoro-4-(4-nitro-1H-pyrazol-1-yl)benzaldehyde



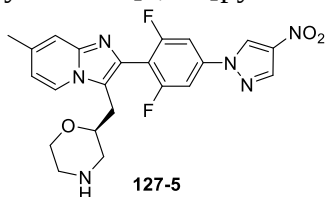
2,4,6-trifluorobenzaldehyde (2.0 g, 12.5 mmol), 4-nitro-1H-pyrazole (1.4 g, 12.5 mmol) and triethylamine (3.8 g, 37.5 mmol) were dissolved in acetonitrile (20 mL) under nitrogen atmosphere. The reaction system was heated to reflux for 3 h. Then the reaction system was cooled to room temperature, added with water (50 mL) to quench the reaction, and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 127-3** (450 mg, 14% yield) in the form of a yellow solid. LC-MS [M+H]⁺ = 254.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(4-nitro-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



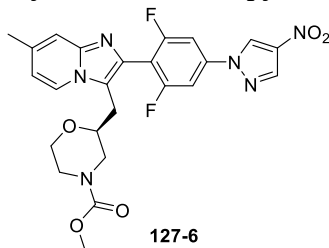
Intermediate 127-3 (450 mg, 1.78 mmol), 4-methylpyridin-2-amine (192 mg, 1.78 mmol), *tert*-butyl (*S*)-2-ethynylmorpholine-4-carboxylate (375 mg, 1.78 mmol), copper(I) chloride (53 mg, 0.53 mmol) and copper (II) trifluoromethanesulfonate (193 mg, 0.53 mmol) were dissolved in toluene (5 mL) under nitrogen atmosphere. The reaction system was heated to 85 °C and stirred for 20 min, added with *N,N*-dimethylacetamide (0.15 mL), and reacted at 85 °C for 5 h. Then the reaction system was cooled to room temperature and reacted overnight. Then the reaction system was added with water (30 mL) to quench the reaction, and extracted with dichloromethane (15 mL × 3). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 127-4** (260 mg, 26% yield) in the form of a brown oily liquid. LC-MS $[M+H]^+ = 555.2$.

Step (3) Preparation of (*S*)-2-((2-(2,6-difluoro-4-(4-nitro-1*H*-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



Intermediate 127-4 (260 mg, 0.5 mmol) was dissolved in dichloromethane (3 mL). The reaction system was cooled to 0 °C in an ice water bath, added dropwise with a solution of HCl in ethanol (33%, 1 mL), and stirred at room temperature for 1 h. Then the reaction system was concentrated by rotary evaporation to give **intermediate 127-5** (280 mg, crude product), which was used directly in the next step. LC-MS $[M+H]^+ = 455.2$.

Step (4) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(4-nitro-1*H*-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

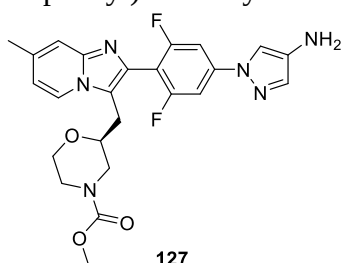


Intermediate 127-5 (200 mg, 0.44 mmol) and *N,N*-diisopropylethylamine (284 mg, 2.20 mmol) were dissolved in dichloromethane (3 mL) under nitrogen atmosphere. The reaction

system was added dropwise with methyl chloroformate (50 mg, 0.52 mmol) at 0 °C in an ice water bath. After the completion of the dropwise addition, the reaction system was reacted for 1 h. Then the reaction system was added with water (20 mL) to quench the reaction, and extracted with dichloromethane (15 mL × 3). The organic phases were combined, washed with saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by silica gel column chromatography to give **intermediate 127-6** (110 mg, 48% yield) in the form of a brown solid. LC-MS $[M+H]^+ = 513.3$.

^1H NMR (400 MHz, CDCl_3) δ 8.71 (s, 1H), 8.34 (s, 1H), 8.25 (d, $J = 7.1$ Hz, 1H), 7.52 – 7.48 (m, 3H), 6.78 (s, 1H), 4.00 – 3.75 (m, 3H), 3.70 (s, 3H), 3.67 – 3.56 (m, 1H), 3.45 – 3.35 (m, 1H), 3.10 – 2.85 (m, 3H), 2.68 – 2.60 (m, 1H), 2.47 (s, 3H).

Step (5) Preparation of methyl (S)-2-((2-(4-(4-amino-1*H*-pyrazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

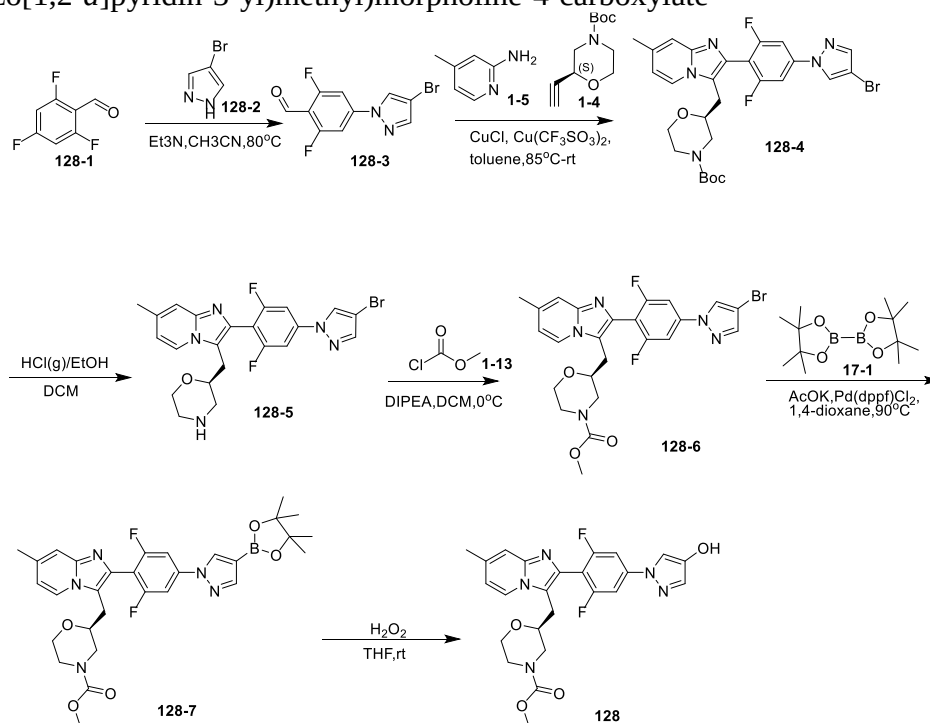


Intermediate 127-6 (80 mg, 0.15 mmol) and saturated ammonium chloride (0.5 mL) were dissolved in ethanol (2 mL) under nitrogen atmosphere. The reaction system was heated to 80 °C, added with iron powder (44 mg, 0.75 mmol), and stirred for 15 min. Then the reaction system was cooled to room temperature, filtered and concentrated. The crude product was purified by preparative high performance liquid chromatography to give **compound 127** (9 mg, 12% yield) in the form of a white solid. LC-MS $[M+H]^+ = 483.2$.

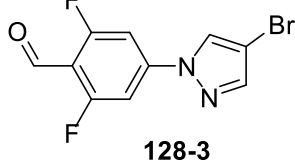
^1H NMR (400 MHz, CDCl_3) δ 8.25 (d, $J = 7.1$ Hz, 1H), 7.53 (s, 1H), 7.52 (s, 1H), 7.46 (s, 1H), 7.34 (d, $J = 8.7$ Hz, 2H), 6.76 (d, $J = 7.0$ Hz, 1H), 4.00 – 3.75 (m, 3H), 3.69 (s, 3H), 3.66 – 3.56 (m, 1H), 3.45–3.35 (m, 1H), 3.08 – 2.85 (m, 3H), 2.67 – 2.61 (m, 1H), 2.46 (s, 3H).

Example 128

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-hydroxy-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

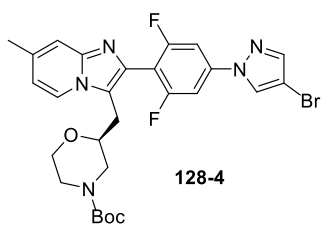


5 Step (1) Preparation of 4-(4-bromo-1H-pyrazol-1-yl)-2,6-difluorobenzaldehyde



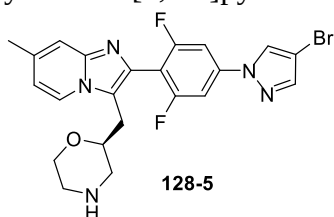
2,4,6-trifluorobenzaldehyde (2.0 g, 12.49 mmol) was dissolved in anhydrous acetonitrile (20 mL), and 4-bromopyrazole (1.8 g, 12.49 mmol) and triethylamine (3.8 g, 37.47 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 80 °C for 1 h. Then the reaction system was cooled to room temperature, added with water (60 mL), and extracted with ethyl acetate (80 mL × 3). The organic phases were combined, washed with saturated brine (50 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 128-3** (800 mg, 22.2% yield) in the form of a white solid. MS [M+H]⁺ = 287.7.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(4-(4-bromo-1H-pyrazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



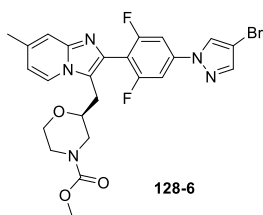
Intermediate 128-3 (600 mg, 2.09 mmol), 4-methylpyridin-2-amine (226 mg, 2.09 mmol), *tert*-butyl (*S*)-2-ethynylmorpholine-4-carboxylate (442 mg, 2.09 mmol), copper(I) chloride (62 mg, 0.63 mmol) and copper(II) trifluoromethanesulfonate (227 mg, 0.63 mmol) were dissolved in toluene solution (6 mL) under nitrogen atmosphere. The reaction system was reacted at 85 °C for 10 min, added with *N,N*-dimethylacetamide (0.2 mL), and reacted at 85 °C for another 5 h. Then the reaction system was reacted at room temperature overnight. After the reaction was completed, the reaction system was filtered, and the filtrate was added with water (30 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (40 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 128-4** (200 mg, 17% yield) in the form of a yellow oil. MS [M + H]⁺ = 588.8.

Step (3) Preparation of (*S*)-2-((2-(4-(4-bromo-1*H*-pyrazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



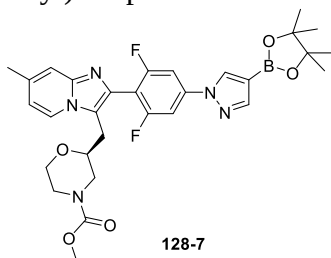
Intermediate 128-4 (200 mg, 0.34 mmol) was dissolved in dichloromethane (2 mL), and a solution of HCl in ethanol (33%, 0.5 mL) was added dropwise with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. Then the reaction system was added dropwise with saturated sodium bicarbonate solution to adjust the pH to 8 with the temperature maintained at 0 °C, added with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give **intermediate 128-5** (180 mg, crude product) in the form of a yellow oil. MS [M + H]⁺ = 488.7.

Step (4) Preparation of methyl (*S*)-2-((2-(4-(4-bromo-1*H*-pyrazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



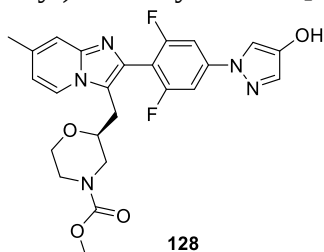
Intermediate 128-5 (180 mg, 0.34 mmol) and *N,N*-diisopropylethylamine (143 mg, 1.11 mmol) were dissolved in dichloromethane (2 mL), and methyl chloroformate (42 mg, 0.44 mmol) was added dropwise under nitrogen atmosphere with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 1 h. Then the reaction system was added with water (10 mL), and extracted with dichloromethane (15 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 128-6** (150 mg, 81% yield) in the form of a yellow solid. MS $[M + H]^+ = 546.7$.

Step (5) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(4-(4,4,5,5-tetramethyl-1,3,2-dioxaborolan-2-yl)-1*H*-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 128-6 (150 mg, 0.27 mmol) was dissolved in anhydrous 1,4-dioxane (2 mL) under nitrogen atmosphere, and 4,4,4',4',5,5,5',5'-octamethyl-2,2'-bis(1,3,2-dioxaborolan) (87 mg, 0.34 mmol), potassium acetate (54 mg, 0.55 mmol) and [1,1'-bis(diphenylphosphino)ferrocene]dichloropalladium (10 mg, 0.014 mmol) were added. The reaction system was reacted at 90 °C for 2 h. After the reaction was completed, the reaction system was cooled to room temperature, filtered and concentrated by rotary evaporation to give **intermediate 128-7** (180 mg, crude product) in a yellow oil, MS $[M + H]^+ = 593.7$.

Step (6) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(4-hydroxy-1*H*-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

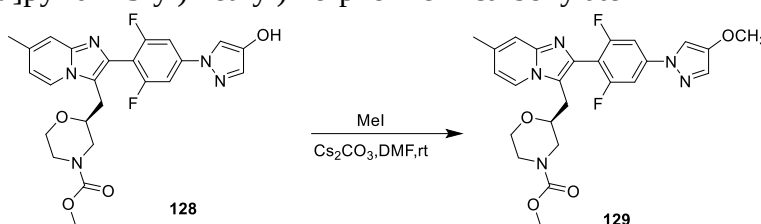


Intermediate 128-7 (180 mg, 0.27 mmol) was dissolved in tetrahydrofuran (2 mL), and hydrogen peroxide (0.5 mL) was added dropwise. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added dropwise with saturated sodium thiosulfate solution (2 mL) to quench the reaction, added with water (10 mL), and extracted with dichloromethane (15 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by high performance liquid chromatography to give **compound 128** (22.9 mg, 17.3% yield) in the form of a white solid. MS [M + H]⁺ = 483.8.

¹H NMR (400 MHz, MeOD) δ 8.86 (d, J = 7.1 Hz, 1H), 7.93 (s, 1H), 7.72 (s, 1H), 7.70 (d, J = 9.6 Hz, 2H), 7.50 (s, 1H), 7.42 (dd, J = 7.1, 1.5 Hz, 1H), 4.03 - 4.00 (m, 1H), 3.85 - 3.76 (m, 3H), 3.69 (s, 3H), 3.40 - 3.36 (m, 1H), 3.28 - 3.26 (m, 2H), 2.99 - 2.90 (m, 1H), 2.80 - 2.72 (m, 1H), 2.66 (s, 3H).

Example 129

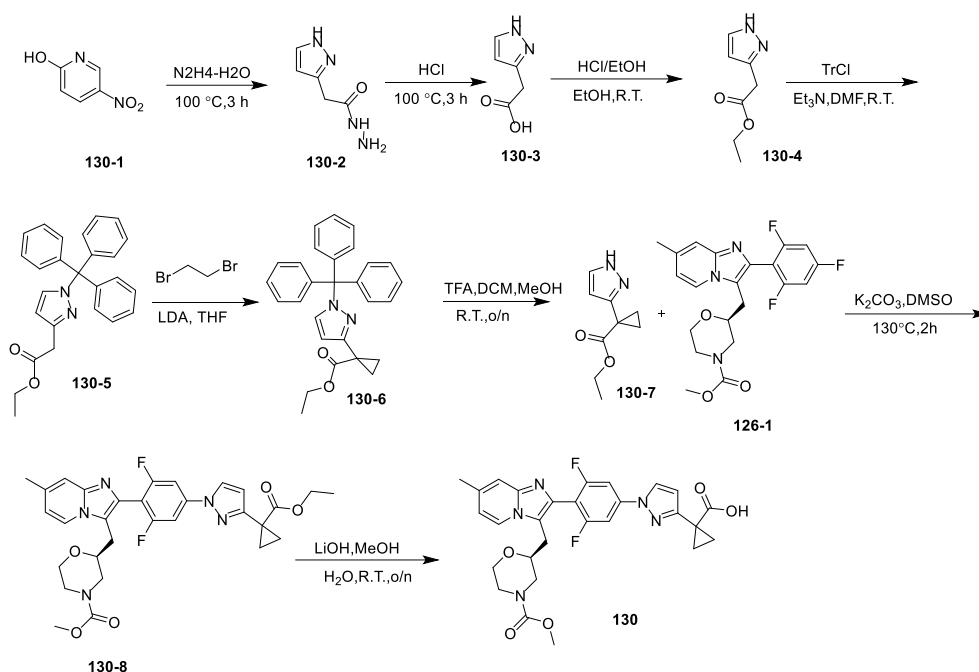
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4-methoxy-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



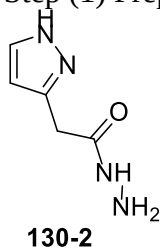
Compound 128 (45 mg, 0.09 mmol) was dissolved in *N,N*-dimethylformamide (1 mL) under nitrogen atmosphere, and cesium carbonate (36 mg, 0.11 mmol) and iodomethane (20 mg, 0.14 mmol) were added sequentially. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added with water (10 mL) to quench the reaction, and extracted with ethyl acetate (15 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by high performance liquid chromatography to give **compound 129** (5.3 mg, 11.4% yield) in the form of a white solid. MS [M + H]⁺ = 497.8.

¹H NMR (400 MHz, MeOD) δ 8.42 (d, J = 7.0 Hz, 1H), 8.15 (s, 1H), 7.59 (d, J = 9.0 Hz, 2H), 7.56 (s, 1H), 7.35 (s, 1H), 6.88 (dd, J = 7.2 Hz, 1.2 Hz, 1H), 3.86 (s, 3H), 3.86 - 3.76 (m, 3H), 3.66 (s, 3H), 3.63 - 3.58 (m, 1H), 3.42 - 3.36 (m, 1H), 3.12 - 3.11 (m, 2H), 2.93 - 2.88 (m, 1H), 2.68 - 2.61 (m, 1H), 2.47 (s, 3H).

Example 130

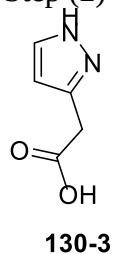


Step (1) Preparation of 2-(1H-pyrazol-3-yl)acethydrazide



- 5 5-nitropyridin-2-ol (10.0 g, 71.38 mmol) was dissolved in hydrazine hydrate (50 mL). The reaction system was stirred at 100°C for 3 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give **intermediate 130-2** (10.0 g, crude product) in the form of a red oil. LC-MS $[\text{M}+\text{H}] = 140.7$.

Step (2) Preparation of 2-(1H-pyrazol-3-yl)acetic acid

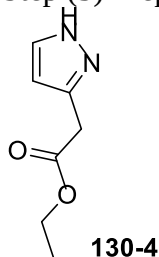


10

Intermediate 130-2 (10 g) was added to concentrated hydrochloric acid (100 mL) under nitrogen atmosphere. The reaction system was heated to 100°C and stirred for 3 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give

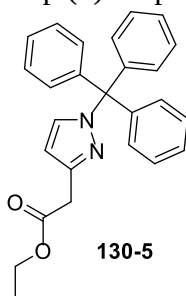
intermediate 130-3 (10 g, crude product) in the form of a colorless transparent oil, MS $[M+H]^+$ = 127.0.

Step (3) Preparation of ethyl 2-(1*H*-pyrazol-3-yl)acetate



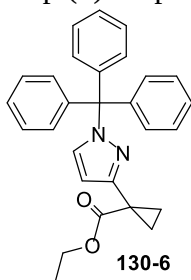
5 **Intermediate 130-3** (10.0 g, crude product) was dissolved in ethanol (15 mL), and a solution of HCl in ethanol (33%, 12.5 mL) was added. The reaction system was stirred at room temperature for 3 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give **intermediate 130-4** (12 g, crude product) in the form of a brown oil, MS $[M+H]^+$ = 155.0.

10 Step (4) Preparation of ethyl 2-(1-trityl-1*H*-pyrazol-3-yl)acetate



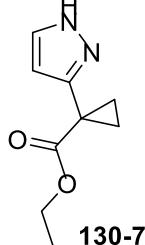
15 **Intermediate 130-4** (12.0 g, crude product) was dissolved in anhydrous DMF (100 mL), and triphenylmethyl chloride (20.0 g, 77.42 mmol) and triethylamine (28.8 g, 285.5 mmol) were added. The reaction system was stirred at room temperature overnight. After the reaction was completed, the reaction system was added with water (300 mL), and extracted with ethyl acetate (100 mL $\times$ 3). The organic phases were combined, washed with saturated brine (150 mL $\times$ 3), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 130-5** (2.3 g, 7.5% yield) in the form of a white solid. MS $[M+H]^+$ = 418.8.

20 Step (5) Preparation of ethyl 1-(1-trityl-1*H*-pyrazol-3-yl)cyclopropane-1-carboxylate



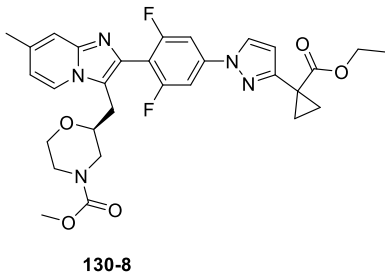
Intermediate 130-5 (1.0 g, 2.52 mmol) was dissolved in anhydrous tetrahydrofuran (20 mL). The reaction system was cooled to -78 °C, added dropwise with lithium diisopropylamide (2.4 mL, 2 N, 4.80 mmol), and stirred at -78 °C for 20 min. Then the reaction system was added with dibromoethane (2.8 g, 15.13 mmol), slowly warmed to room temperature, and stirred overnight. After the reaction was completed, the reaction system was added with saturated ammonium chloride (10 mL), diluted with water (20 mL), and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered, and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 130-6** (550 mg, 51.6% yield) in the form of a yellow oil. MS [M+H]⁺ = 444.8.

Step (6) Preparation of ethyl 1-(1*H*-pyrazol-3-yl)cyclopropane-1-carboxylate



Intermediate 130-6 (400 mg, 1.01 mmol) was dissolved in dichloromethane (2 mL) and methanol (2 mL), and trifluoroacetic acid (1.2 mL) was added. The reaction system was stirred at room temperature overnight. After the reaction was completed, the reaction system was adjusted to pH 10 with saturated sodium carbonate solution, and extracted with dichloromethane (15 mL × 3). The organic phases were combined, dried over sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 130-7** (65 mg, 35.6% yield) in the form of a white solid, MS [M+H]⁺ = 181.0.

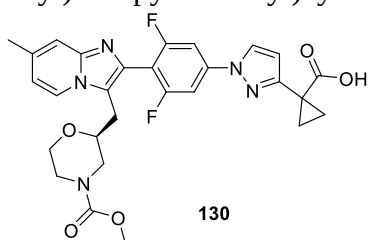
Step (7) Preparation of methyl (S)-2-((2-(4-(3-(1-(carbethoxyethoxycarbonyl)cyclopropyl)-1*H*-pyrazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 130-7 (35 mg, 0.19 mmol) was dissolved in anhydrous DMSO (1.5 mL), and **intermediate 126-1** (87 mg, 0.21 mmol) and potassium carbonate (66 mg, 0.48 mmol) were

added. The reaction system was stirred at 130 °C overnight. After the reaction was completed, the reaction system was added with water (15 mL), and extracted with ethyl acetate (10 mL × 3). The organic phases were combined, washed with saturated brine (10 mL), dried over sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 130-8** (25 mg, 22.7% yield) in the form of a white solid. MS $[M + H]^+ = 579.8$.

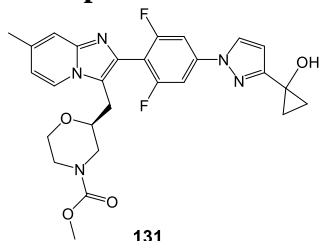
Step (8) Preparation of (S)-1-(1-(3,5-difluoro-4-(3-((4-(carbomethoxy<methoxycarbonyl>)morpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)phenyl)-1*H*-pyrazol-3-yl)cyclopropane-1-carboxylic acid



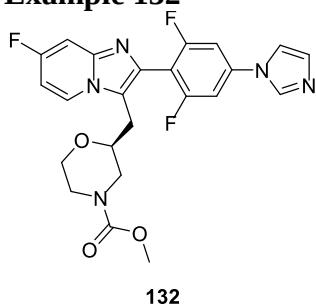
Intermediate 130-8 (25 mg, 0.044 mmol) was dissolved in tetrahydrofuran (1 mL)/methanol (1 mL), and lithium hydroxide solution (0.5 mL, 1 N) was added. The reaction system was stirred at room temperature overnight. After the reaction was completed, the reaction system was adjusted to pH 4 with diluted hydrochloric acid, and extracted with ethyl acetate (5 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was separated by high performance liquid chromatography, and lyophilized to give **compound 130** (5.0 mg, 20.6% yield) in the form of a white solid. MS $[M + H]^+ = 551.7$.

^1H NMR (400 MHz, CDCl_3) δ 8.31 (d, $J = 6.6$ Hz, 1H), 7.93 (d, $J = 2.6$ Hz, 1H), 7.63 (s, 1H), 7.38 (d, $J = 8.2$ Hz, 2H), 6.84 ((d, $J = 6.8$ Hz, 1H), 6.33 (s, 1H), 3.05 – 3.77 (m, 3H), 3.70 (s, 3H), 3.66 – 2.59 (m, 1H), 3.46 – 3.37 (m, 1H), 3.10 – 2.99 (m, 2H), 2.99 – 2.87 (m, 1H), 2.70 – 2.63 (m, 1H), 2.50 (s, 3H), 1.94 (dd, $J = 7.6, 4.0$ Hz, 2H), 1.47 (dd, $J = 7.6, 4.1$ Hz, 2H)

Example 131



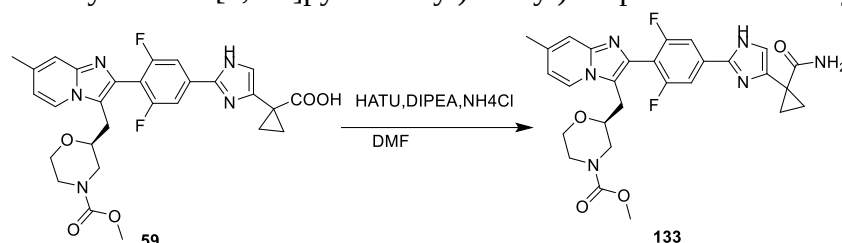
The compound was prepared according to the preparation method as described in Example 126. LC-MS: $[M + H]^+ = 524.2$.

Example 132

The compound was prepared according to the preparation method as described in Example 5. LC-MS: $[M+H]^+ = 472.2$.

5 **Example 133**

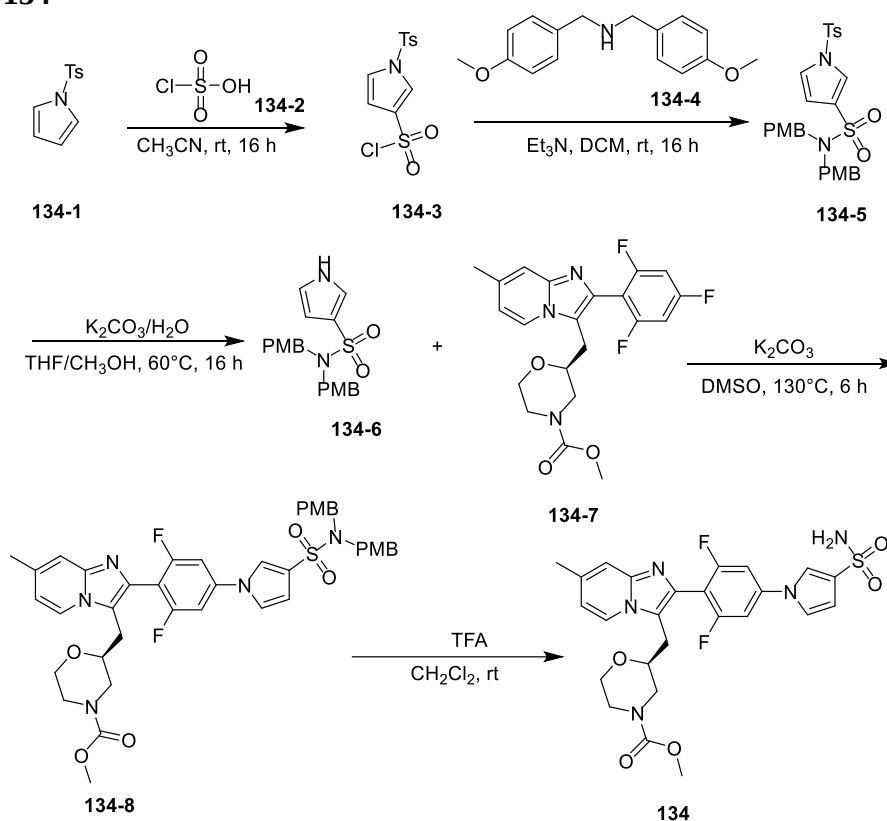
Preparation of methyl (S)-2-((2-(4-(4-(1-carbamylcyclopropyl)-1H-imidazol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



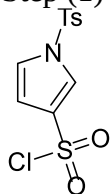
Compound 59 (400 mg, 0.72 mmol) and 2-(7-azabenzotriazol-1-yl)-*N,N,N',N'*-tetramethyluronium hexafluorophosphate (410 mg, 1.08 mmol) were dissolved in *N,N*-dimethylformamide (5 mL). The reaction system was stirred at room temperature for 10 min under nitrogen atmosphere, added with *N,N*-diisopropylethylamine (279 mL, 2.16 mmol) and ammonium chloride (58 mg, 1.1 mmol), and stirred at room temperature for another 2 h. Then the reaction system was added with water (30 mL) to quench the reaction, and extracted with ethyl acetate (15 mL $\times$ 3). The organic phases were combined, washed with saturated brine (10 mL $\times$ 3), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated by silica gel column chromatography to give a crude product (200 mg) in the form of a brown solid. The crude product (50 mg) was purified by preparative chromatography to give **compound 133** (11.1 mg) in the form of a white solid. LC-MS $[M+H]^+$: 551.

¹H NMR (400 MHz, CDCl₃) δ 8.27 (d, *J* = 6.9 Hz, 1H), 7.47 (s, 1H), 7.39 (d, *J* = 8.3 Hz, 2H), 6.96 (s, 1H), 6.80 (d, *J* = 6.9 Hz, 1H), 5.56 (brs, 1H), 3.90 – 3.80 (m, 3H), 3.66 (s, 3H), 3.62 – 3.54 (m, 1H), 3.43 – 3.34 (m, 1H), 3.04 – 2.82 (m, 3H), 2.67 – 2.56 (m, 1H), 2.51 (s, 3H), 1.65 – 1.59 (m, 2H), 1.15 – 1.09 (m, 2H).

Example 134



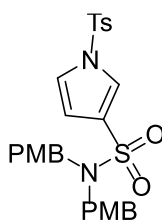
Step (1) Preparation of 1-tosyl-1*H*-pyrrol-3-sulfonyl chloride



134-3

- 5 *N*-*p*-toluenesulfonylpyrrole (2.00 g, 9.01 mmol) was dissolved in acetonitrile (20 mL), and chlorosulfonic acid (5 mL) was added dropwise slowly at 0°C in an ice water bath. The reaction system was reacted at room temperature overnight. Then the reaction system was added dropwise slowly to an ice (100 g), and extracted with dichloromethane ($80\text{ mL} \times 2$). The organic phases were combined, washed with saturated sodium bicarbonate ($30\text{ mL} \times 2$), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 134-3** (600 mg, 21% yield) in the form of a white solid.
- 10

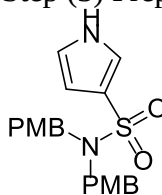
Step (2) Preparation of *N*,*N*-bis(4-methoxybenzyl)-1-tosyl-1*H*-pyrrol-3-sulfonamide



134-5

Bis-(4-methoxybenzyl)-amine (195 mg, 0.76 mmol) and triethylamine (191 mg, 1.89 mmol) were dissolved in dichloromethane (5 mL) under nitrogen atmosphere, and a solution of **intermediate 134-3** (200 mg, 0.63 mmol) in dichloromethane (1 mL) was added dropwise at 0 °C in an ice water bath. The reaction system was stirred at room temperature overnight. Then the reaction system was poured into water (20 mL), and extracted with dichloromethane (30 mL × 2). The organic phases were combined, washed with saturated brine (30 mL), filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 134-5** (300 mg, 88% yield) in the form of a white solid. MS $[M+Na]^+ = 562$.

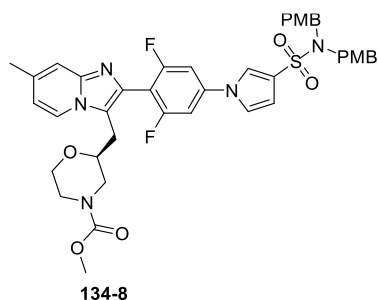
Step (3) Preparation of *N,N*-bis(4-methoxybenzyl)-1*H*-pyrrol-3-sulfonamide



134-6

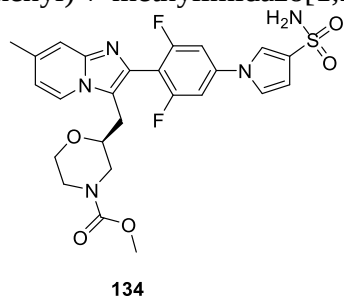
Intermediate 134-5 (800 mg, 1.48 mmol) was dissolved in methanol (10 mL) and tetrahydrofuran (10 mL), and aqueous potassium carbonate solution (3 mL, 7.40 mmol) was added. The reaction system was stirred at 60 °C overnight. Then the reaction system was concentrated under reduced pressure. The residue was adjusted to pH 4 with acetic acid, diluted with ethyl acetate (50 mL), washed with saturated brine (30 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 134-6** (500 mg, 88% yield) in the form of a white solid. MS $[M + Na]^+ = 408.8$.

Step (4) Preparation of methyl (*S*)-2-((2-(4-(3-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-1*H*-pyrrol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Methyl (S)-2-((7-methyl-2-(2,4,6-trifluorophenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate (271 mg, 0.65 mmol, **intermediate 134-7**, prepared according to the preparation method for **intermediate 2-7**, LC-MS: $[M+H]^+ = 420.2$) was dissolved in dimethyl sulfoxide (5 mL), and potassium carbonate (269 mg, 1.95 mmol) and *N,N*-bis(4-methoxybenzyl)-1*H*-pyrrol-3-sulfonamide (251 mg, 0.65 mmol) were added. The reaction system was reacted at 130 °C for 6 h under nitrogen atmosphere. Then the reaction system was added with water (30 mL), and extracted with ethyl acetate (30 mL × 2). The organic phases were combined, washed with saturated brine (20 mL × 2), filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 134-8** (320 mg, 44% yield) in the form of a brown solid. MS $[M+H]^+ = 786.2$.

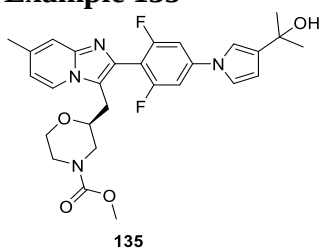
Step (5) Preparation of methyl (S) 2-((2-(2,6-difluoro-4-(3-sulfamoyl-1*H*-pyrrol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 134-8 (280 mg, 0.36 mmol) was dissolved in dichloromethane (3 mL), and trifluoroacetic acid (3 mL) was added. The reaction system was reacted at room temperature for 16 h. Then the reaction system was concentrated under reduced pressure. The crude product was dissolved in dichloromethane (40 mL), washed with aqueous sodium bicarbonate solution (10%, 10 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative chromatography to give **compound 134** (22.6 mg, 12% yield) in the form of a white solid. MS $[M+H]^+ = 545.7$.

¹H NMR (400 MHz, CDCl₃) δ 8.23 (d, *J* = 7.0 Hz, 1H), 7.73-7.63 (m, 1H), 7.45 (s, 1H), 7.16 – 7.13 (m, 1H), 7.11 (d, *J* = 7.7 Hz, 2H), 6.80 – 6.69 (m, 2H), 4.96 (s, 2H), 4.00 – 3.82 (m, 3H), 3.70 (s, 3H), 3.64 – 3.58 (m, 1H), 3.45 – 3.38 (m, 1H), 3.12 – 2.99 (m, 2H), 2.98 – 2.88 (m, 1H), 2.69 – 2.59 (m, 1H), 2.46 (s, 3H).

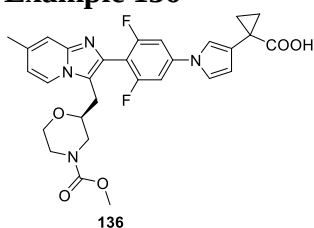
Example 135



The compound was prepared according to the preparation method as described in Example 126. LC-MS: $[M+H]^+ = 525.2$.

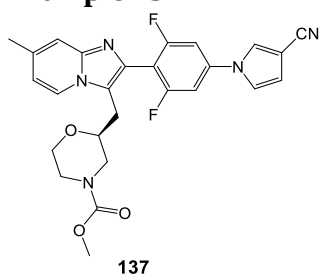
5

Example 136



The compound was prepared according to the preparation method for compound 5. LC-MS: $[M+H]^+ = 551.2$.

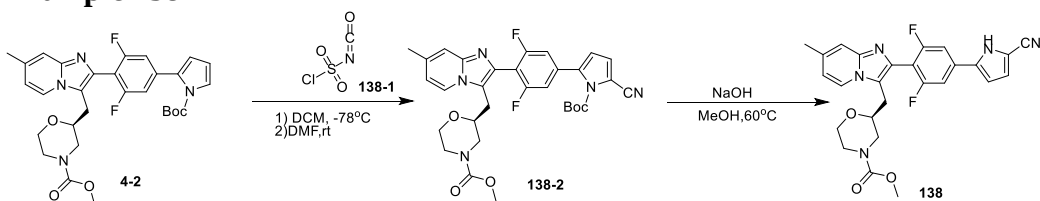
Example 137



10

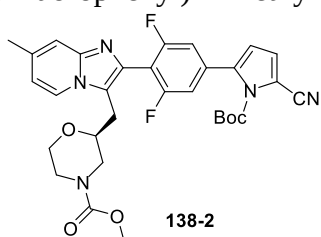
The compound was prepared according to the preparation method for compound 5. LC-MS: $[M+H]^+ = 492.7$.

Example 138



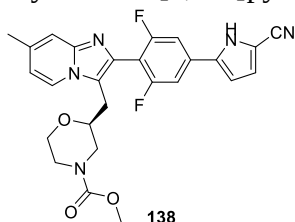
15

Step (1) Preparation of methyl (S)-2-((2-(4-(1-(tert-butoxycarbonyl)-5-cyano-pyrrol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate



Intermediate 4-2 (290 mg, 0.51 mmol) was dissolved in anhydrous dichloromethane (5 mL) under nitrogen atmosphere, and chlorosulfonyl isocyanate (58 mg, 0.41 mmol) was added dropwise slowly at -78 °C. After the completion of the dropwise addition over 2 min, the reaction system was reacted at -78 °C for 1 h, added dropwise with *N,N*-dimethylformamide (0.44 mL) over 1 min, and reacted at room temperature for 1 h. Then the reaction system was added with water (20 mL), and extracted with dichloromethane (50 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 138-2** (200 mg, 66% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 592.8$.

Step (2) Preparation of methyl (*S*)-2-((2-(2,6-difluoro-4-(5-cyano-1*H*-pyrrol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

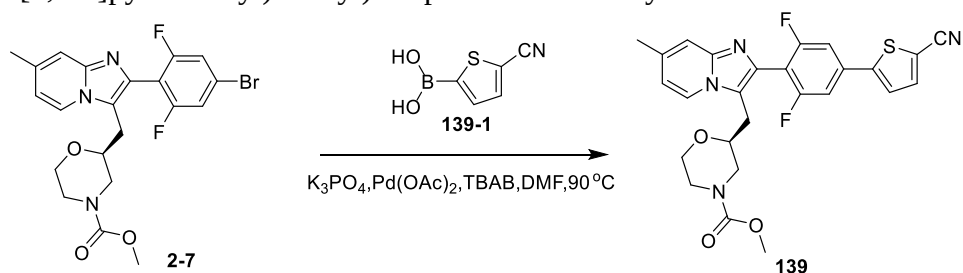


Intermediate 138-2 (200 mg, 0.41 mmol) was dissolved in absolute methanol (3 mL), and sodium hydroxide solution (3 M, 0.4 mL) was added. The reaction system was reacted at 60 °C for 1 h. After the reaction was completed, the reaction system was added dropwise with diluted hydrochloric acid (1 M) to adjust the pH to 6, added with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 138** (41.2 mg, 25% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 492.7$.

^1H NMR (400 MHz, MeOD) δ 8.85 (d, $J = 7.3$ Hz, 1H), 7.72 (s, 1H), 7.64 (d, $J = 9.3$ Hz, 2H), 7.42 - 7.41 (m, 1H), 6.99 (d, $J = 4.0$ Hz, 1H), 6.91 (d, $J = 4.0$ Hz, 1H), 4.02 - 3.99 (m, 1H), 3.85 - 3.75 (m, 3H), 3.69 (s, 3H), 3.39 - 3.34 (m, 1H), 3.28 - 3.26 (m, 2H), 3.00 - 2.88 (m, 1H), 2.81 - 2.71 (m, 1H), 2.65 (s, 3H).

Example 139

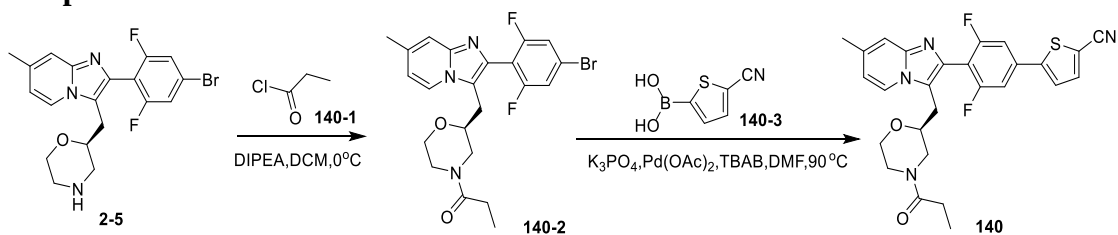
Preparation of methyl (S)-2-((2-(4-(5-cyanothiophen-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



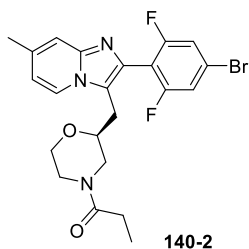
Intermediate 2-7 (100 mg, 0.21 mmol) was dissolved in anhydrous *N,N*-dimethylformamide (2 mL), and (5-cyanothiophen-2-yl)boronic acid (32 mg, 0.21 mmol), potassium phosphate (132 mg, 0.62 mmol), palladium acetate (9.4 mg, 0.042 mmol) and tetrabutylammonium bromide (20 mg, 0.062 mmol) were added under nitrogen atmosphere. The reaction system was reacted at $90^\circ C$ for 12 h. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (20 mL), and extracted with ethyl acetate (20 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 139** (3.8% yield) in the form of a yellow solid.

LC-MS: $[M+H]^+ = 509.7$. ^1H NMR (400 MHz, MeOD): δ 8.47 (d, $J = 6.8$ Hz, 1H), 7.83 (d, $J = 4.0$ Hz, 1H), 7.71 (d, $J = 4.0$ Hz, 1H), 7.59 (d, $J = 8.3$ Hz, 2H), 7.40 (s, 1H), 6.95 (d, $J = 7.0$ Hz, 1H), 3.79-3.89 (m, 3H), 3.67 (s, 3H), 3.63-3.59 (m, 1H), 3.40-3.37 (m, 1H), 3.14-3.13 (m, 2H), 2.93-2.88 (m, 1H), 2.70-2.61 (m, 1H), 2.49 (s, 3H).

Example 140

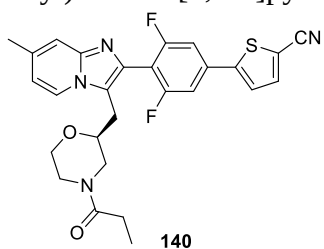


Step (1) Preparation of (S)-1-(2-((2-(4-bromo-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholino)propan-1-one



Intermediate 2-5 (500 mg, 1.18 mmol) was dissolved in anhydrous dichloromethane (5 mL), and under nitrogen atmosphere, *N,N'*-diisopropylethylamine (304 mg, 2.36 mmol) was added, followed by the dropwise addition of propionyl chloride (131 mg, 1.42 mmol) with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 2 h. After the reaction was completed, the reaction system was added with water (40 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (40 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 140-2** (450 mg, 79% yield) in the form of a brown solid, LC-MS: [M+H]⁺ = 478.7.

Step (2) Preparation of (S)-5-(3,5-difluoro-4-(7-methyl-3-((4-propionylmorpholin-2-yl)methyl)imidazo[1,2-a]pyridin-2-yl)phenyl)thiophene-2-carbonitrile

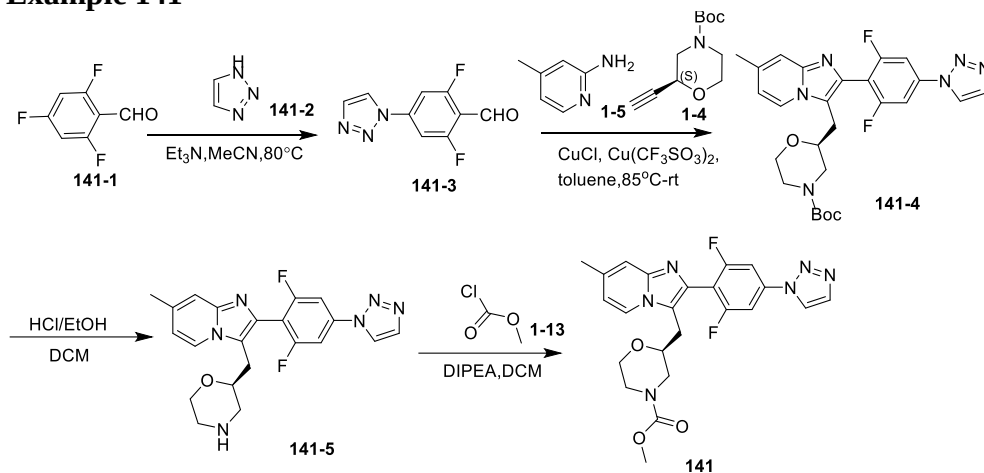


Intermediate 140-2 (100 mg, 0.21 mmol) was dissolved in anhydrous *N,N*-dimethylformamide (2 mL), and (5-cyanothiophen-2-yl)boronic acid (32 mg, 0.21 mmol), potassium phosphate (134 mg, 0.63 mmol), palladium acetate (9.6 mg, 0.043 mmol) and tetrabutylammonium bromide (22 mg, 0.068 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 90 °C for 12 h. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (20 mL), and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 140** (14.7 mg, 13.9% yield) in the form of a yellow solid.

LC-MS: [M + H]⁺ = 507.6. ¹H NMR (400 MHz, MeOD): δ8.45 (brs, 1H), 7.83 (d, J = 4.0 Hz, 1H), 7.70 (d, J = 3.8 Hz, 1H), 7.58 (t, 2H), 7.37 (brs, 1H), 6.95 – 6.87 (m, 1H), 4.32-4.23

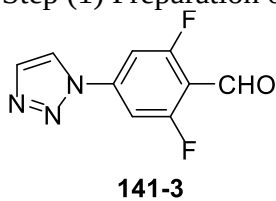
(m, 1H), 3.84-3.81(m, 1H), 3.77 – 3.68 (m, 1H), 3.65-3.56 (m, 1H), 3.44 – 3.36 (m, 1H), 3.16-3.15 (m, 2H), 2.96 – 2.85 (m, 1H), 2.76 – 2.63 (m, 1H), 2.48 (s, 3H), 2.40 – 2.27 (m, 2H), 1.09-1.03 (m, 3H).

Example 141



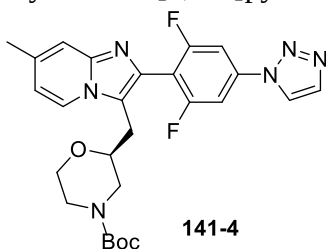
5

Step (1) Preparation of 2,6-difluoro-4-(1H-1,2,3-triazol-1-yl)benzaldehyde



2,4,6-trifluorobenzaldehyde (4.0 g, 24.98 mmol) was dissolved in acetonitrile (40 mL), and 1H-1,2,3-triazole (1.7 g, 24.98 mmol) and triethylamine (7.6 g, 74.95 mmol) were added. The reaction system was stirred at 80 °C overnight. After the reaction was completed, the reaction system was concentrated by rotary evaporation. The residue was purified by silica gel column chromatography to give **intermediate 141-3** in the form of a light yellow oil (530 mg, 10.2% yield). LC-MS: $[M+H]^+ = 209.9$.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(1H-1,2,3-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

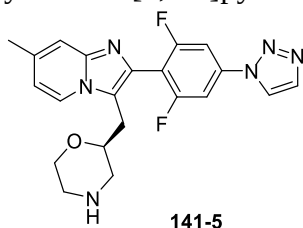


Intermediate 141-3 (520 mg, 2.49 mmol), **intermediate 1-5** (268 mg, 2.49 mmol), **intermediate 1-4** (525 mg, 2.49 mmol), copper(I) chloride (74 mg, 0.75 mmol) and copper(II) trifluoromethanesulfonate (271 mg, 0.75 mmol) were dissolved in anhydrous toluene (5 mL) under nitrogen atmosphere. The reaction system was heated to 85 °C, added with *N,N'*-

20

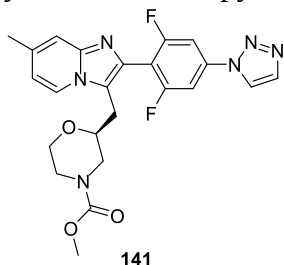
dimethylacetamide (0.2 mL), and stirred at 85 °C overnight. After the reaction was completed, the reaction system was filtered, and the filtrate was added with water (20 mL), and extracted with dichloromethane (15 mL × 2). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 141-4** (80 mg, 6.7% yield) in the form of a colorless oil, LC-MS: $[M+H]^+ = 510.8$.

Step (3) Preparation of (S)-2-((2-(2,6-difluoro-4-(1*H*-1,2,3-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



Intermediate 141-4 (80 g, 0.16 mmol) was dissolved in anhydrous dichloromethane (1 mL), and a solution of HCl in ethanol (0.3 mL) was added. The reaction system was stirred at room temperature for 3 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give **intermediate 141-5** (128 mg, crude product) in the form of a white solid, LC-MS $[M+H]^+ = 410.8$.

Step (4) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1*H*-1,2,3-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

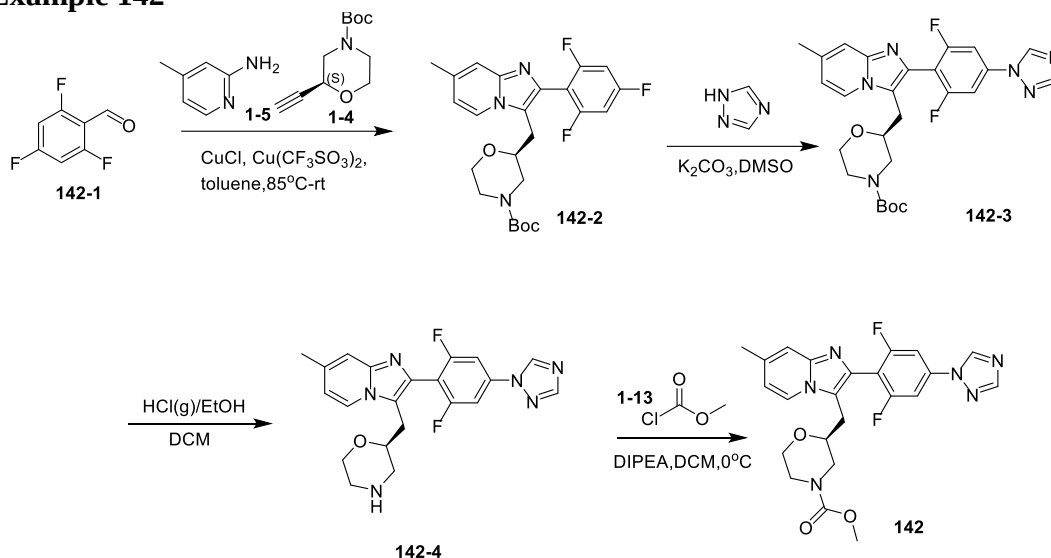


Intermediate 141-5 (128 mg, 0.16 mmol, crude product) was dissolved in anhydrous dichloromethane (5 mL), and under nitrogen atmosphere, *N,N'*-diisopropylethylamine (200 mg, 1.56 mmol) was added, followed by the dropwise addition of a solution of methyl chloroformate in dichloromethane (10 mg/mL, 1.8 mL, 0.19 mmol) with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 2 h. After the reaction was completed, the reaction system was added with water (20 mL), and extracted with dichloromethane (10 mL × 3). The organic phases were combined, washed with saturated brine (10 mL), dried over sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by silica gel column chromatography to give a product.

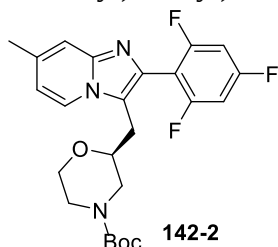
The product was separated by preparative by high performance liquid chromatography to give **compound 141** (14 mg, 19.35% yield) in the form of a white solid, LC-MS: $[M+H]^+ = 468.8$.

^1H NMR (400 MHz, CDCl_3) δ 8.23 (d, $J = 7.1$ Hz, 1H), 8.08 (d, $J = 1.0$ Hz, 1H), 7.93 (d, $J = 1.0$ Hz, 1H), 7.55 (d, $J = 7.6$ Hz, 2H), 7.47 (s, 1H), 6.75 (d, $J = 6.7$ Hz, 1H), 4.05 – 3.80 (m, 3H), 3.70 (s, 3H), 3.65 – 3.57 (m, 1H), 3.46 – 3.36 (m, 1H), 3.10 – 2.85 (m, 3H), 2.70 – 2.60 (m, 1H), 2.46 (s, 3H).

Example 142



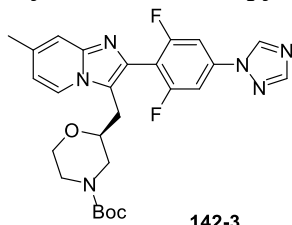
Step (1) Preparation of *tert*-butyl (S)-2-((7-methyl-2-(2,4,6-trifluorophenyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



2,4,6-trifluorobenzaldehyde (3.0 g, 18.74 mmol), 4-methylpyridin-2-amine (2.0 g, 18.74 mmol), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (4 g, 18.74 mmol), copper(I) chloride (562 mg, 5.68 mmol) and copper(II) trifluoromethanesulfonate (2.1 g, 5.68 mmol) were dissolved in toluene solution (30 mL) under nitrogen atmosphere. The reaction system was reacted at 85 °C for 10 min, added with *N,N*-dimethylacetamide (0.5 mL), and reacted at 85 °C for another 5 h. Then the reaction system was reacted at room temperature overnight. After the reaction was completed, the reaction system was filtered, and the filtrate was added with water (50 mL), and extracted with dichloromethane (100 mL $\times$ 3). The organic phases were combined, washed with saturated brine (50 mL $\times$ 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by

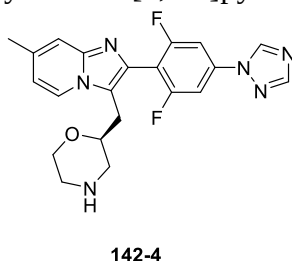
silica gel column chromatography to give **intermediate 142-2** (3.0 g, 34.8% yield) in the form of a yellow oil, MS $[M+H]^+ = 461.8$.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(1*H*-1,2,4-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



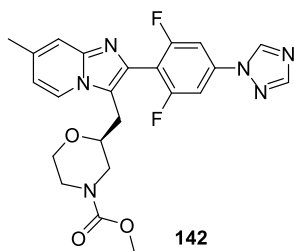
Intermediate 142-2 (300 mg, 0.65 mmol) was dissolved in anhydrous dimethyl sulfoxide (3 mL), and 1,2,4-triazole (40 mg, 0.58 mmol) and potassium carbonate (180 mg, 1.30 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 85 °C for 1 h. Then the reaction system was cooled to room temperature, added with water (20 mL), and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 142-3** (150 mg, 45.2% yield) in the form of a yellow oil. MS $[M+H]^+ = 510.8$.

Step (3) Preparation of (S)-2-((2-(2,6-difluoro-4-(1*H*-1,2,4-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



Intermediate 142-3 (150 mg, 0.29 mmol) was dissolved in dichloromethane (2 mL), and a solution of HCl in ethanol (33%, 0.5 mL) was added dropwise with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. Then the reaction system was adjusted to pH 8 with saturated sodium bicarbonate solution, diluted with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give **intermediate 142-4** (130 mg, crude product) in the form of a yellow solid, MS $[M+H]^+ = 410.8$.

Step (4) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(1*H*-1,2,4-triazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

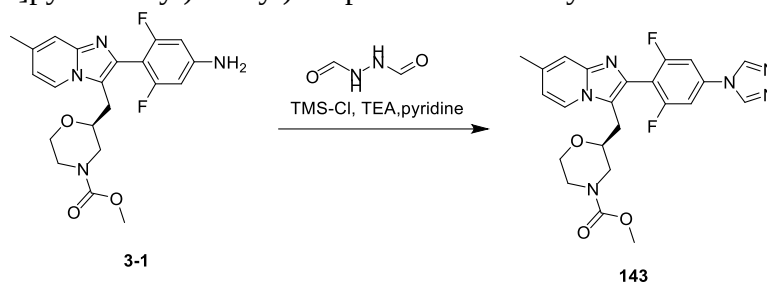


Intermediate 142-4 (130 mg, 0.29 mmol) and *N,N*-diisopropylethylamine (123 mg, 0.95 mmol) were dissolved in dichloromethane (2 mL), and methyl chloroformate (36 mg, 0.38 mmol) was added dropwise under nitrogen atmosphere with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 1 h. Then the reaction system was added with water (20 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 142** (30.4 mg, 20.5% yield) in the form of a yellow solid. MS $[M+H]^+ = 468.7$.

¹H NMR (400 MHz, MeOD) δ 9.35 (s, 1H), 8.88 (d, *J* = 7.1 Hz, 1H), 8.29 (s, 1H), 7.97 (d, *J* = 8.7 Hz, 2H), 7.75 (s, 1H), 7.44 (dd, *J* = 7.1, 1.4 Hz, 1H), 4.05 – 4.02 (m, 1H), 3.85 – 3.74 (m, 3H), 3.69 (s, 3H), 3.39 – 3.35 (m, 1H), 3.31 – 3.22 (m, 2H), 2.98 – 2.88 (m, 1H), 2.81 – 2.71 (m, 1H), 2.67 (s, 3H).

Example 143

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(4*H*-1,2,4-triazol-4-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



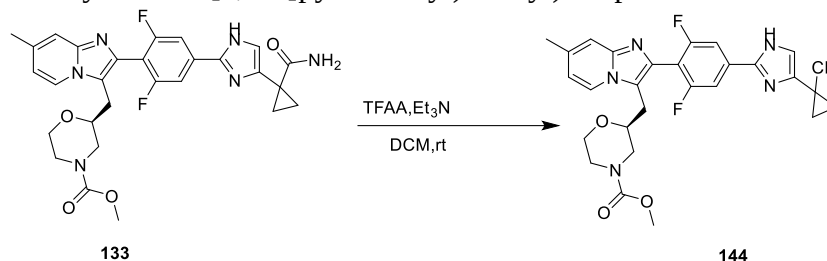
Intermediate 3-1 (75 mg, 0.18 mmol) and diformylhydrazide (486 mg, 0.54 mmol) were dissolved in anhydrous pyridine (10 mL). The reaction system was cooled to 0 °C, added dropwise with triethylamine (128 mg, 1.26 mmol) and trimethylchlorosilane (294 mg, 2.7 mmol) sequentially under nitrogen atmosphere, and stirred under reflux for 24 h. Then the reaction system was concentrated under reduced pressure, added with saturated brine (10 mL), adjusted to pH 7-8 with saturated sodium bicarbonate solution, and extracted with DCM (10 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated.

The crude product was purified by preparative thin-layer chromatography to give **compound 143** (30 mg, 36% yield) in the form of a white solid. MS $[M+H]^+ = 469.2$.

¹H NMR (400 MHz, d₆-DMSO) δ 9.31 (s, 2H), 8.44 (d, $J = 7.1$ Hz, 1H), 7.85 (d, $J = 8.2$ Hz, 2H), 7.37 (s, 1H), 6.85 (d, $J = 7.1$ Hz, 1H), 3.82 – 3.84 (m, 1H), 3.70 – 3.61 (m, 2H), 3.56 (s, 3H), 3.52 – 3.45 (m, 1H), 3.29 – 3.21 (m, 1H), 3.12 – 3.00 (m, 2H), 2.90 – 2.71 (m, 1H), 2.59 – 2.45 (m, 1H), 2.39 (s, 3H).

Example 144

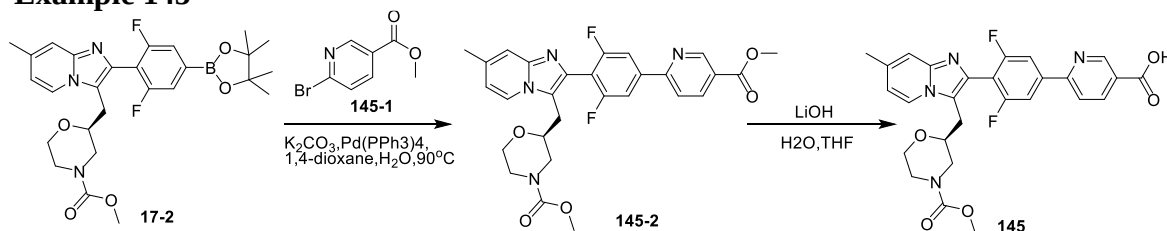
Preparation of methyl (S)-2-((2-(4-(4-(1-cyanochloropropyl)-1H-imidazol-2-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



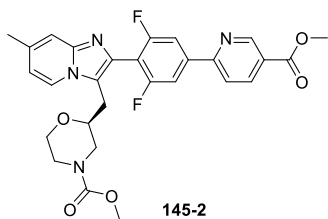
Compound 133 (150 mg, 0.27 mmol) and triethylamine (121 mg, 1.20 mmol) were dissolved in dichloromethane (3 mL). Under nitrogen atmosphere, the reaction system was cooled to 0 °C, added dropwise with trifluoroacetic anhydride (125 mg, 0.60 mmol), and stirred at room temperature overnight. Then the reaction system was added with water (10 mL) to quench the reaction, and extracted with dichloromethane (10 mL $\times$ 3). The organic phases were combined, washed with saturated brine (10 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by preparative chromatography to give **compound 144** (23.7 mg, 16% yield) in the form of a white solid. LC-MS $[M+H]^+ : 533$.

¹H NMR (400 MHz, CDCl₃) δ 8.35 (s, 1H), 7.62 (s, 1H), 7.44 (d, $J = 8.2$ Hz, 2H), 7.23 (s, 1H), 6.91 (d, $J = 7.1$ Hz, 1H), 4.00 – 3.75 (m, 3H), 3.68 (s, 3H), 3.62 – 3.53 (m, 1H), 3.44 – 3.33 (m, 1H), 3.03 – 2.82 (m, 3H), 2.66 – 2.56 (m, 1H), 2.55 (s, 3H), 1.68 – 1.58 (m, 4H).

Example 145

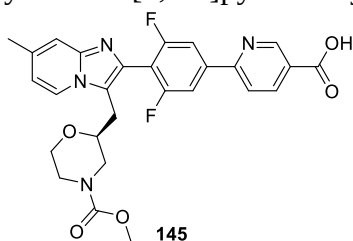


Step (1) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methoxycarbonyl)pyridin-2-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 17-2 (130 mg, 0.25 mmol) was dissolved in anhydrous 1,4-dioxane (5 mL) under nitrogen atmosphere, and methyl 6-bromonicotinate (64 mg, 0.30 mmol), potassium carbonate (103 mg, 0.74 mmol), and tetrakis(triphenylphosphine)palladium (30 mg, 0.025 mmol) were added, followed by the addition of water (0.5 mL). The reaction system was reacted at 90 °C for 2 h. Then the reaction system was cooled to room temperature, added with water (20 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous magnesium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 145-2** (70 mg, 53% yield) in the form of a yellow solid. LC-MS $[M+H]^+ = 553.2$.

Step (2) Preparation of methyl (S)-2-((2,6-difluoro-4-nicotinic acid)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)-methyl)morpholine-4-carboxylate

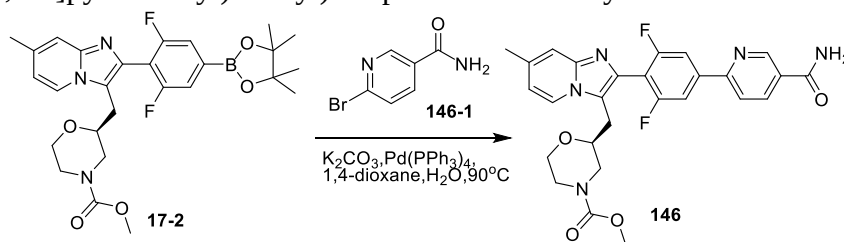


Intermediate 145-2 (70 mg, 0.13 mmol) was dissolved in tetrahydrofuran (1.5 mL), and lithium hydroxide solution (3 M, 0.13 mL) was added dropwise. The reaction system was reacted at room temperature for 1 h. After the reaction was completed, the reaction system was added dropwise with diluted hydrochloric acid (1 M) to adjust the pH to 6, added with water (10 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 145** (30.1 mg, 44% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 523.6$.

^1H NMR (400 MHz, MeOD) δ 9.30 (d, $J = 2.1$ Hz, 1H), 8.88 (d, $J = 7.1$ Hz, 1H), 8.53 (dd, $J = 8.3, 2.1$ Hz, 1H), 8.21 (d, $J = 8.3$ Hz, 1H), 8.13 (d, $J = 9.3$ Hz, 2H), 7.75 (s, 1H), 7.44 (dd, $J = 7.1, 1.5$ Hz, 1H), 4.03-4.01 (m, 1H), 3.87-3.70 (m, 3H), 3.69 (s, 3H), 3.40-3.36 (m, 1H), 3.31-3.23 (m, 2H), 2.98-2.89 (m, 1H), 2.82-2.71 (m, 1H), 2.67 (s, 3H).

Example 146

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(formamido)pyridinyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

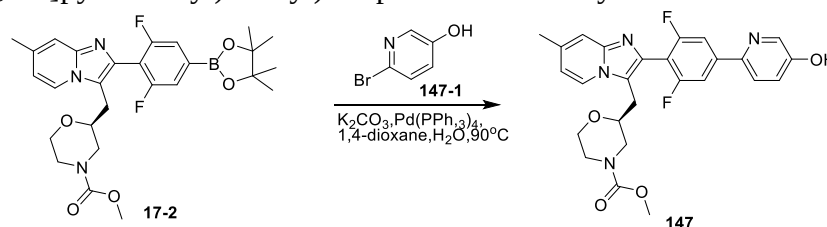


Intermediate 17-2 (120 mg, 0.23 mmol) was dissolved in 1,4-dioxane (3 mL) under nitrogen atmosphere, and 6-bromonicotinamide (55 mg, 0.27 mmol), potassium carbonate (95 mg, 0.69 mmol) and tetrakis(triphenylphosphine)palladium (27 mg, 0.023 mmol) were added, followed by the addition of water (0.5 mL). The reaction system was reacted at 90 °C for 2 h. Then the reaction system was cooled to room temperature, added with water (20 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous magnesium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 146** (16.9 mg, 14% yield) in the form of a yellow solid.

LC-MS: $[M+H]^+ = 522.6$. ^1H NMR (400 MHz, MeOD) : δ 9.21 (d, $J = 2.1$ Hz, 1H), 8.88 (d, $J = 7.2$ Hz, 1H), 8.42 (dd, $J = 8.3, 2.3$ Hz, 1H), 8.20 (d, $J = 8.3$ Hz, 1H), 8.12 (d, $J = 9.4$ Hz, 2H), 7.75 (s, 1H), 7.44 (dd, $J = 7.1, 1.4$ Hz, 1H), 4.04 – 4.01 (m, 1H), 3.86 – 3.71 (m, 3H), 3.70 (s, 3H), 3.40 – 3.36 (m, 1H), 3.31 – 3.25 (m, 2H), 2.99 – 2.90 (m, 1H), 2.83 – 2.73 (m, 1H), 2.67 (s, 3H).

Example 147

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-hydroxypyridinyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 17-2 (130 mg, 0.25 mmol) was dissolved in 1,4-dioxane (5 mL) under nitrogen atmosphere, and 2-bromo-5-hydroxypyridine (51 mg, 0.30 mmol), potassium carbonate (102 mg, 0.74 mmol) and tetrakis(triphenylphosphine)palladium (28 mg, 0.025 mmol) were added, followed by the addition of water (0.5 mL). The reaction system was reacted at 90 °C for

2 h. After the reaction was completed, the reaction system was cooled to room temperature, added with water (20 mL), and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (30 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by high performance liquid chromatography to give **compound 147** (62.7 mg, 51.4% yield) in the form of a yellow solid.

LC-MS: $[M + H]^+ = 495.2$. ^1H NMR (400 MHz, DMSO) : δ 10.49 (s, 1H), 8.91 (d, $J = 7.1$ Hz, 1H), 8.30 (d, $J = 2.8$ Hz, 1H), 8.08 (d, $J = 8.7$ Hz, 1H), 8.01 (d, $J = 9.7$ Hz, 2H), 7.80 (s, 1H), 7.44 (dd, $J = 7.1, 1.2$ Hz, 1H), 7.34 (dd, $J = 8.7, 2.9$ Hz, 1H), 3.94 (d, $J = 12.8$ Hz, 1H), 3.64-3.62 (m, 3H), 3.58 (s, 3H), 3.34-3.12 (m, 3H), 2.88-2.76 (m, 1H), 2.68 – 2.62 (m, 1H), 2.58 (s, 3H).

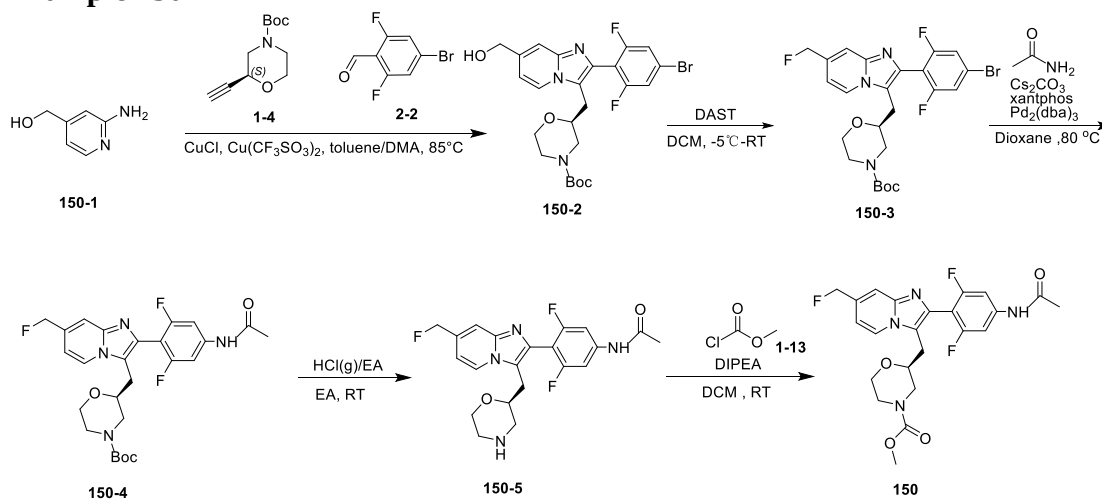
Examples 148-170

Examples 148, 151, 152, 154, 155, 156 and 157 were prepared using the corresponding starting materials according to the method as described in Example 2. Examples 165-166 were prepared according to the synthesis method as described in Example 10; and Example 170 was prepared according to the synthesis method as described in Example 25.

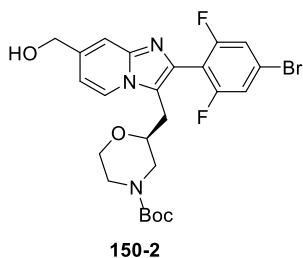
Number	Compound structure	ESI-MS [M + H] ⁺
Example 148		463.2
Example 151		495.1
Example 152		495.4
Example 154		477.2
Example 155		475.6
Example 156		457.2

Example 157		493.2
Example 165		453.4
Example 166		473.5
Example 170		530.3

Example 150

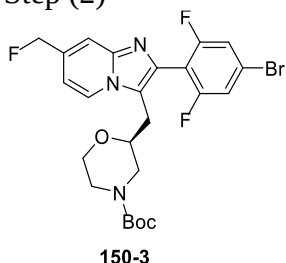


Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-(hydroxymethyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



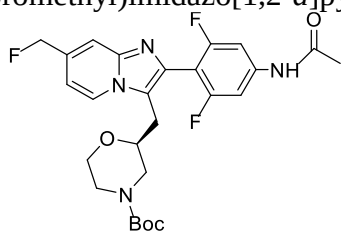
Intermediate 150-1 (2 g, 16 mmol), **intermediate 1-4** (3.4 g, 16 mmol), **intermediate 2-2** (3.6 g, 16 mmol), CuCl (0.5 g, 6 mmol) and Cu(CF₃SO₃)₂ (1.75 g, 6 mmol) were added to toluene (60 mL) sequentially. The reaction system was purged with N₂ four times, and stirred at room temperature for 15 min. Then the reaction system was heated to 85 °C, added with DMA (1 mL), heated and reacted for 5 h, and cooled and stirred overnight (16.5 h) after the heating was stopped. After the reaction was completed as detected by LCMS, the reaction system was diluted with DCM (100 mL), added with water (80 mL) and 25% aqueous ammonia (15 mL), stirred for 30 min, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (100 mL × 2). The DCM phases were combined, washed with saturated aqueous ammonium chloride solution and saturated brine sequentially, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give a crude product. The crude product was purified by column chromatography to give **intermediate 150-2** (3.4 g) in the form of a light yellow oil.

Step (2)



Intermediate 150-2 (1.3 g, 2.42 mmol) was dissolved in DCM (30 mL). The reaction system was purged with N₂ three times, and cooled to -5 °C below. Then DAST (0.40 mL, 3.03 mmol) was diluted with DCM (5 mL), and added dropwise to the reaction system. The resulting reaction system was warmed to room temperature and reacted for 16 h. After the reaction was completed as detected by TLC, the reaction system was cooled to 5 °C below, diluted with DCM, slowly poured into saturated NaHCO₃ solution at 5 °C below, stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (30 mL × 2). The DCM phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 150-3** (0.32 g) in the form of a yellow oil.

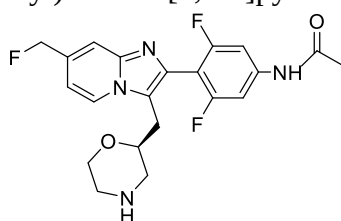
Step (3) Preparation of *tert*-butyl (S)-2-((2-(4-acetamido-2,6-difluorophenyl)-7-(fluoromethyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



150-4

Intermediate 150-3 (0.32 g, 0.59 mmol), acetamide (0.05 g, 0.71 mmol), Cs₂CO₃ (0.29 g, 0.71 mmol), and Xantphos (0.07 g, 0.12 mmol) were added to dioxane (8 mL) sequentially. The reaction system was purged with N₂ three times, added with Pd₂(dba)₃ (0.05 g, 0.06 mmol), purged with N₂ three times, and reacted at 80 °C for 19 h. After the reaction was completed as detected by TLC, the reaction system was diluted with DCM (15 mL) and water (30 mL), stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (15 mL × 2). The DCM phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give **intermediate 150-4** (0.47 g, crude product) in the form of a red oil, which was used directly in the next step.

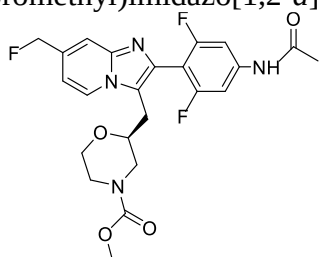
Step (4) Preparation of (S)-N-(3,5-difluoro-4-(7-(fluoromethyl)-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)phenyl)acetamide



150-5

Intermediate 150-4 (0.47 g, crude product) was dissolved in EA (5 mL), and a solution of HCl in EA (15 mL) was added. The reaction system was reacted at room temperature for 5.5 h. After the reaction was completed as detected by TLC, the reaction system was diluted with EA (10 mL) and water (10 mL), back-extracted with stirring to give an aqueous phase, and the EA phase was back-extracted with water (10 mL × 1). The aqueous phases were combined, added with DCM (15 mL), adjusted to pH 9-10 with Na₂CO₃ solid, stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (15 mL × 2). The DCM phases were combined, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give **intermediate 150-5** (0.2 g) in the form of a red oil.

Step (5) Preparation of methyl (S)-2-((2-(4-acetamido-2,6-difluorophenyl)-7-(fluoromethyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

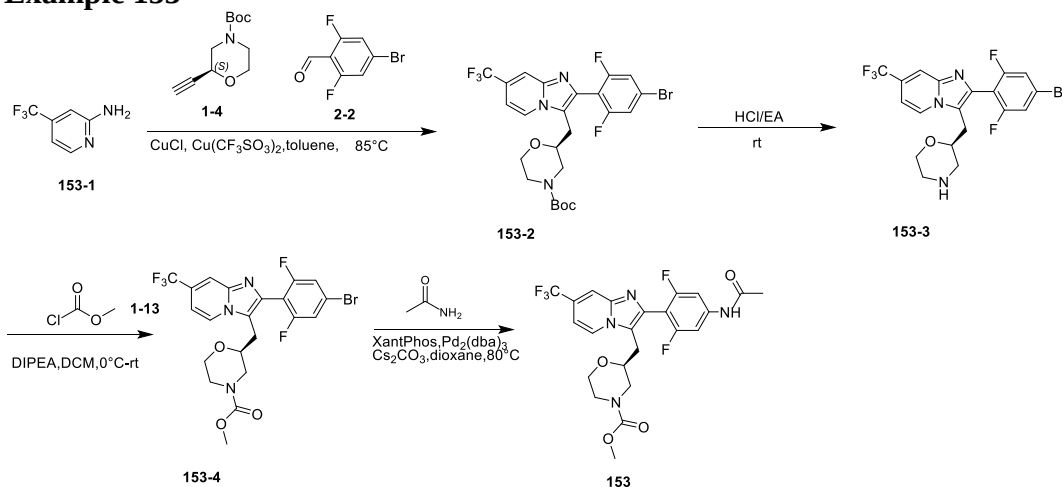


150

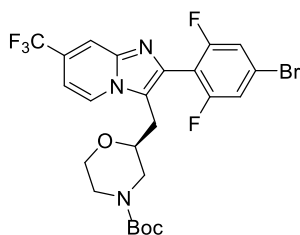
Intermediate 150-5 (100 mg, 0.227 mmol) was dissolved in DCM (4 mL), and DIPEA (0.4 mL) was added. The reaction system was purged with N₂ once, and cooled to 5 °C below. Then the reaction system was added dropwise with methyl chloroformate (43 mg, 0.504 mmol), warmed to room temperature and reacted for 1 h. After the reaction was completed as detected by TLC, the reaction system was diluted with DCM (15 mL) and water (20 mL), stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (15 mL × 2). The DCM phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give a crude product. The crude product was separated by prep-HPLC, and lyophilized to give **compound 150** (20.2 mg, 98.34% purity) in the form of a white solid.

LC-MS: [M+H]⁺=477.25. ¹H NMR (400 MHz, CD₃OD) δ 8.52 (d, J = 7.1 Hz, 1H), 8.52 (d, J = 7.1 Hz, 8H), 8.52 (d, J = 7.1 Hz, 1H), 7.54 (s, 7H), 7.46 – 7.37 (m, 13H), 6.99 (d, J = 7.1 Hz, 8H), 5.54 (s, 7H), 5.42 (s, 7H), 4.98 – 4.83 (m, 169H), 4.79 (d, J = 15.2 Hz, 2H), 4.58 (s, 9H), 3.80 (t, J = 13.0 Hz, 25H), 3.66 (d, J = 11.2 Hz, 19H), 3.58 (s, 8H), 3.33 (tt, J = 6.6, 3.2 Hz, 27H), 3.21 – 3.04 (m, 15H), 2.89 (s, 10H), 2.63 (s, 15H), 2.17 (s, 18H).

Example 153



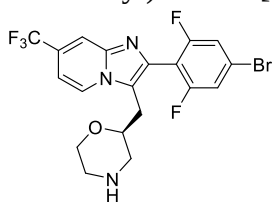
Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-(trifluoromethyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



153-2

Compound 153-1 (0.77 g, 4.74 mmol), **intermediate 1-4** (1.0 g, 4.74 mmol), **intermediate 2-2** (1.04 g, 4.74 mmol), CuCl (0.14 g, 1.42 mmol) and Cu(CF₃SO₃)₂ (0.51 g, 1.42 mmol) were dissolved in toluene (10 mL) sequentially. The reaction system was purged with N₂ four times, and stirred at room temperature for 15 min. Then the reaction system was heated to 85 °C, added with DMA (0.3 mL), heated and reacted for 6 h, and cooled and stirred overnight (16 h) after the heating was stopped. After the reaction was completed as detected by LCMS, the reaction system was diluted with DCM (100 mL), added with water (100 mL) and 25% aqueous ammonia (10 mL), stirred for 10 min, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (50 mL × 3). The DCM phases were combined, washed with saturated aqueous ammonium chloride solution and saturated brine sequentially, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give a crude product. The crude product was purified by silica gel column chromatography (a small amount of methanol) to give **intermediate 153-2** (0.8 g, 86.37% purity) in the form of a yellow foamy solid. LC-MS: [M+H]⁺ = 578.03.

Step (2) Preparation of (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-(trifluoromethyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine

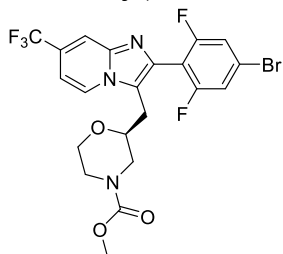


153-3

Intermediate 153-2 (0.8 mg, 1.4 mmol) was dissolved in EA (10 mL). The reaction system was cooled to °C below. Then the reaction system was added with a solution of HCl in EA (3 M, 15 mL), and reacted at room temperature for 1 h and 50 min. After the reaction was completed as detected by LCMS, the reaction system was diluted with EA (10 mL) and back-extracted with water (20 mL × 2). The aqueous phases were combined, added with DCM (20 mL), adjusted to pH 8-9 with Na₂CO₃ solid, stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (20 mL × 2). The DCM phases were combined, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure

to give **intermediate 153-3** (0.65 g, 94.48% purity) in the form of a yellow oil. LC-MS: $[M+H]^+ = 476.0$.

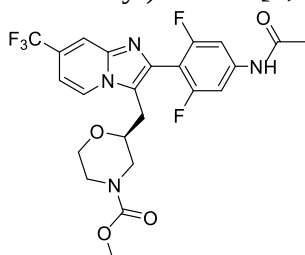
Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-(trifluoromethyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



153-4

Intermediate 153-3 (650 mg, 1.4 mmol) was dissolved in DCM (10 mL), and DIPEA (0.21 mL) was added. The reaction system was purged with N_2 once, and cooled to 5 °C below. Then the reaction system was added dropwise with methyl chloroformate (150 mg, 1.6 mmol), and reacted at room temperature for 2 h. After the reaction was completed as detected by LCMS, the reaction system was diluted with DCM (10 mL) and water (30 mL), stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (15 mL $\times$ 2). The DCM phases were combined, dried over anhydrous sodium sulfate, and concentrated at 40 °C to give **intermediate 153-4** (650 mg, 93.67% purity, crude product) in the form of a yellow foamy solid. LC-MS: $[M+H]^+ = 534.05$.

Step (4) Preparation of methyl (S)-2-((2-(4-acetamido-2,6-difluorophenyl)-7-(trifluoromethyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



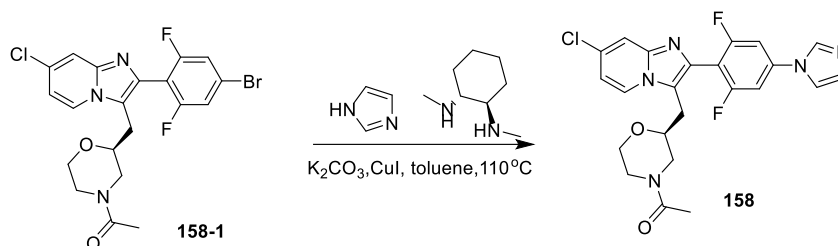
153

Intermediate 153-4 (300 mg, 0.56 mmol), acetamide (70 mg, 0.84 mmol), Cs_2CO_3 (350 mg, 0.84 mmol), Xantphos (40 mg, 0.056 mmol), and $Pd_2(dba)_3$ (60 mg, 0.056 mmol) were dissolved in dioxane (10 mL) sequentially. The reaction system was purged with N_2 four times and reacted at 80 °C for 20 h. After the reaction was completed as detected by LCMS, the reaction system was diluted with DCM (15 mL) and water (30 mL), stirred, and left to stand to separate the DCM phase. The aqueous phase was extracted with DCM (15 mL $\times$ 2). The DCM phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and filtered, and the filtrate was concentrated under reduced pressure to give a crude product. The crude

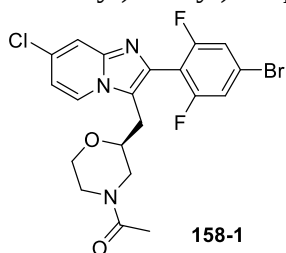
product was separated by prep-HPLC (H₂O/CH₃CN), and lyophilized to give **compound 153** (60 mg, 99.01% purity) in the form of a white solid. LC-MS: [M+H]⁺ = 513.15.

¹H NMR (400 MHz, d-DMSO) δ 10.45 (s, 1H), 8.74 (d, J = 7.3 Hz, 1H), 8.08 (s, 1H), 7.45 (d, J = 9.9 Hz, 2H), 7.24 (dd, J = 7.3, 1.8 Hz, 1H), 3.78 (d, J = 12.9 Hz, 1H), 3.65 (t, J = 10.7 Hz, 2H), 3.56 (s, 3H), 3.51 (s, 1H), 3.23 (td, J = 11.8, 2.8 Hz, 1H), 3.10 (dd, J = 15.4, 7.2 Hz, 2H), 2.81 (s, 1H), 2.54 (s, 1H), 2.11 (s, 3H).

Example 158

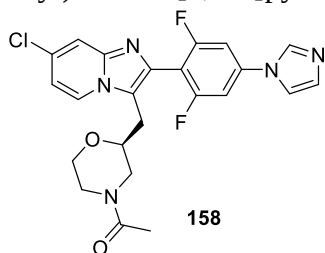


Step (1) Preparation of (S)-1-(2-((2-(4-bromo-2,6-difluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholino)ethan-1-one



The compound was prepared according to the preparation method for intermediate 2-7. LC-MS: [M+H]⁺ = 484.0.

Step (2) Preparation of (S)-1-(2-((7-chloro-2-(2,6-difluoro-4-(1H-imidazol-1-yl)phenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholino)ethan-1-one



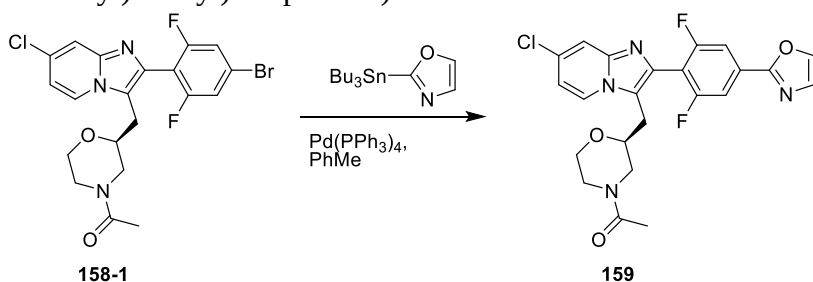
Intermediate 158-1 (120 mg, 0.25 mmol) was dissolved in toluene (2 mL), and imidazole (25 mg, 0.37 mmol), potassium carbonate (103 mg, 0.75 mmol), copper(I) iodide (4.7 mg, 0.025 mmol) and *trans*-(1R,2R)-N,N'-dimethyl-1,2-cyclohexanediamine (3.5 mg, 0.025 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 110 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (20 mL) and extracted with dichloromethane (50 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous

sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by preparative chromatography to give **compound 158** (13.6 mg, 12% yield) in the form of a white solid. MS $[M+H]^+ = 471.7$.

^1H NMR (400 MHz, MeOD) δ 8.59 (t, 1H), 8.39 (s, 1H), 7.79 (s, 1H), 7.65 – 7.63 (m, 1H), 7.61 – 7.57 (m, 2H), 7.23 (s, 1H), 7.09 – 7.00 (m, 1H), 4.33 – 4.24 (m, 1H), 3.92 – 3.54 (m, 3H), 3.47 – 3.36 (m, 1H), 3.23 – 3.12 (m, 2H), 3.01 – 2.63 (m, 1H), 2.53 – 2.45 (m, 1H), 2.06 (d, 3H).

Example 159

Preparation of (S)-1-(2-((7-chloro-2-(2,6-difluoro-4-(oxazol-2-yl)phenyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholino)ethan-1-one



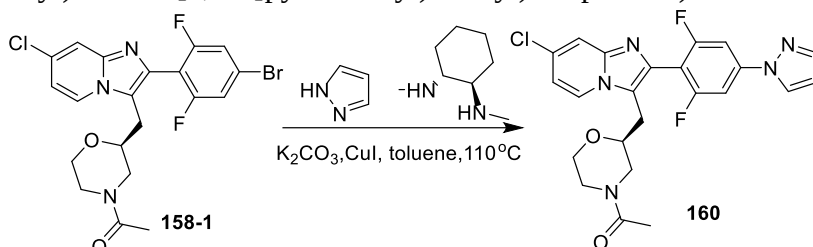
Intermediate 158-1 (4.2 g, 8.66 mmol) and tetrakis(triphenylphosphine)palladium (2.0 g, 1.73 mmol) were added to toluene (125 mL) sequentially. The reaction system was stirred under nitrogen atmosphere. Then the reaction system was injected with 2-(tributylstannyl)oxazole (4.9 g, 13.00 mmol) via a syringe, and heated to 80 °C and reacted overnight. Little **intermediate 158-1** was remained as detected by TLC, and the work-up was directly performed. The reaction system was cooled, added with saturated aqueous potassium fluoride solution (100 mL), and stirred for 30 min. Then the reaction was extracted with DCM (100 mL), followed by liquid separation. The aqueous phase was back-extracted with DCM (50 mL $\times$ 2). The organic phases were combined, washed with saturated brine (200 mL), dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The crude product was purified by preparative chromatography to give **compound 159** (0.49 g, 99% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 473.05$.

^1H NMR (400 MHz, dmso) δ 8.66 – 8.55 (m, 18H), 8.64 – 8.56 (m, 17H), 8.34 (s, 16H), 8.34 (s, 14H), 7.77 (dd, $J = 7.6, 2.5$ Hz, 45H), 7.77 (dd, $J = 7.6, 2.5$ Hz, 47H), 7.50 (s, 15H), 7.50 (s, 14H), 7.08 (td, $J = 7.1, 1.9$ Hz, 17H), 7.08 (td, $J = 7.1, 1.9$ Hz, 17H), 4.20 (d, $J = 12.8$ Hz, 13H), 4.20 (d, $J = 12.8$ Hz, 11H), 4.05 (d, $J = 12.0$ Hz, 9H), 4.05 (d, $J = 12.0$ Hz, 13H), 3.74 (d, $J = 12.8$ Hz, 9H), 3.70 (dd, $J = 30.3, 12.1$ Hz, 29H), 3.67 (d, $J = 11.5$ Hz, 17H), 3.55 (d, $J = 13.5$ Hz, 19H), 3.45 (dd, $J = 7.2, 3.2$ Hz, 11H), 3.58 – 3.20 (m, 61H), 3.26 (dd, $J = 11.6, 9.5$ Hz, 11H), 3.26 (dd, $J = 11.6, 9.5$ Hz, 31H), 3.26 (dd, $J = 11.6, 9.5$ Hz, 29H), 3.20 – 2.94 (m, 51H), 3.23 – 3.09 (m, 38H), 2.80 (dd, $J = 12.7, 10.7$ Hz, 9H), 2.80 (dd, $J = 12.7, 10.7$ Hz, 10H), 2.55 (d, $J =$

3.1 Hz, 6H), 2.64 – 2.45 (m, 22H), 2.39 – 2.22 (m, 12H), 2.36 – 2.24 (m, 11H), 1.92 (s, 47H), 1.92 (s, 48H).

Example 160

Preparation of (S)-1-(2-((7-chloro-2-(2,6-difluoro-4-(1H-pyrazol-1-yl)phenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholino)ethan-1-one

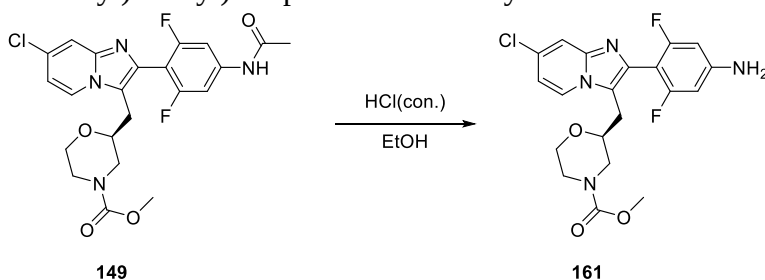


Intermediate 158-1 (170 mg, 0.35 mmol) was dissolved in toluene (2 mL), and pyrazole (36 mg, 0.53 mmol), potassium carbonate (145 mg, 1.05 mmol), copper(I) iodide (7 mg, 0.037 mmol) and *trans*-(1*R*,2*R*)-*N,N'*-dimethyl-1,2-cyclohexanediamine (5 mg, 0.035 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 110 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filter cake was washed with dichloromethane. The filtrate was added with water (30 mL) and extracted with dichloromethane (50 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by preparative chromatography to give **compound 160** (5.8 mg, 3.5% yield) in the form of a white solid. MS $[M+H]^+ = 471.7$.

¹H NMR (400 MHz, MeOD) δ 8.93 (dd, *J* = 7.4, 3.2 Hz, 1H), 8.47 (d, *J* = 2.6 Hz, 1H), 8.02 – 7.98 (m, 1H), 7.85 – 7.82 (m, 3H), 7.51 (dd, *J* = 7.4, 2.0 Hz, 1H), 6.65 – 6.64 (m, 1H), 4.43 – 4.40 (m, 0.5H), 4.29 – 4.26 (m, 0.5H), 3.92 – 3.89 (m, 1H), 3.86 – 3.80 (m, 1H), 3.72 – 3.69 (m, 1H), 3.46 – 3.36 (m, 1H), 3.31 – 3.21 (m, 2H), 3.08 – 3.02 (m, 1H), 2.75 – 2.70 (m, 0.5H), 2.58 – 2.52 (m, 0.5H), 2.08 (s, 3H).

Example 161

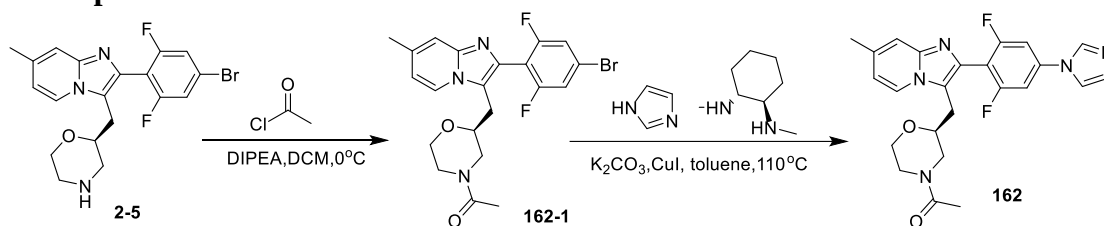
Preparation of methyl (S)-2-((2-(4-amino-2,6-difluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



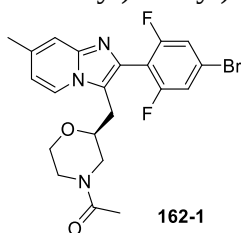
Compound 149 (0.3 g) was dissolved in EtOH (4 mL), and concentrated HCl (6 mL) was added. The reaction system was stirred at 100 °C for 2 h. The starting material was consumed, and a new spot was generated as detected by TLC. Then the reaction system was concentrated. The crude product was separated by preparative reverse phase chromatography to give **compound 161** (0.14 g) in the form of a white solid, MS $[M+H]^+ = 437.2$.

$^1\text{H-NMR}$ ($^1\text{H NMR}$ (400 MHz, DMSO) δ 8.44, 7.56, 6.88, 6.68, 6.68, 4.07, 4.05, 3.96, 3.93, 3.82, 3.78, 3.69, 3.51, 3.46, 2.73, 2.48.) (NB190027-03-02).

Example 162

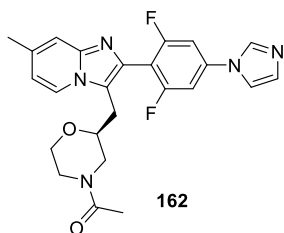


Step (1) Preparation of (S)-1-(2-((2-(4-bromo-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholino)ethan-1-one



Intermediate 2-5 (1.20 g, 2.84 mmol) and *N,N*-diisopropylethylamine (1.10 g, 8.51 mmol) were dissolved in dichloromethane (6 mL), and acetyl chloride (268 mg, 3.41 mmol) was added dropwise in an ice water bath with the temperature maintained at 0 °C. After the completion of the dropwise addition was completed, the reaction system was reacted at 0 °C for 2 h. After the reaction was completed, the reaction system was added with water (30 mL) and extracted with dichloromethane (50 mL $\times$ 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography (petroleum ether:ethyl acetate = 5:1) to give **intermediate 162-1** (700 mg, 53.8% yield) in the form of a yellow oil. MS $[M+H]^+ = 464.9$.

Step (2) Preparation of (S)-1-(2-((2-(2,6-difluoro-4-(1*H*-imidazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholino)ethan-1-one

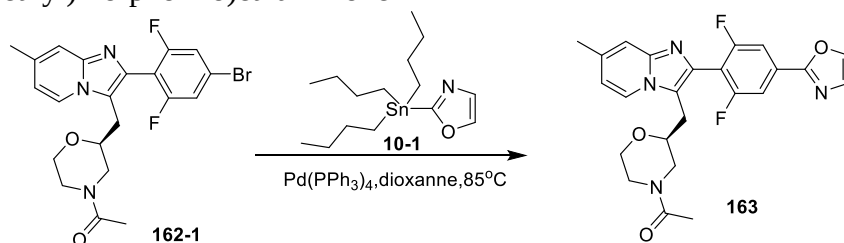


Intermediate 162-1 (150 mg, 0.32 mmol) was dissolved in toluene (2 mL), and imidazole (33 mg, 0.48 mmol), potassium carbonate (135 mg, 0.98 mmol), copper(I) iodide (6 mg, 0.032 mmol) and *trans*-(1*R*,2*R*)-*N,N'*-dimethyl-1,2-cyclohexanediamine (4.6 mg, 0.032 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 110 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (20 mL) and extracted with dichloromethane (30 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by preparative chromatography to give **compound 162** (21.3 mg, 14.6% yield) in the form of a white solid. MS [M+H]⁺ = 451.8.

¹H NMR (400 MHz, MeOD) δ 8.48 (t, 1H), 8.42 (s, 1H), 7.80 (s, 1H), 7.59 (t, 2H), 7.39 (s, 1H), 7.24 (s, 1H), 6.97 – 6.91 (m, 1H), 4.33 – 4.23 (m, 1H), 3.84 – 3.81 (m, 1H), 3.79 – 3.68 (m, 1H), 3.67 – 3.56 (m, 1H), 3.45 – 3.34 (m, 1H), 3.22 – 3.21 (m, 0.5H), 3.17 – 3.15 (m, 2H), 2.98 – 2.92 (m, 0.5H), 2.73 – 2.67 (m, 0.5H), 2.49 (s, 3H), 2.47 – 2.44 (m, 0.5H), 2.05 (d, 3H).

Example 163

Preparation of (S)-1-(2-((2-(2,6-difluoro-4-(oxazol-2-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholino)ethan-1-one



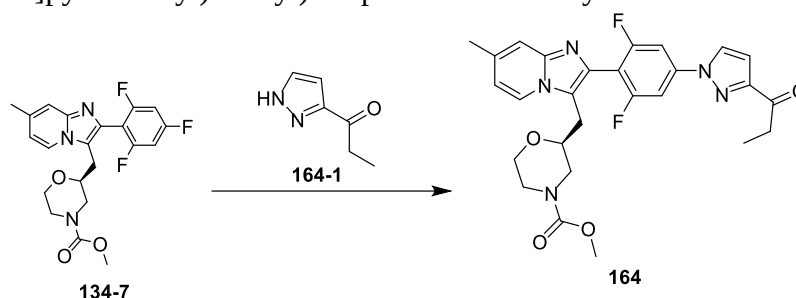
Intermediate 162-1 (150 mg, 0.32 mmol) was dissolved in 1,4-dioxane (2 mL), and 2-(tributylstannyl)oxazole (174 mg, 0.48 mmol) and tetrakis(triphenylphosphine)palladium (37 mg, 0.032 mmol) were added under nitrogen atmosphere. The reaction system was reacted at 85 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature and filtered, and the filtrate was added with water (30 mL) and extracted with dichloromethane (50 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation

to give a crude product. The crude product was purified by preparative chromatography to give **compound 163** (78.8 mg, 54% yield) in the form of a white solid. MS $[M+H]^+ = 452.8$.

^1H NMR (400 MHz, MeOD) δ 8.47 (t, 1H), 8.11 (s, 1H), 7.85 – 7.75 (m, 2H), 7.42 (s, 1H), 7.39 (s, 1H), 6.93 (d, $J = 6.1$ Hz, 1H), 4.32 – 4.22 (m, 1H), 3.84 – 3.76 (m, 1H), 3.69 – 3.65 (m, 1H), 3.61 – 3.54 (m, 1H), 3.45 – 3.35 (m, 1H), 3.17 – 3.16 (m, 2H), 2.99 – 2.88 (m, 1H), 2.74 – 2.63 (m, 1H), 2.49 (s, 3H), 2.04 (d, 3H).

Example 164

Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(3-propionyl-1H-pyrazol-1-yl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

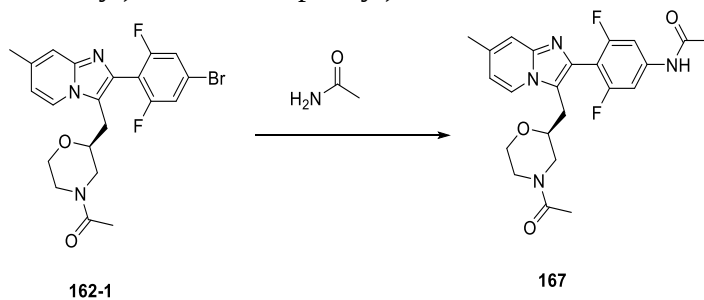


Intermediate 134-7 (50 mg, 0.12 mmol), 1-(1H-pyrazol-3-yl)propan-1-one (30 mg, 0.12 mmol) and potassium carbonate (30 mg, 0.36 mmol) were added to dimethyl sulfoxide (5 mL). The reaction system was stirred at 130 °C for 4 h under nitrogen atmosphere. The the reaction system was cooled to room temperature and filtered, and the filtrate was dispersed in ethyl acetate (15 mL), washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by preparative chromatography to give **compound 164** (5 mg, 7.5% yield) in the form of a yellow solid.

MS $[M+H]^+ = 523.8$. ^1H NMR (400 MHz, d_4 -MeOD) δ 8.51 (d, $J = 2.7$ Hz, 1H), 8.43 (d, $J = 6.9$ Hz, 1H), 7.80 (d, $J = 8.5$ Hz, 2H), 7.37 (s, 1H), 7.03 (d, $J = 2.6$ Hz, 1H), 6.89 (dd, $J = 7.0$, 1.5 Hz, 1H), 3.89 – 3.78 (m, 3H), 3.66 (s, 3H), 3.68 – 3.58 (m, 1H), 3.42 – 3.37 (m, 1H), 3.21 – 3.12 (m, 4H), 2.94 – 2.85 (m, 1H), 2.66 – 2.62 (m, 1H), 2.48 (s, 3H), 1.25 (t, $J = 7.3$ Hz, 3H).

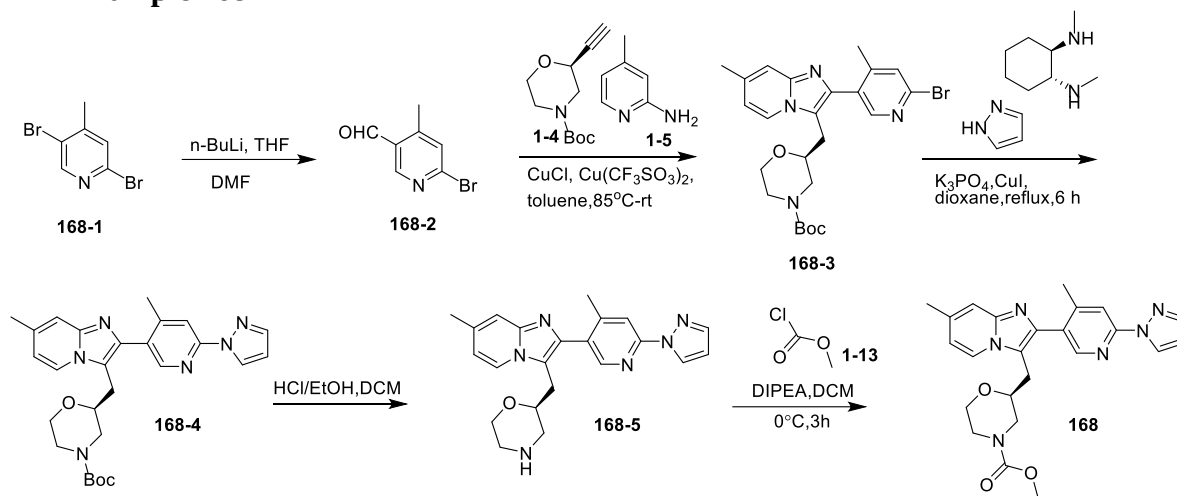
Example 167

Preparation of (S)-N-(4-(3-((4-acetylmorpholin-2-yl)methyl)-7-methylimidazo[1,2-*a*]pyridin-2-yl)-3,5-difluorophenyl)acetamide

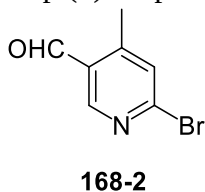


Intermediate 162-1 (240 mg, 0.518 mmol) and acetamide (45.9 mg, 0.778 mmol), cesium carbonate (150 mg, 0.778 mmol) and 4,5-bis(diphenylphosphino)-9,9-dimethylxanthene (6.00 mg, 0.010 mmol) were added to a reaction flask, and 1,4-dioxane was added. The reaction system was purged with N₂, added with tris(dibenzylideneacetone)dipalladium (9.50 mg, 0.010 mmol), purged with N₂ again, and heated to 80 °C and reacted for 4 h. After the reaction was completed as monitored by LCMS, the reaction system was added with water, and extracted with dichloromethane (30 mL × 2). The ester phase was dried over anhydrous sodium sulfate, and concentrated to remove dichloromethane. The residue was purified by preparative high performance liquid chromatography to give **compound 167** (102 mg, 52.4% yield) in the form of a white solid. LC-MS [M + H]⁺ = 443.1. ¹H NMR (400 MHz, DMSO-d₆) δ 10.40 (s, 1H), 8.42 – 8.35 (m, 1H), 7.41 (dd, *J* = 9.8, 1.7 Hz, 2H), 7.32 (s, 1H), 6.80 (dd, *J* = 9.6, 4.1 Hz, 1H), 4.07 (dd, *J* = 23.0, 13.0 Hz, 1H), 3.73 – 3.56 (m, 2H), 3.51 – 3.36 (m, 1H), 3.32 – 3.22 (m, 1H), 3.15 (td, *J* = 11.7, 2.6 Hz, 1H), 3.01 (dd, *J* = 5.8, 2.7 Hz, 2H), 2.77 (dd, *J* = 12.9, 10.5 Hz, 1H), 2.59 – 2.52 (m, 1H), 2.37 (s, 3H), 2.26 (dd, *J* = 12.6, 11.1 Hz, 1H), 2.10 (s, 3H), 1.91 (d, *J* = 12.7 Hz, 3H).

Example 168



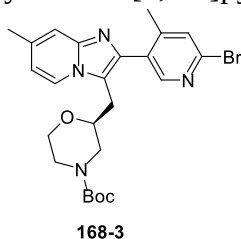
Step (1) Preparation of 6-bromo-4-methyl nicotinaldehyde



2,5-dibromo-4-methylpyridine (5.0 g, 17.19 mmol) was dissolved in tetrahydrofuran (60 mL) under nitrogen atmosphere. The reaction system was cooled to -78 °C, added with *n*-butyllithium (7.5 mL, 2.4 N), and stirred at this temperature for 30 min. Then the reaction system was added with *N,N*-dimethylformamide (2.5 g, 34.38 mmol), and reacted at -78 °C for 3 h. The

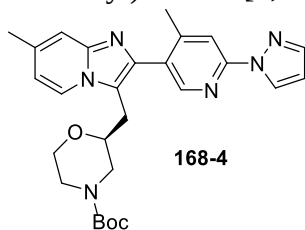
resulting reaction system was added with diluted hydrochloric acid (1 N, 20 mL) to quench the reaction, diluted with ethyl acetate (100 mL), washed with saturated sodium bicarbonate solution (80 mL) and saturated brine (80 mL) sequentially, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 168-2** (2.3 g, 54% yield) in the form of a white solid. LC-MS $[M+H]^+$: 199.8.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(6-bromo-4-methylpyridin-3-yl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 168-2 (320 mg, 1.50 mmol), *tert*-butyl (2*S*)-2-ethynylmorpholin-4-ylcarboxylate (317 mg, 1.50 mmol), 4-methylpyridin-2-amine (162 mg, 1.50 mmol), copper(I) chloride (45 mg, 0.45 mmol) and copper(II) trifluoromethanesulfonate (163 mg, 0.45 mmol) were added to anhydrous toluene (6 mL) under nitrogen atmosphere. The reaction system was heated to 85 °C, added dropwise with *N,N*-dimethylacetamide (0.1 mL), and stirred at 85 °C overnight. After the reaction was completed, the reaction system was cooled to room temperature, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 168-3** (324 mg, 41% yield) in the form of a brown powdery solid. MS $[M+H]^+$ = 500.7.

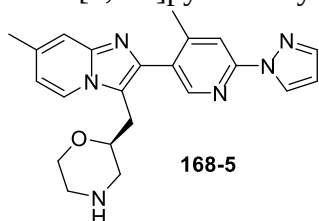
Step (3) Preparation of *tert*-butyl (S)-2-((7-methyl-2-(4-methyl-6-(1*H*-pyrazol-1-yl)pyridin-3-yl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 168-3 (320 mg, 0.64 mmol), 1*H*-pyrazole (130 mg, 1.92 mmol), copper(I) iodide (12 mg, 0.06 mmol), (1*R*,2*R*)-*N*₁,*N*₂-dimethylcyclohexane-1,2-diamine (18 mg, 0.13 mmol) and potassium phosphate (408 mg, 1.92 mmol) were added to dioxane (5 mL) under nitrogen atmosphere. The reaction system was reacted at 100 °C for 6 h. Then the reaction system was cooled to room temperature, added with water (15 mL) to quench the reaction, filtered, and extracted with ethyl acetate (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary

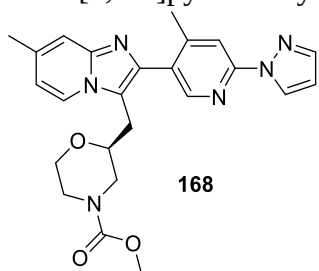
evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 168-4** (170 mg, 55% yield) in the form of a brownish-yellow oil. MS $[M+H]^+ = 488.9$.

Step (4) Preparation of (S)-2-((7-methyl-2-(4-methyl-6-(1H-pyrazol-1-yl)pyridin-3-yl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine



Intermediate 168-4 (170 g, 0.35 mmol) was dissolved in dichloromethane (5 mL), and a solution of HCl in ethanol (33%, 1.0 mL) was added. The reaction system was stirred at room temperature for 3 h. After the reaction was completed, the reaction system was concentrated by rotary evaporation to give **intermediate 168-5** (145 mg) in the form of a brown solid, MS $[M+H]^+ = 388.9$.

Step (5) Preparation of methyl (S)-2-((7-methyl-2-(4-methyl-6-(1H-pyrazol-1-yl)pyridin-3-yl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



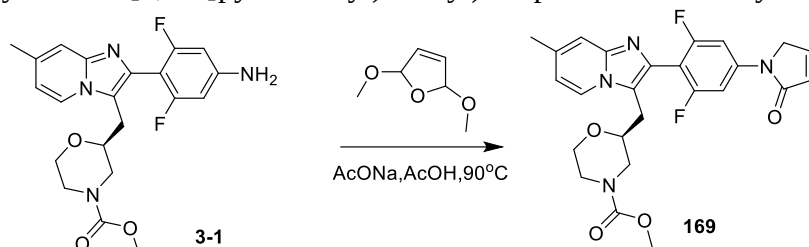
Intermediate 168-5 (135 mg, 0.35 mmol) was dissolved in anhydrous dichloromethane (5 mL) under nitrogen atmosphere, and diisopropylethylamine (225 mg, 1.75 mmol) was added at 0 °C, followed by the dropwise addition of a solution of methyl chloroformate in dichloromethane (10 mg/mL, 3.5 mL). The reaction system was stirred at 0 °C for 2 h. After the reaction was completed, the reaction system was added with water (20 mL), and extracted with dichloromethane (20 mL × 3). The organic phases were combined, washed with saturated brine (20 mL), dried over sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by silica gel column chromatography (dichloromethane:ethyl acetate = 2:1) to give a product. The product was purified by preparative by high performance liquid chromatography to give **compound 168** (45 mg, 21% yield) in the form of a white solid.

MS $[M+H]^+ = 446.8$. ^1H NMR (400 MHz, CDCl_3) δ 8.60 (d, $J = 2.2$ Hz, 1H), 8.33 (s, 1H), 8.09 (d, $J = 7.1$ Hz, 1H), 7.96 (s, 1H), 7.77 (s, 1H), 7.39 (s, 1H), 6.71 (d, $J = 7.1$ Hz, 1H), 6.51 –

6.47 (m, 1H), 4.00 – 3.75 (m, 3H), 3.67 (s, 3H), 3.62 – 3.52 (m, 1H), 3.45 – 3.34 (m, 1H), 3.08 – 2.98 (m, 2H), 2.96 – 2.86 (m, 1H), 2.63 – 2.52 (m, 1H), 2.44 (s, 3H), 2.41 (s, 3H).

Example 169

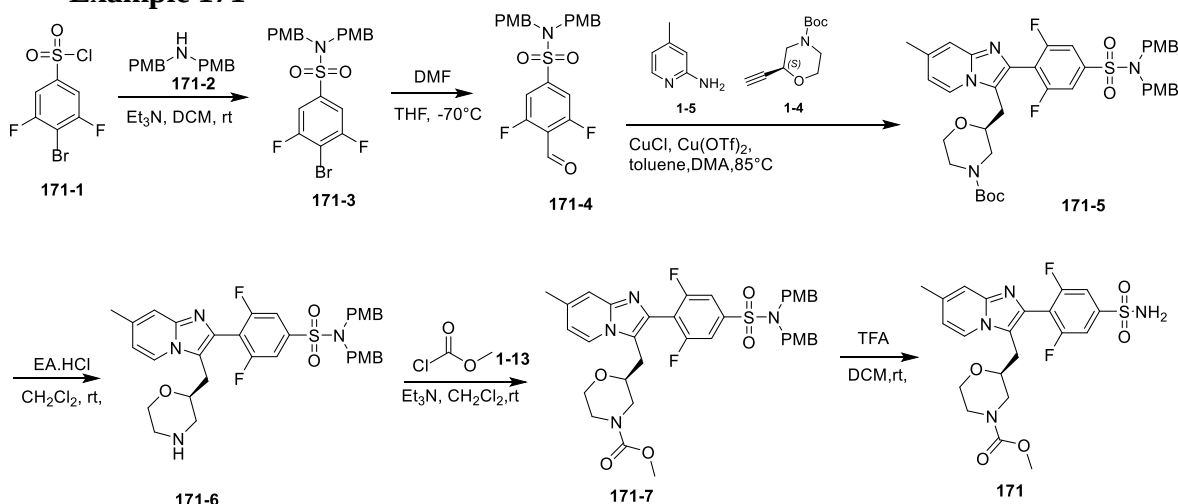
Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(2-carbonyl-2,5-dihydro-1H-pyrrol-1-yl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



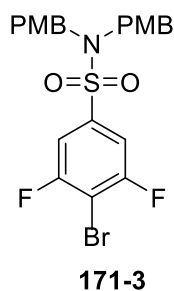
Intermediate 3-1 (100 mg, 0.24 mmol) was dissolved in acetic acid (1 mL) under nitrogen atmosphere, and 2,5-dimethoxy-2,5-dihydrofuran (35 mg, 0.26 mmol) and sodium acetate (50 mg, 0.48 mmol) were added. The reaction system was reacted at 90 °C for 2 h. After the reaction was completed, the reaction system was cooled to room temperature, filtered and concentrated by rotary evaporation. The crude product was separated by preparative by high performance liquid chromatography to give **compound 169** (20.2 mg, 17.4% yield) in the form of a white solid. MS $[M+H]^+ = 482.8$.

^1H NMR (400 MHz, MeOD) δ 8.42 (d, $J = 7.0$ Hz, 1H), 7.70 (d, $J = 10.1$ Hz, 2H), 7.50 (dt, $J = 6.1, 1.9$ Hz, 1H), 7.35 (s, 1H), 6.89 (dd, $J = 7.1, 1.4$ Hz, 1H), 6.29 (dt, $J = 4.3, 1.9$ Hz, 1H), 4.66 (t, $J = 1.8$ Hz, 2H), 3.82 – 3.79 (m, 3H), 3.67 (s, 3H), 3.63 – 3.57 (m, 1H), 3.41 – 3.38 (m, 1H), 3.15 – 3.08 (m, 2H), 2.95 – 2.86 (m, 1H), 2.70 – 2.59 (m, 1H), 2.47 (s, 3H).

Example 171

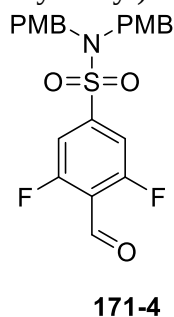


Step (1) Preparation of 4-bromo-3,5-difluoro-N,N-bis(4-methoxybenzyl)benzenesulfonamide



Bis(4-methoxybenzyl)amine (36.0 g, 140 mmol, **compound 171-2**) and Et₃N (21.2 g, 210 mmol) were dissolved in dichloromethane (1000 mL) sequentially. The reaction system was stirred in an ice bath, and added with 4-bromo-3,5-difluorobenzenesulfonyl chloride (40.5 g, 139 mmol, **compound 171-1**) in portions (an exothermic reaction) with the internal temperature maintained at less than 10 °C. After the addition of **compound 171-1**, the ice bath was removed and the reaction system was reacted at room temperature for 1.0 h. After **compound 171-1** was consumed as detected by TLC (PE/EA = 10/1), the DCM phase was washed with aqueous citric acid solution (400 mL × 2, 10% in wt.) and saturated brine (400 mL) sequentially, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation to give **intermediate 171-3** (68.3 g, 96% yield) in the form of a white solid. LC-MS: [M+H]⁺ = 511.1.

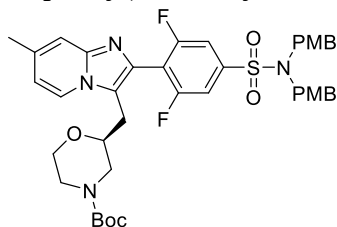
Step (2) Preparation of 3,5-difluoro-4-formyl-*N,N*-bis(4-methoxybenzyl)benzenesulfonamide



Intermediate 171-3 (68.3 g, 133.3 mmol) was dissolved in THF (800 mL). The reaction system was heated to 40 °C to dissolve the undissolved solid, purged with N₂ three times, and cooled to -70 °C below. Then the reaction system was added dropwise with *n*-butyllithium (2.5 M, 64 mL, 1.2 eq) slowly with the internal temperature maintained at less than -70 °C. After the completion of the dropwise addition of *n*-butyllithium, the reaction was reacted for 45 min, added with DMF (12.7 g, 1.3 eq), and reacted for another 2.0 h. The reaction system was added with hydrochloric acid (0.5 M) to quench the reaction and adjust the pH to 4, followed by liquid separation. The organic phase was concentrated by rotary evaporation. The aqueous phase was extracted with DCM (300 mL × 2). The residue after the rotary evaporation of THF was dissolved in the combined DCM phases, which was then washed with saturated brine, dried and concentrated by rotary evaporation to give a crude product (74 mg). The crude product was

purified by column chromatography to give **intermediate 171-4** (55 g, 90% yield, pure product) in the form of a white solid. LC-MS: $[M+H]^+ = 461.2$.

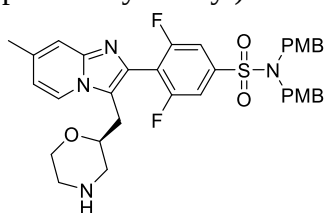
Step (3) Preparation of *tert*-butyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



171-5

To a 100 mL round-bottom flask were added **intermediate 171-4** (18.2 g, 39.4 mmol, 1.0 eq), 4-methylpyridin-2-amine (4.26 g, 39.4 mmol, 1.0 eq, **compound 1-5**), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (8.33 g, 39.4 mmol, 1.0 eq, **compound 1-4**), CuCl (1.17 g, 11.8 mmol, 0.3 eq), Cu(OTf)₂ (4.34 g, 11.8 mmol, 0.3 eq), toluene (200 mL) and DMA (12 mL) sequentially. The reaction system was purged with nitrogen 3 times, and heated in an oil bath at 85 °C overnight (12 h). The starting material (**compound 1-5**) was consumed, and **intermediate 171-4** was remained as detected by TLC. Then the reaction system was cooled to room temperature, added with aqueous ammonia (100 mL) and water (150 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with DCM (150 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 171-5** (11.2 g, 96% purity, 37% yield) in the form of a yellow solid. LC-MS: $[M+H]^+ = 761.9$.

Step (4) Preparation of (S)-3,5-difluoro-*N,N*-bis(4-methoxybenzyl)-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)benzenesulfonamide

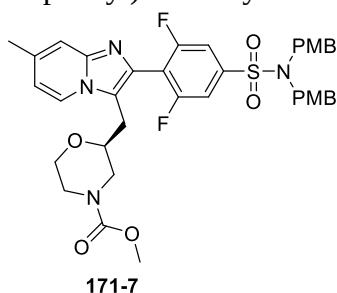


171-6

Intermediate 171-5 (11.2 g, 14.7 mol) was dissolved in DCM (33 mL), and a solution of HCl in dioxane (4 M, 33 mL) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (100 mL) and dichloromethane (200 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a DCM phase.

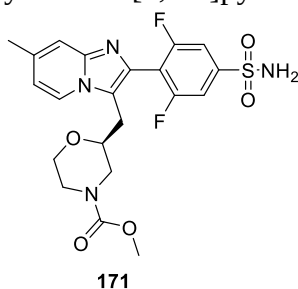
The aqueous phase was extracted with DCM (100 mL × 2). The DCM phases were combined, washed with saturated brine, dried and concentrated by rotary evaporation to give **intermediate 171-6** (9.7 g) in the form of a white solid, which was used directly in the next step. LC-MS: $[M+H]^+ = 622.2$.

- 5 Step (5) Preparation of methyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2,6-difluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



- Intermediate 171-6** (9.7 g, 14.7 mmol, 1.0 eq) was dissolved in DCM (40 mL), and Et₃N (2.96 g, 29.3 mmol, 2.0 eq) was added, followed by the dropwise addition of methyl chloroformate (2.08 g, 22 mmol, 1.5 eq). The reaction system was reacted for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (150 mL) and DCM (200 mL), and stirred for 10 min, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (50 mL × 2). The DCM phases were combined, washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 171-7** (9.4 g, 89% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 720.9$.
- 10
- 15

Step (6) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-sulfamoylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

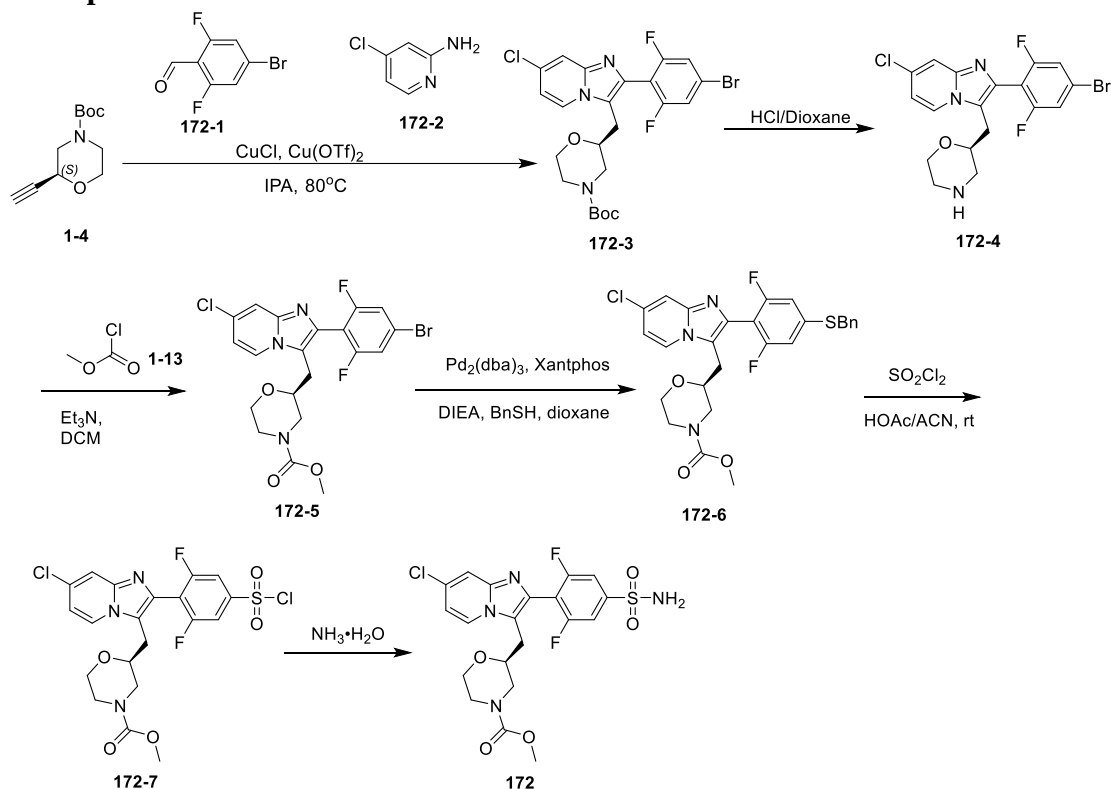


- Intermediate 171-7** (9.4 g) was dissolved in DCM (30 mL), and TFA (30 mL) was added. The reaction system was stirred at 35 °C overnight. A product (80% yield) was generated as detected by LCMS. Then the reaction system was supplemented with TFA (10 mL), and reacted at 40 °C for 4.0 h. The resulting reaction system was concentrated by rotary evaporation. The crude product was dissolved with acetonitrile and separated by preparative chromatography to
- 20

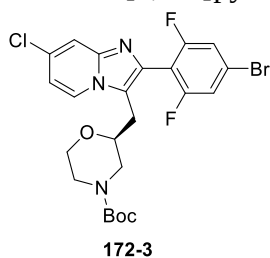
give **compound 171** (3.165 g, 50.6% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 481.1$.

^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 8.41 (d, $J = 7.11$ Hz, 1H), 7.69 (s, 2H), 7.59 (d, $J = 6.6$, 2H), 7.33 (s, 1H), 6.84 – 6.78 (m, 1H), 3.76 (d, $J = 12.5$ Hz, 1H), 3.60 (d, $J = 11.6$ Hz, 2H), 3.53 (s, 3H), 3.44 (d, $J = 3.6$ Hz, 1H), 3.31 (s, 1H), 3.24 – 3.14 (m, 1H), 3.02 (qd, $J = 15.7, 6.1$ Hz, 2H), 2.77 (s, 1H), 2.34 (s, 3H).

Example 172



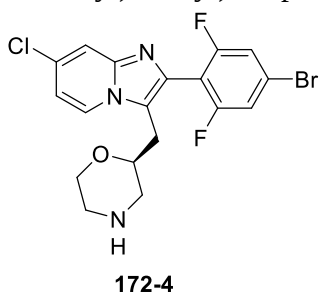
Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



To a 100 mL round-bottom flask were added *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (3.1 g, 1.0 eq, **intermediate 1-4**), 4-bromo-2,6-difluorobenzaldehyde (2.76 g, 1.0 eq, **compound 172-1**), 4-chloropyridin-2-amine (1.61 g, 1.0 eq, **compound 172-2**), CuCl (0.37 g, 0.3 eq), Cu(OTf)₂ (1.36 g, 0.3 eq) and isopropanol (50 mL) sequentially. The reaction system was purged with nitrogen 3 times, and reacted in an oil bath at 80 °C overnight. The starting material (**compound 172-2**) was consumed as detected by TLC. The reaction system was

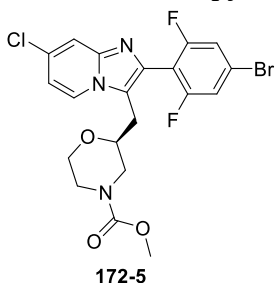
concentrated by rotary evaporation to remove isopropanol, and extracted with EA and aqueous ammonia sequentially. The EA phase was washed with saturated brine and citric acid sequentially, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 172-3** (3.0 g, 78% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 542.2$.

Step (4) Preparation of (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine



Intermediate 172-3 (2.67 g) was dissolved in dichloromethane (24 mL), and a solution of HCl in dioxane (24 mL) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was concentrated by rotary evaporation, added with water (15 mL) and dichloromethane (15 mL), and extracted to give an aqueous phase. The aqueous phase was adjusted to pH 8-9 (weak alkalinity) with aqueous sodium carbonate solution, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 ml $\times$ 2). The dichloromethane phases were combined, washed with saturated brine, and concentrated by rotary evaporation to give **intermediate 172-4** (1.70 g, 88.6% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 442.1$.

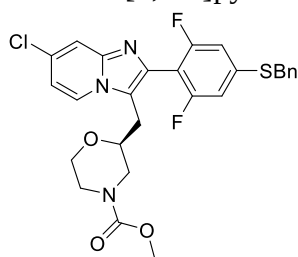
Step (5) Preparation of methyl (S)-2-((2-(4-bromo-2,6-difluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 172-4 (1.4 g, 1.0 eq) was dissolved in dichloromethane (10 mL), and triethylamine (480 mg, 1.5 eq) was added, followed by the dropwise addition of methyl chloroformate (388 mg, 1.3 eq). The reaction system was reacted for 1.0 h, and a product was generated as detected by LC-MS. After the reaction was completed, the reaction system was

added with water (10 mL), and stirred for 30 min, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 172-5** (1.01 g, 93.02% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 499.8$.

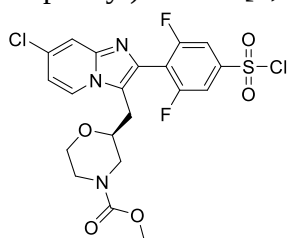
Step (6) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2,6-difluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



172-6

Intermediate 172-5 (0.73 g, 1.0 eq) was dissolved in dioxane (4 mL), and BnSH (0.24 g, 1.3 eq), $Pd_2(dba)_3$ (0.04 g, 0.03 eq), Xantphos (0.04 g, 0.05 eq) and DIEA (0.60 g, 3.0 eq) were added. The reaction system was purged with N_2 three times and reacted at 80 °C overnight. The starting material was consumed completely as monitored by LCMS. The reaction system was added with dichloromethane (10 mL) and water (10 mL), followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 172-6** (0.82 g, 91.53% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 544.2$.

Step (7) Preparation of methyl (S)-2-((7-chloro-2-(4-(chlorosulfonyl)-2,6-difluorophenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

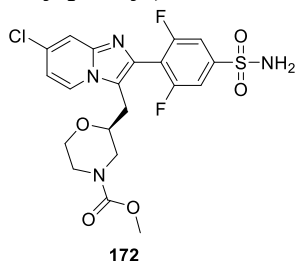


172-7

Intermediate 172-6 (510 mg) was added to a reaction flask, and dissolved with acetonitrile (3 mL), followed by the addition of glacial acetic acid (281 mg, 5.0 eq). The reaction system was added dropwise with SO_2Cl_2 (506 mg, 4.0 eq) in an ice bath, and reacted at 0 °C for 1 h. The

starting material was consumed, and **intermediate 172-7** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

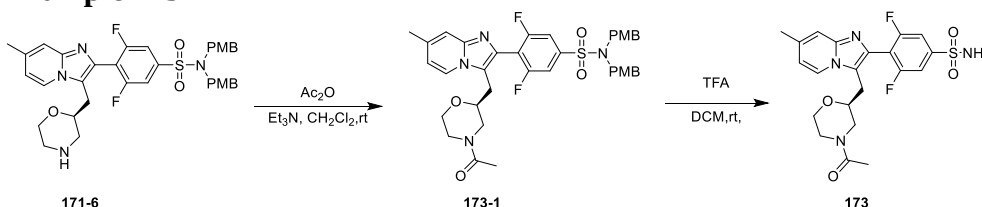
Step (8) Preparation of methyl (S)-2-((7-chloro-2-(2,6-difluoro-4-sulfamoylphenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



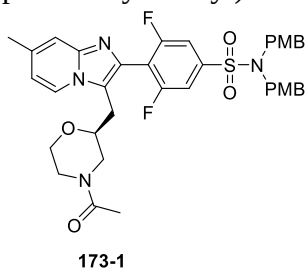
Aqueous ammonia (2 mL) was diluted with acetonitrile (1 mL) and added dropwise to the above reaction system at 0 °C. The reaction system was reacted at room temperature for 0.5 h. The starting material was consumed completely, and a target product was generated as detected by LCMS. The reaction system was extracted with water and ethyl acetate twice, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by preparative chromatography to give **compound 172** (185 mg, 99.74% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 501.1$.

^1H NMR (400 MHz, DMSO- d_6) δ = 8.11 (d, $J=7.4$, 1H), 7.29 (d, $J=1.6$, 1H), 7.22 (s, 2H), 7.14 (d, $J=6.6$, 2H), 6.60 (dd, $J=7.4$, 2.1, 1H), 3.33 (d, $J=12.8$, 1H), 3.13 (d, $J=11.3$, 2H), 3.07 (s, 3H), 2.97 (d, $J=7.8$, 1H), 2.77 – 2.69 (m, 1H), 2.69 – 2.61 (m, 1H), 2.53 (dd, $J=15.5$, 8.3, 1H).

Example 173



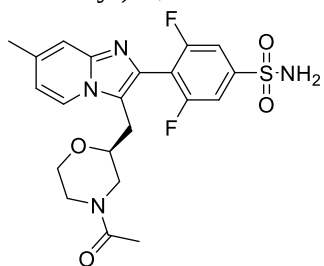
Step (1) Preparation of (S)-3,5-difluoro-N,N-bis(4-methoxybenzyl)-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-a]pyridin-2-yl)benzenesulfonamide



Intermediate 171-6 was prepared according to the steps in Example 1. **Intermediate 171-6** (620 mg, 0.94 mmol, 1.0 eq) was dissolved in DCM (5 mL), and Et_3N (236 mg, 2.34 mmol, 2.5 eq) was added, followed by the dropwise addition of acetic anhydride (191 mg, 1.88 mmol,

2 eq). The reaction system was reacted for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (30 mL) and DCM (50 mL), and stirred for 10 min, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (20 mL $\times$ 2). The DCM phases were combined, washed with saturated sodium chloride solution, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 173-1** (450 mg, 68.3% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 705.2$.

Step (2) Preparation of (S)-4-(3-((4-acetylmorpholin-2-yl)methyl)-7-methylimidazo[1,2-a]pyridin-2-yl)-3,5-difluorobenzenesulfonamide

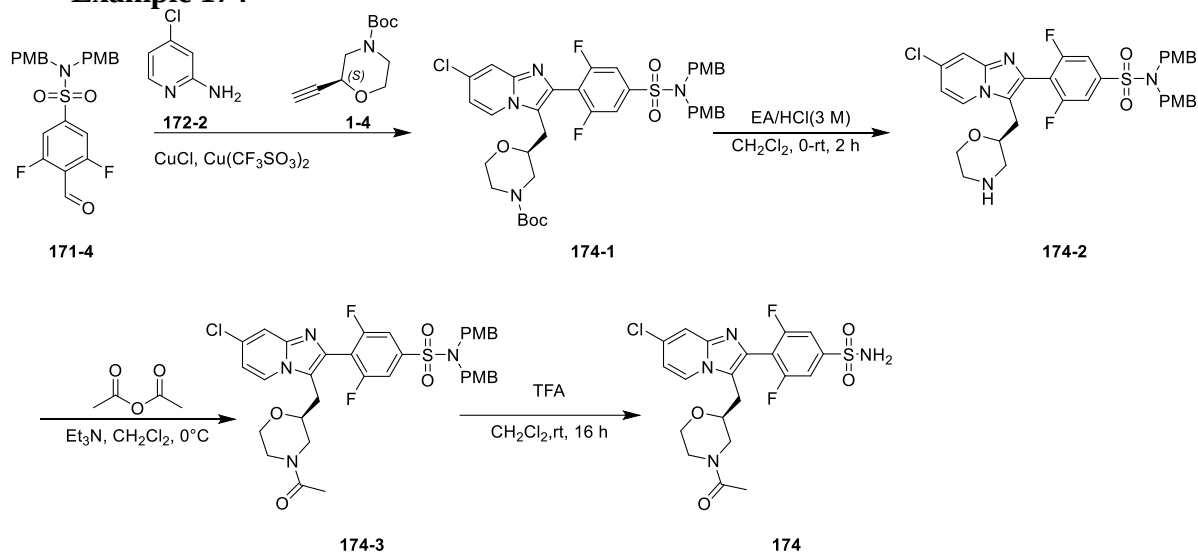


173

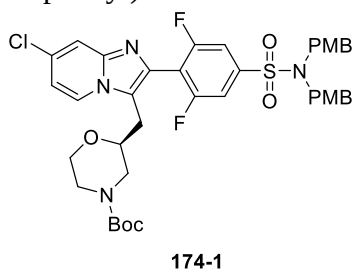
Intermediate 173-1 (420 mg) was dissolved in DCM (2 mL), and TFA (2 mL) was added. The reaction system was stirred at room temperature overnight. Then the reaction system was heated to 35 °C and reacted for 4.0 h, and the work-up was performed. The reaction system was added with water (20 mL) and DCM (20 mL). The aqueous phase was adjusted to pH 9 with sodium carbonate solution, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (20 mL $\times$ 2). The DCM phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give a product, which was then slurried with DCM/PE (1:10) to give **compound 173** (104 mg, 37.5% yield) in the form of a white solid.

^1H NMR (400 MHz, CDCl_3) δ 8.23 (1 H, dd, J 36.9, 6.8), 7.60 (1 H, s), 7.53 (2 H, t, J 6.0), 6.80 (2 H, t, J 12.6), 6.57 (1 H, s), 4.37 (1 H, t, J 11.4), 3.91 – 3.72 (1 H, m), 3.51 (2 H, d, J 12.1), 3.35 (1 H, dd, J 21.8, 11.9), 3.16 (0.5 H, t, J 11.6), 3.09 – 2.78 (2 H, m), 2.66 (0.5 H, t, J 11.1), 2.47 (3 H, d, J 3.8), 2.44 – 2.32 (0.5 H, m), 2.04 (2 H, s), 1.98 (1 H, s), 1.82 (2 H, s).

Example 174



Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2,6-difluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



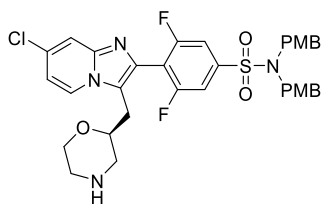
5

3,5-difluoro-4-formyl-*N,N*-bis(4-methoxybenzyl)benzenesulfonamide (29 g, 62.84 mmol), 4-chloropyridin-2-amine (8.08 g, 62.84 mmol) and *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (13.28 g, 62.84 mmol) were dissolved in toluene (150 mL) and dimethylacetamide (150 mL). The reaction system was stirred well, and into which copper(I) chloride (1.87 g, 18.85 mmol) and copper (II) trifluoromethanesulfonate (6.82 g, 18.85 mmol) were added, and reacted at 85 °C for 16 h under nitrogen atmosphere. Then the reaction system was added with water (750 mL), and extracted with ethyl acetate (150 mL × 2). The organic phase was washed with aqueous ammonia (100 mL × 1) and saturated brine (100 mL × 2) sequentially, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography (petroleum ether:ethyl acetate = 10:1-3:1) to give **intermediate 174-1** (18 g, 85% purity, 39% yield) in the form of a brownish-yellow foamy solid. LC-MS: $[M+H]^+ = 783.0$.

10

15

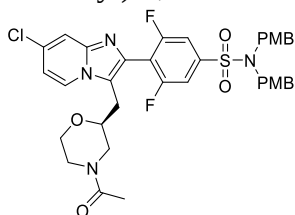
Step (2) Preparation of (S)-4-(7-chloro-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)-3,5-difluoro-*N,N*-bis(4-methoxybenzyl)benzenesulfonamide



174-2

Intermediate 174-1 (17 g, 21.7 mmol) was dissolved in dichloromethane (60 mL), and a solution of HCl in ethyl acetate (60 mL, 3 M) was added dropwise with the temperature maintained at 0 °C. After the completion of the dropwise addition, the reaction system was reacted at room temperature for 2 h. The starting material was consumed completely as detected by TLC (PE/EA = 3/1, R_f = 0.1). The reaction system was concentrated under reduced pressure. The crude product was adjusted to pH 8 with aqueous sodium bicarbonate solution, extracted with ethyl acetate (150 mL × 2), dried over anhydrous sodium sulfate with stirring for 20 min, filtered and concentrated to give **intermediate 174-2** (13 g, 90% purity, 87% yield) in the form of a brown foamy solid, LC-MS (M+H)⁺ = 683.0.

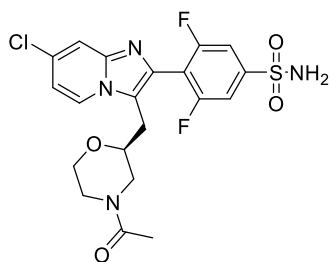
Step (3) Preparation of (S)-4-(3-((4-acetylmorpholin-2-yl)methyl)-7-chloroimidazo[1,2-a]pyridin-2-yl)-3,5-difluoro-N,N-bis(4-methoxybenzyl)benzenesulfonamide



174-3

Intermediate 174-2 (1.5 g, 2.2 mmol) was dissolved in dichloromethane (15 mL), and triethylamine (444 mg, 4.39 mmol) was added at 0 °C, followed by the dropwise addition of acetic anhydride (336 g, 3.29 mmol). The reaction system was reacted at 0 °C for 1 h. Then the reaction system was added with saturated aqueous sodium chloride solution (40 mL), extracted with dichloromethane (50 mL × 2), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography (PE/EA = 1/4-1/2) to give **intermediate 174-3** (1.1 g, 69% yield) in the form of a yellowish-white solid, LC-MS: [M+H]⁺ = 725.1.

Step (4) Preparation of (S)-4-(3-((4-acetylmorpholin-2-yl)methyl)-7-chloroimidazo[1,2-a]pyridin-2-yl)-3,5-difluorobenzenesulfonamide

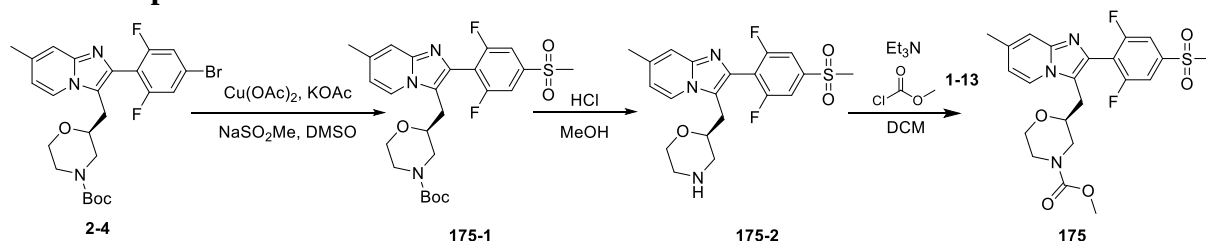


174

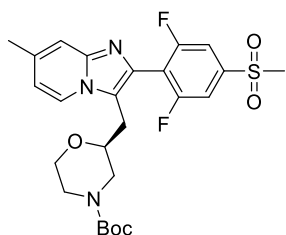
Intermediate 174-3 (1.1 g, 1.52 mmol) was dissolved in dichloromethane (15 mL), and trifluoroacetic acid (3 mL) was added at 0°C. The reaction system was reacted at room temperature for 16 h. The reaction system was added dropwise to an aqueous sodium bicarbonate solution to adjust the pH to 8, and extracted with dichloromethane (50 mL × 3). The organic phase was washed twice with saturated brine, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by preparative chromatography. The resulting acetonitrile, water, and trifluoroacetic acid phases were concentrated, adjusted to pH 8 with sodium bicarbonate solution, extracted with ethyl acetate (300 mL × 3), washed with saturated brine (100 mL × 2), dried over anhydrous sodium sulfate with stirring for 20 min, filtered and concentrated. The solid after concentration was washed three times with acetonitrile, added with acetonitrile (10 mL) and pure water (30 mL), and lyophilized to give **compound 174** (358 mg, 97% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 484.2$.

^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 8.62 (t, $J = 7.3$ Hz, 1H), 7.82 – 7.77 (m, 1H), 7.73 (d, $J = 2.3$ Hz, 2H), 7.64 (d, $J = 6.6$ Hz, 2H), 7.10 (td, $J = 7.3, 2.3$ Hz, 1H), 4.26 – 4.19 (m, 1H), 4.05 (d, $J = 13.3$ Hz, 1H), 3.78 (d, $J = 12.7$ Hz, 1H), 3.69 – 3.60 (m, 1H), 3.56 (d, $J = 13.3$ Hz, 1H), 3.52 – 3.37 (m, 1H), 3.25 (td, $J = 11.5, 2.3$ Hz, 1H), 3.20 – 2.97 (m, 3H), 2.80 (dd, $J = 12.9, 10.5$ Hz, 1H), 2.54 (dd, $J = 12.7, 2.9$ Hz, 1H), 2.29 (dd, $J = 12.8, 10.9$ Hz, 1H), 1.94 (d, $J = 4.2$ Hz, 3H).

Example 175



Step (1) Preparation of *tert*-butyl (S)-2-((2-(2,6-difluoro-4-(methanesulfonyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

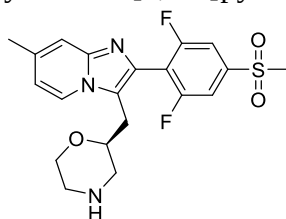


175-1

Intermediate 2-4 was prepared according to the steps in Example 2. **Intermediate 2-4** (90 mg), NaSO₂Me (22 mg), Cu(OAc)₂ (32 mg) and KOAc (34 mg) were added to a reaction flask containing DMSO (5 mL) sequentially. The reaction system was reacted at 100 °C overnight.

- 5 The starting material was consumed, and a new spot was generated as detected by TLC (PE/EtOAc = 1/1). The reaction system was added with EtOAc (10 mL), washed with NH₄Cl (10 mL × 3) and saturated brine (10 mL × 3) sequentially, dried over Na₂SO₄, and concentrated to give crude product. The crude product was purified by prep-TLC to give **intermediate 175-1** (28 mg) in the form of a white solid. LC-MS: [M+H]⁺ = 522.1.

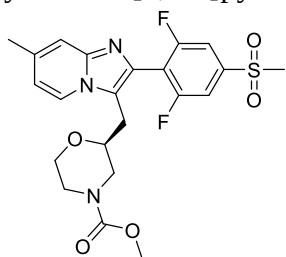
- 10 Step (2) Preparation of (S)-2-((2-(2,6-difluoro-4-(methylsulfonyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine



175-2

- 15 To **intermediate 175-1** (0.1 g) was added a solution of HCl in methanol (3 mL, 4 M) at room temperature. The reaction system was reacted at room temperature for 1 h. Then the reaction mixture was concentrated by rotary evaporation to give **intermediate 175-2** (80 mg) in the form of a yellow solid, which was used directly in the next step without purification.

Step (3) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-(methylsulfonyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



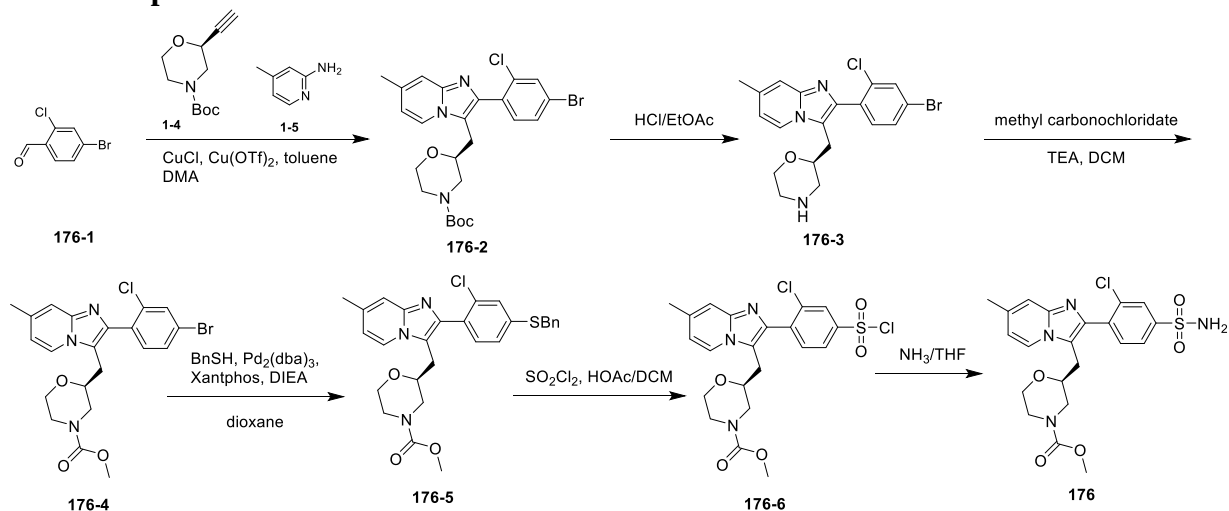
175

- 20 **Intermediate 175-2** (80 mg) was added into a reaction flask containing DCM (5 mL), and triethylamine (30 mg) was added. The reaction system was added with methyl chloroformate (27

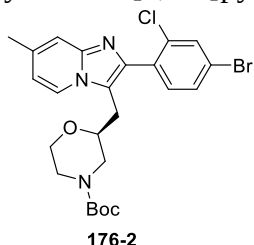
mg) at 0 °C and reacted at this temperature for 1 h. The starting material was consumed, and a target product was generated as detected by TLC. The product was purified by preparative chromatography to give **compound 175** (20 mg, 99.1% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 480.1$.

- 5 ^1H NMR (400 MHz, DMSO- d_6) δ 8.44 (d, $J = 7.2$ Hz, 1H), 7.81 (d, $J = 6.1$ Hz, 1H), 7.37 (s, 1H), 6.85 (d, $J = 6.9$ Hz, 1H), 3.77 (d, $J = 11.6$ Hz, 1H), 3.63 (d, $J = 11.9$ Hz, 2H), 3.55 (s, 2H), 3.46 (s, 1H), 3.39 (s, 2H), 3.28 – 3.16 (m, 1H), 3.13 – 2.97 (m, 1H), 2.38 (s, 2H).

Example 176



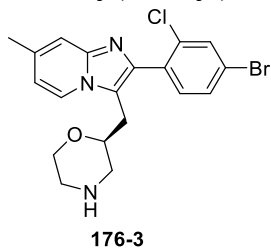
- 10 Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chlorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



- To a 100 mL round-bottom flask were added 4-bromo-2-chlorobenzaldehyde (2.0 g, 1 eq, **compound 176-1**), **compound 1-4** (1.9 g, 1 eq), **compound 1-5** (1.0 g, 1 eq), CuCl (0.27 g, 0.3 eq), Cu(OTf)₂ (0.99 g, 0.3 eq), toluene (20 mL) and DMA (2 mL) sequentially. The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material (**compound 176-1**) was consumed as detected by TLC. Then the reaction system was cooled to room temperature, added with aqueous ammonia (10 mL) and water (15 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with DCM (15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation to
- 15
- 20

give **intermediate 176-2** (1.46 g, 95% purity) in the form of a yellow oil. LC-MS: $[M+H]^+ = 520.2$.

Step (2) Preparation of (S)-2-((2-(4-bromo-2-chlorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine

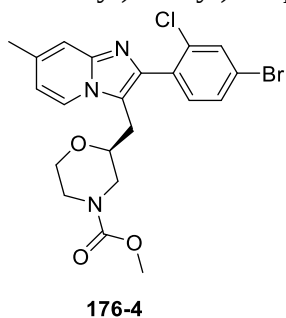


Intermediate 176-2 (1.46 g) was dissolved in methanol (5 mL), and a solution of HCl in EtOAc (4 M, 10 mL) was added. The reaction system was stirred at room temperature for 1 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (10 mL $\times$ 2). The DCM phases were combined, washed with saturated brine, dried and concentrated by rotary evaporation to give **intermediate 176-3** (1.05 g, 90% purity) in the form of a yellow oil. LC-MS: $[M+H]^+ = 420.1$.

10

Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2-chlorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

15

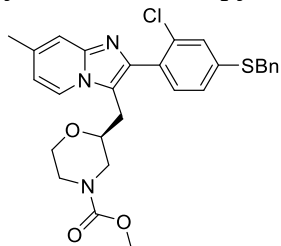


Intermediate 176-3 (1.05 g, 1.0 eq) was dissolved in DCM (10 mL), and DIPEA (0.65 g, 2.0 eq) was added, followed by the dropwise addition of methyl chloroacetate (0.27 g, 1.5 eq). The reaction system was reacted for 1 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and DCM (20 mL), and stirred for 10 min, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (10 mL $\times$ 2). The DCM phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 176-4** (0.88 g, 96% purity). LC-MS: $[M+H]^+ = 478.2$.

20

25

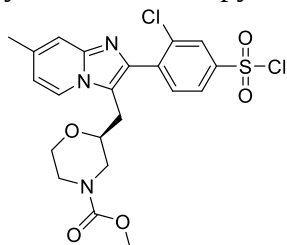
Step (4) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2-chlorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



176-5

Intermediate 176-4 (0.78 g, 1.0 eq) was dissolved in dioxane (10 mL), and BnSH (0.23 g, 1.1 eq), Pd₂(dba)₃ (0.15 g, 0.1 eq), Xantphos (0.1 g, 0.1 eq) and DIPEA (0.42 g, 2.0 eq) were added. The reaction system was purged with N₂ three times, and reacted at 85 °C overnight. The starting material was consumed completely as monitored by TLC. After the reaction was completed, the reaction system was added with water (10 mL) and EA (20 mL), and stirred for 10 min, followed by liquid separation to give an EtOAc phase. The aqueous phase was extracted with EtOAc (10 mL × 2). The EtOAc phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 176-5** (0.85 g, 92% purity) in the form of a yellow oil. LC-MS: [M+H]⁺ = 522.0.

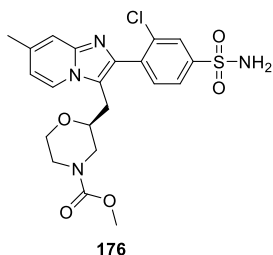
Step (5) Preparation of methyl (S)-2-((2-(2-chloro-4-(chlorosulfonyl)phenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



176-6

Intermediate 176-5 (150 mg) was added to a reaction flask containing DCM (1 mL), glacial acetic acid (86.3 mg, 5.0 eq) was added, and SO₂Cl₂ (154.9 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at room temperature for 2 h. The starting material was consumed, and **intermediate 176-6** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

Step (6) Preparation of methyl (S)-2-((2-(2-chloro-4-sulfamoylphenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



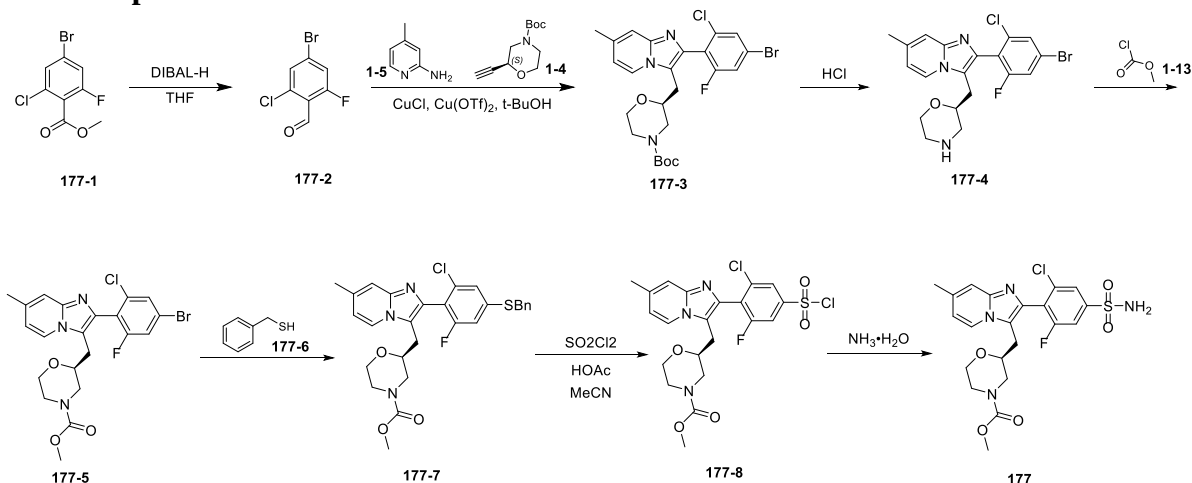
A solution of NH_3 in THF (0.4 M, 5 mL) was added dropwise slowly to the above reaction system at 0 °C. The reaction system was reacted at room temperature for 2 h. The starting material was consumed completely, and a target product was generated as detected by LCMS.

5 The reaction system was concentrated to dryness, and purified by preparative chromatography to give **compound 176** (18 mg) in the form of a white solid. LC-MS: $[\text{M}+\text{H}]^+ = 479.0$.

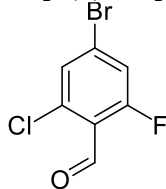
1H NMR (400 MHz, DMSO- d_6) δ 8.43 (d, $J = 7.1$ Hz, 1H), 7.96 (d, $J = 1.7$ Hz, 1H), 7.83 (dd, $J = 8.0, 1.8$ Hz, 1H), 7.71 (d, $J = 8.0$ Hz, 1H), 7.59 (s, 2H), 7.36 (s, 1H), 6.83 (dd, $J = 7.1, 1.6$ Hz, 1H), 3.79 – 3.59 (m, 3H), 3.56 (s, 3H), 3.51 (s, 1H), 3.23 (dd, $J = 11.8, 9.1$ Hz, 1H), 3.05 (qd, $J = 15.6, 6.3$ Hz, 2H), 2.82 (s, 1H), 2.51 (s, 1H), 2.38 (s, 3H).

10

Example 177



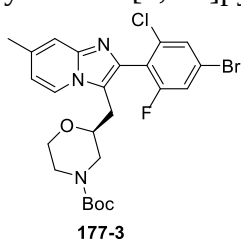
Step (1) Preparation of 4-bromo-2-chloro-6-fluorobenzaldehyde



15 To a 100 mL round-bottom flask were added methyl 4-bromo-2-chloro-6-fluorobenzoate (1.0 g, 1.0 eq, **compound 177-1**) and anhydrous tetrahydrofuran sequentially. The reaction system was cooled to -70 °C, then slowly added with DIBAL-H (3.8 mL, 1.2 eq) and reacted at this temperature for 2 h. The starting material was consumed completely and a new spot was

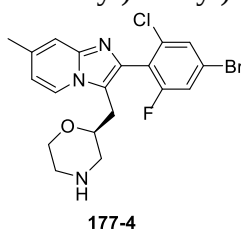
generated as detected by TLC. The reaction system was added with aqueous HCl solution (1 M) to quench the reaction, then stirred at room temperature for 20 min, and extracted with dichloromethane (30 mL × 3). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 177-2** (0.9 g, 80% purity) in the form of a colorless oil. LC-MS: $[M+H]^+ = 237.1$.

Step (2) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 177-2 (0.75 g, 1.0 eq), **compound 1-5** (0.67 g, 1.0 eq), **compound 1-4** (0.34 g, 1.0 eq), CuCl (0.19 g, 0.3 eq) and Cu(OTf)₂ (0.69 g, 0.3 eq) were added to toluene (20 mL) and DMA (2 mL). The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material **compound 1-4** was consumed as detected by TLC. Then the reaction system was cooled to room temperature, added with aqueous ammonia (10 mL) and water (15 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with dichloromethane (15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 177-3** (0.2 g, 99% purity) in the form of a brown foamy solid. LC-MS: $[M+H]^+ = 538.1$.

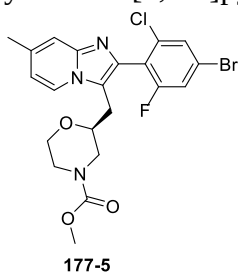
Step (3) Preparation of (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



Intermediate 177-3 (0.2 g) was dissolved in methanol (5 mL), and a solution of HCl in EA (10 mL) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a dichloromethane phase. The

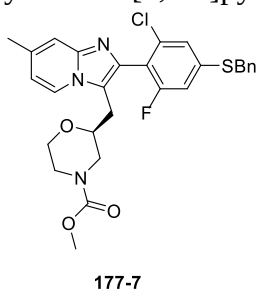
aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated brine, dried, filtered and concentrated by rotary evaporation to give **intermediate 177-4** (170 mg) in the form of a yellow solid. The crude product was used directly in the next step. LC-MS: $[M+H]^+ = 438.1$.

- 5 Step (4) Preparation of methyl (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



- Intermediate 177-4** (170 mg, 1.0 eq) was dissolved in dichloromethane (2 mL), and triethylamine (120 mg, 2.0 eq) was added, followed by the dropwise addition of methyl chloroacetate (55 mg, 1.5 eq). The reaction system was reacted for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL), and stirred for 10 min, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 177-5** (175 mg, 96% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 496.1$.

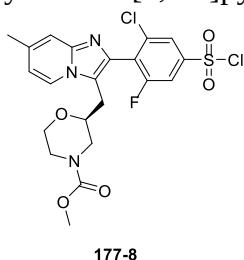
Step (5) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2-chloro-6-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



- Intermediate 177-5** (175 mg, 1.0 eq) was dissolved in dioxane (10 mL), and BnSH (53 mg, 1.1 eq), $Pd_2(dba)_3$ (36 mg, 0.1 eq), Xantphos (25 mg, 0.1 eq) and DIPEA (95 mg, 2.0 eq) were added. The reaction system was purged with N_2 three times and reacted at 85 °C overnight. The starting material was consumed completely as monitored by TLC. After the reaction was completed, the reaction system was added with water (10 mL) and ethyl acetate (20 mL), and stirred for 10 min, followed by liquid separation to give an ethyl acetate phase. The aqueous

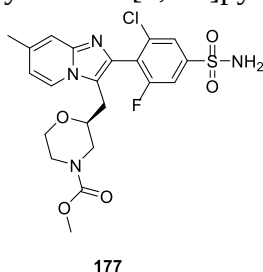
phase was extracted with ethyl acetate (10 mL × 2). The ethyl acetate phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 177-7** (165 mg, 95% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 540.0$.

- 5 Step (6) Preparation of methyl (S)-2-((2-(2-chloro-4-(chlorosulfonyl)-6-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



- Intermediate 177-7** (160 mg) was added to a reaction flask containing dichloromethane (5 mL), glacial acetic acid (90 mg, 5.0 eq) was added, and SO₂Cl₂ (160 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at 0 °C for 1 h. The starting material was consumed, and **intermediate 177-8** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

Step (7) Preparation of methyl (S)-2-((2-(2-chloro-6-fluoro-4-sulfamoylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

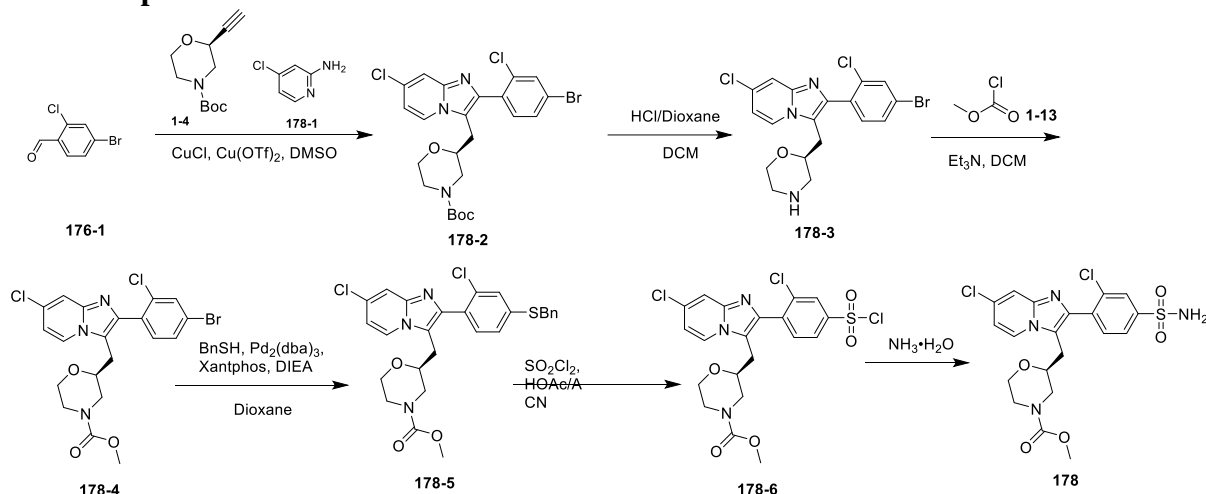


- 15 Aqueous ammonia (10 mL) was added dropwise slowly to the above reaction system at 0 °C. The reaction system was reacted at 0 °C for 0.2 h. The starting material was consumed completely, and a target product was generated as detected by LCMS. The reaction system was concentrated to dryness, and purified by preparative chromatography. to give **compound 177** (80 mg, 98.982% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 497.2$.

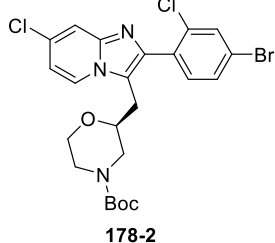
20 ¹H NMR (400 MHz, DMSO-*d*₆) δ 8.44 (d, *J* = 7.1 Hz, 1H), 7.85 (s, 1H), 7.72 (s, 2H), 7.70 (s, 1H), 7.37 (s, 1H), 6.86 (d, *J* = 7.1 Hz, 1H), 3.77 (d, *J* = 11.6 Hz, 1H), 3.63 (d, *J* = 11.9 Hz, 2H), 3.55 (s, 3H), 3.43 (s, 1H), 3.19 (dt, *J* = 11.8, 5.8 Hz, 1H), 3.01 (qd, *J* = 15.5, 6.0 Hz, 2H), 2.78 (s, 1H), 2.52 (s, 1H), 2.38 (s, 3H).

25

Example 178



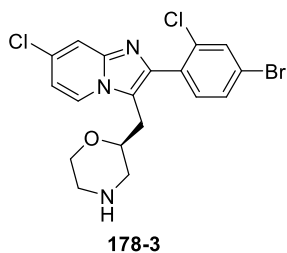
Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chlorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



5

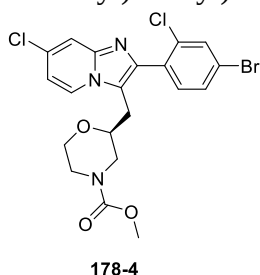
To a 100 mL round-bottom flask were added **compound 1-4** (680 mg, 1.0 eq), **compound 176-1** (706 mg, 1.0 eq), 4-chloropyridin-2-amine (413 mg, 1.0 eq, **compound 178-1**), CuI (61 mg, 0.1 eq), Cu(OTf)₂ (116 mg, 0.1 eq) and DMSO (7 mL) sequentially. The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material **compound 1-4** was consumed as detected by TLC. After the reaction system was cooled to room temperature, aqueous ammonium chloride solution (10 mL) and ethyl acetate (15 mL) were added. The reaction system was shaken for 5 min, followed by liquid separation to give an organic phase. The aqueous phase was extracted with ethyl acetate (15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 178-2** (0.65 g) in the form of a yellow oil. LC-MS: [M+H]⁺ = 540.2.

Step (2) Preparation of (S)-2-((2-(4-bromo-2-chlorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



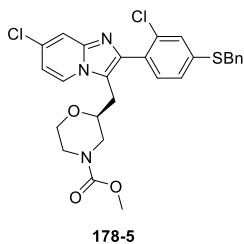
Intermediate 178-2 (0.65 g) was dissolved in dichloromethane (2 mL), and a solution of HCl in dioxane (6 mL) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, The reaction system was concentrated by rotary evaporation, added with water (10 mL) and dichloromethane (20 mL), and extracted to give an aqueous phase. The aqueous phase was adjusted to pH 8-9 (weak alkalinity) with aqueous sodium carbonate solution, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated brine, dried, filtered and concentrated by rotary evaporation to give **intermediate 178-3** (512 mg, 95% purity) in the form of a yellow oil. LC-MS: $[M+H]^+ = 440.1$.

Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2-chlorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



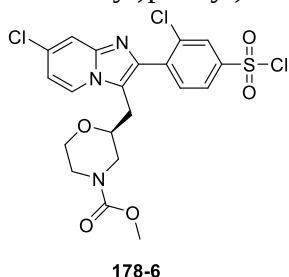
Intermediate 178-3 (500 mg, 1.0 eq) was dissolved in dichloromethane (5 mL), and triethylamine (206 mg, 2.0 eq) was added, followed by the dropwise addition of methyl chloroacetate (115 mg, 1.2 eq). The reaction system was reacted for 1.0 h and a product was generated as detected by LC-MS. After the reaction was completed, the reaction system was added with water (15 mL), and stirred for 30 min, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation to give **intermediate 178-4** (0.52 g) in the form of a yellow oil. LC-MS: $[M+H]^+ = 498.2$.

Step (4) Preparation of methyl (S)-2-((2-(4-benzylthio-2-chlorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



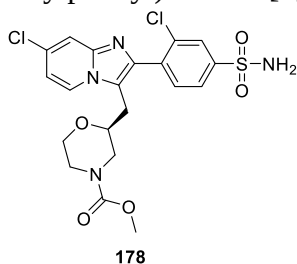
Intermediate 178-4 (0.52 g, 1.0 eq) was dissolved in dioxane (6 mL), and BnSH (0.16 g, 1.2 eq), Pd₂(dba)₃ (0.05 g, 0.05 eq), Xantphos (0.06 g, 0.1 eq) and DIEA (0.41 g, 3.0 eq) were added. The reaction system was purged with N₂ three times and reacted at 80 °C overnight. The starting material was consumed completely as monitored by LCMS. After the reaction was completed, the reaction system was added with water (10 mL) and ethyl acetate (20 mL), followed by liquid separation to give an ethyl acetate phase. The aqueous phase was extracted with ethyl acetate (10 mL × 2). The ethyl acetate phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 178-5** (0.48 g, 86% purity) in the form of a yellow oil. LC-MS: [M+H]⁺ = 542.2.

Step (5) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-4-(chlorosulfonyl)phenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 178-5 (220 mg) was added to a reaction flask of acetonitrile (3 mL), glacial acetic acid (121 mg, 5.0 eq) was added, and SO₂Cl₂ (218 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at 0 °C for 1 h. The starting material was consumed, and **intermediate 178-6** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

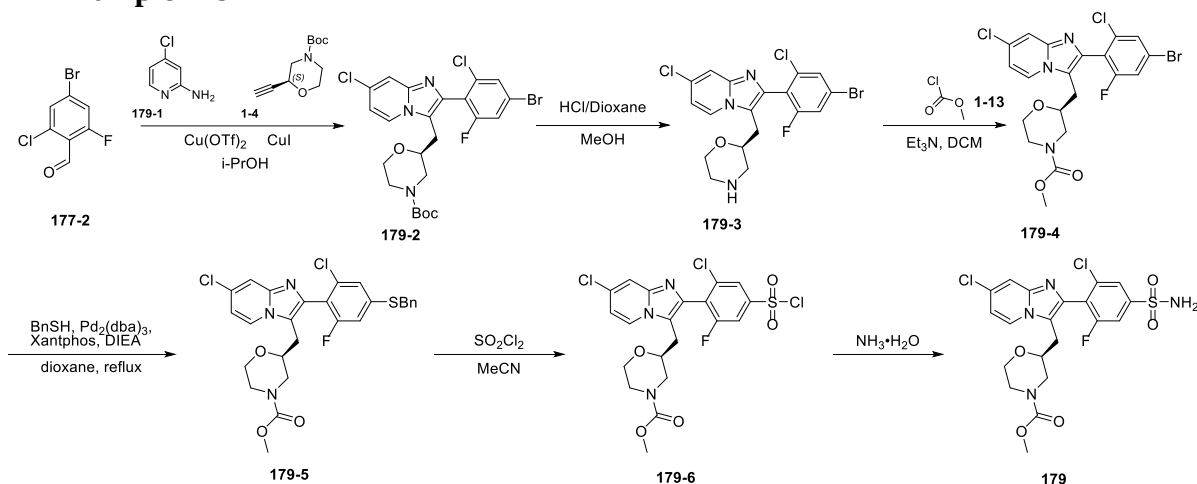
Step (6) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-4-sulfamoylphenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



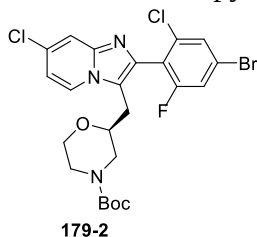
Aqueous ammonia (1.5 mL) was diluted with acetonitrile (1 mL) and added dropwise to the above reaction system at 0 °C. The reaction system was reacted at room temperature for 0.5 h. The starting material was consumed completely and a target product was generated as detected by LC-MS. The reaction system was extracted with water and ethyl acetate twice, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by preparative chromatography to give **compound 178** (30 mg, 99.6% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 499.1$.

^1H NMR (400 MHz, DMSO- d_6) δ 8.61 (d, $J = 7.4$ Hz, 1H), 7.98 (s, 1H), 7.85 (d, $J = 9.6$ Hz, 1H), 7.79 (d, $J = 2.1$ Hz, 1H), 7.73 (d, $J = 8.0$ Hz, 1H), 7.61 (s, 2H), 7.09 (dd, $J = 7.4, 2.1$ Hz, 1H), 3.79 (d, $J = 12.5$ Hz, 1H), 3.63 (d, $J = 11.5$ Hz, 3H), 3.56 (s, 4H), 3.51 (s, 1H), 3.26 – 3.21 (m, 2H), 3.20 – 3.14 (m, 1H), 3.12 (s, 1H), 3.03 (dd, $J = 15.6, 8.4$ Hz, 1H), 2.81 (s, 1H).

Example 179



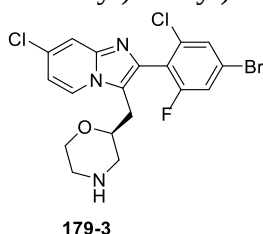
Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 177-2 (2.16 g, 1.0 eq), 4-chloropyridin-2-amine (1.17 g, 1.0 eq, **compound 179-1**), **compound 1-4** (1.92 g, 1.0 eq), CuI (0.19 g, 0.1 eq) and Cu(OTf)₂ (0.33 g, 0.1 eq) were added to isopropanol (25 mL). The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material **compound 1-4** was consumed as detected by TLC. work-up: the reaction system was cooled to room temperature, added with aqueous ammonia (10 mL) and water (15 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with dichloromethane

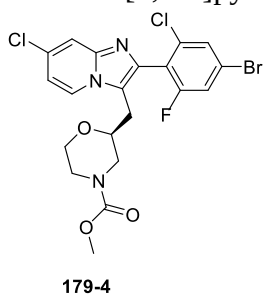
(15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 179-2** (0.93 g, 92% purity) in the form of a light yellow foamy solid. LC-MS: $[M+H]^+ = 558.0$.

5 Step (2) Preparation of (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



Intermediate 179-2 (0.93 g) was dissolved in methanol (10 mL), and a solution of HCl in dioxane (10 mL) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated brine, dried, and concentrated by rotary evaporation to give **intermediate 179-3** (0.78 g) in the form of a white solid, LC-MS: $[M+H]^+ = 458.1$. The crude product was used directly in the next step.

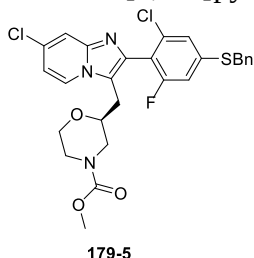
Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2-chloro-6-fluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 179-3 (690 mg, 1.0 eq) was dissolved in dichloromethane (10 mL), and triethylamine (380 mg, 2.0 eq) was added, followed by the dropwise addition of methyl chloroacetate (213 mg, 1.5 eq). The reaction system was reacted for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL), and stirred for 10 min, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride, dried over

anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 179-4** (472 mg, 96% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 515.9$.

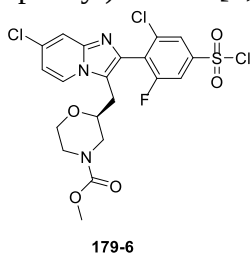
Step (4) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2-chloro-6-fluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 179-4 (420 mg, 1.0 eq) was dissolved in dioxane (10 mL), and BnSH (130 mg, 1.2 eq), $Pd_2(dba)_3$ (84 mg, 0.1 eq), Xantphos (50 mg, 0.1 eq) and DIPEA (210 mg, 2.0 eq) were added. The reaction system was purged with N_2 three times and reacted at 85 °C overnight.

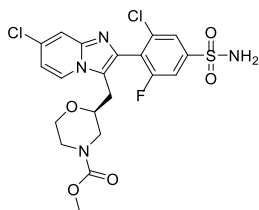
The starting material was consumed completely as monitored by TLC. After the reaction was completed, the reaction system was added with water (10 mL) and ethyl acetate (20 mL), and stirred for 10 min, followed by liquid separation to give an ethyl acetate phase. The aqueous phase was extracted with ethyl acetate (10 mL $\times$ 2). The ethyl acetate phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 179-5** (465 mg, 94% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 560.1$.

Step (5) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-4-(chlorosulfonyl)-6-fluorophenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 179-5 (415 mg) was added to a reaction flask of dichloromethane (5 mL), glacial acetic acid (224 mg, 5.0 eq) was added, and SO_2Cl_2 (415 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at 0 °C for 1 h. The starting material was consumed, and **intermediate 179-6** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

Step (6) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-6-fluoro-4-sulfamoylphenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

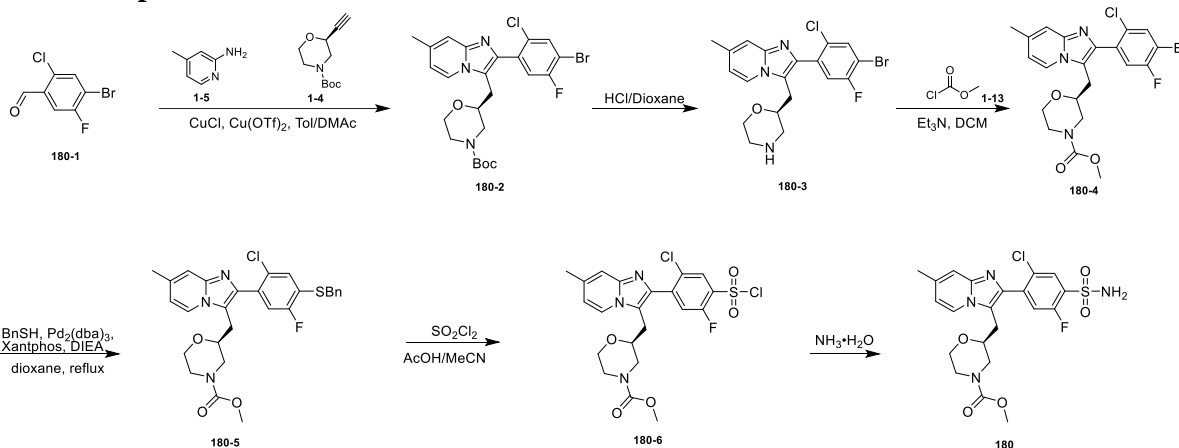


179

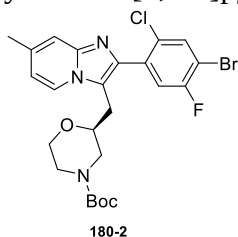
Aqueous ammonia (10 mL) was added dropwise slowly to the above reaction system at 0 °C. The reaction system was reacted at 0 °C for 0.2 h. The starting material was consumed completely, and a target product was generated as detected by LCMS. The reaction system was concentrated to dryness. The residue was purified by preparative chromatography to give **compound 179** (170 mg, 99.778 % purity) in the form of a white solid. LC-MS: $[M+H]^+ = 517.2$.

^1H NMR (400 MHz, DMSO- d_6) δ 8.61 (d, $J = 7.4$ Hz, 1H), 7.79 (dd, $J = 6.5, 4.4$ Hz, 2H), 7.64 (dt, $J = 8.0, 4.0$ Hz, 1H), 7.14 – 7.05 (m, 1H), 6.94 (s, 1H), 3.81 (d, $J = 11.7$ Hz, 1H), 3.63 (d, $J = 11.6$ Hz, 2H), 3.56 (s, 3H), 3.45 (d, $J = 3.5$ Hz, 1H), 3.20 (td, $J = 11.4, 2.2$ Hz, 1H), 3.04 (ddd, $J = 23.5, 15.5, 5.9$ Hz, 2H), 2.79 (s, 1H), 2.51 (s, 1H).

Example 180



Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

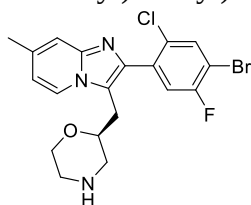


180-2

To a 100 mL round-bottom flask were added 4-bromo-2-chloro-5-fluorobenzaldehyde (1.5 g, 6.4 mmol, **compound 180-1**), **compound 1-4** (2.67 g, 12.6 mmol) and **compound 1-5** (0.68 g, 6.4 mmol) sequentially, and then toluene (15 mL) and *N,N*-dimethylacetamide (1 mL) were added, followed by the addition of catalysts copper(I) chloride (187.6 mg, 1.9 mmol) and copper(II) trifluoromethanesulfonate (685.4 mg, 1.9 mmol). The reaction system was purged

with nitrogen three times, and reacted at 85 °C for 4 h. After the reaction was completed, a target product was generated as detected by LC-MS. The reaction system was concentrated by rotary evaporation to remove the solvent, dissolved with dichloromethane (50 mL), added with aqueous ammonia (50 mL), and stirred, followed by liquid separation to give an organic phase. The aqueous phase was extracted with DCM (150 mL × 3). The organic phase was washed with saturated citric acid solution and saturated brine 3 times, respectively, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 180-2** (1.9 g, 59.9% yield) in the form of a light yellow foamy solid powder. LC-MS: $[M+H]^+ = 538.1$.

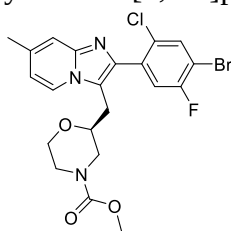
Step (2) Preparation of (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine



180-3

Intermediate 180-2 (0.7 g, 1.4 mmol) was dissolved in dichloromethane (5 mL). The reaction system was added with a solution of HCl in 1,4-dioxane solution (4 mL, 4.0 mol/L) in an ice bath, and then reacted at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was concentrated by rotary evaporation to remove the solvent, added with water (5 mL) and ethyl acetate (5 mL), and left to stand for liquid separation to give an aqueous phase. The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with saturated aqueous sodium bicarbonate solution, and then extracted with ethyl acetate (5 mL × 3). The organic phases were combined, washed with saturated saline 3 times, and concentrated to give **intermediate 180-3** (560 mg, 91.5% yield) in the form of a light yellow oily liquid, which was used directly in the next step. LC-MS: $[M+H]^+ = 438.1$.

Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

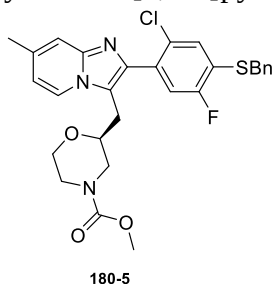


180-4

Intermediate 180-3 (560 mg, 1.28 mmol) was dissolved in dichloromethane (5 mL), and triethylamine (260 mg, 2.56 mmol) was added. The reaction system was added with methyl

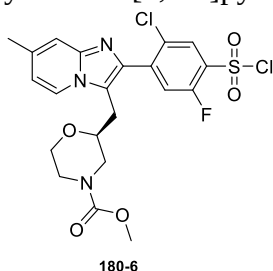
chloroformate (145.5 mg, 1.54 mmol) in an ice bath, reacted at room temperature for 0.5 h, and a product was generated as detected by LC-MS. After the reaction was completed, the reaction system was added with water (5 mL) to quench the reaction and left to stand for liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (5 mL × 2). The dichloromethane phases were combined, washed with saturated brine 3 times, dried over anhydrous sodium sulfate, and concentrated to give **intermediate 180-4** (520 mg, 81.9% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 496.1$.

Step (4) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2-chloro-5-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 180-4 (520 mg, 1.05 mmol) was dissolved in 1,4-dioxane (10 mL), and $Pd_2(dba)_3$ (28.8 mg, 0.0315 mmol), Xantphos (30.3 mg, 0.0525 mmol), DIEA (401.7 mg, 3.15 mmol) and BnSH (169.6 mg, 1.36 mmol) were added. The reaction system was purged with nitrogen 3 times and reacted at 80 °C overnight. The starting material was consumed completely as detected by LC-MS. After the reaction was completed, water (10 mL) and ethyl acetate (20 mL) were added. The reaction system was stirred and left to stand for liquid separation to give an ethyl acetate phase. The aqueous phase was extracted with ethyl acetate (10 mL × 2). The ethyl acetate phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by column chromatography to give **intermediate 180-5** (500 mg, 88.5% yield) in the form of a light yellow solid. LC-MS: $[M+H]^+ = 440.2$.

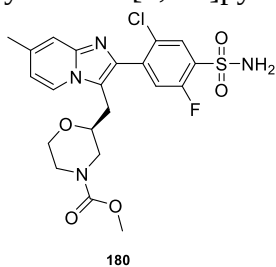
Step (5) Preparation of methyl (S)-2-((2-(2-chloro-4-(chlorosulfonyl)-5-fluorophenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Substrate **intermediate 180-5** (500 mg, 0.92 mmol) was added to a reaction flask of acetonitrile (15 mL), glacial acetic acid (276 mg, 4.6 mmol) was added, and sulfonyl chloride

(500 mg, 3.7 mmol) was added dropwise slowly in an ice bath. The reaction system was reacted at 0 °C for 1 h. The starting material was consumed and **intermediate 180-6** was generated as detected by LC-MS, LC-MS: $[M+H]^+ = 515.9$. The reaction system was used directly in the next step without treatment.

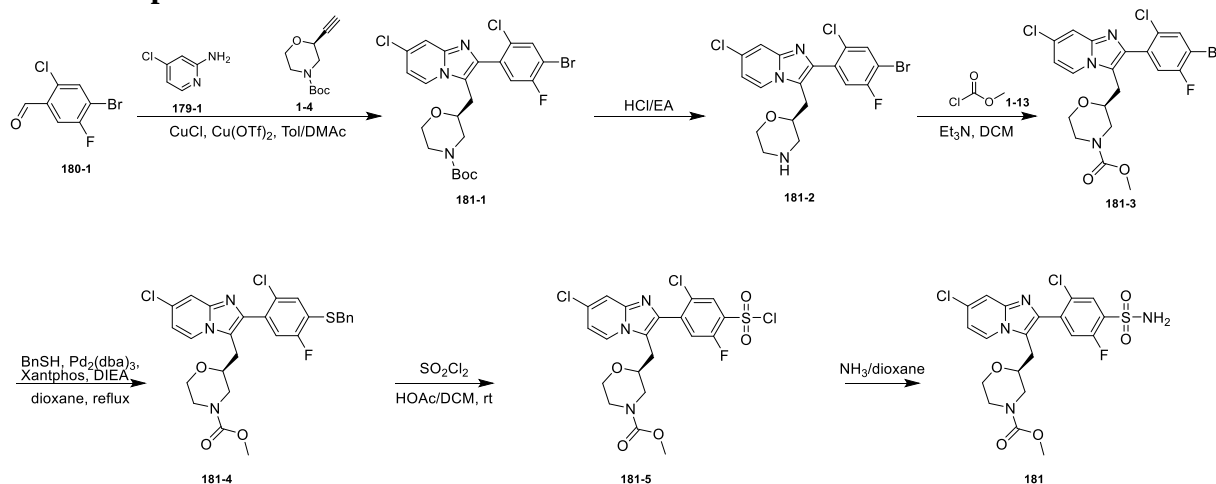
- 5 Step (6) Preparation of methyl (S)-2-((2-(2-chloro-5-fluoro-4-sulfamoylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



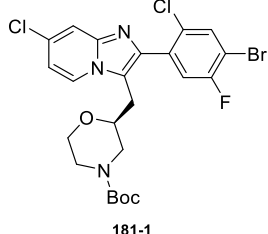
- 10 Aqueous ammonia (15 mL) was diluted with acetonitrile (10 mL) and added dropwise to the reaction system of the previous step at 0 °C. The reaction system was reacted at room temperature for 0.5 h. The starting material was consumed completely and a target product was generated as detected by LC-MS. The reaction system was left to stand for liquid separation to give an organic phase. The aqueous phase was extracted with ethyl acetate 2 times. The organic phase was washed with saline solution, dried over anhydrous sodium sulfate, and concentrated. The residue was purified by preparative chromatography, and separated by preparative TLC to give **compound 180** (140 mg, 99.5% purity) in the form of a white solid, LC-MS: $[M+H]^+ = 497.1$.

- 20 ^1H NMR (400 MHz, DMSO- d_6) δ : 8.42 (d, $J = 7.1$ Hz, 1H), 7.89 (s, 2H), 7.87 (d, $J = 6.6$ Hz, 1H), 7.62 (d, $J = 10.2$ Hz, 1H), 7.35 (s, 1H), 6.83 (dd, $J = 7.1, 1.6$ Hz, 1H), 3.84 – 3.73 (m, 1H), 3.64 (t, $J = 11.0$ Hz, 2H), 3.58 – 3.52 (m, 3H), 3.53 – 3.45 (m, 1H), 3.22 (td, $J = 11.7, 2.7$ Hz, 1H), 3.12 (dd, $J = 15.6, 4.1$ Hz, 1H), 3.00 (dd, $J = 15.6, 8.2$ Hz, 1H), 2.81 (s, 1H), 2.53 (d, $J = 7.6$ Hz, 1H), 2.42 – 2.29 (m, 3H).

Example 181



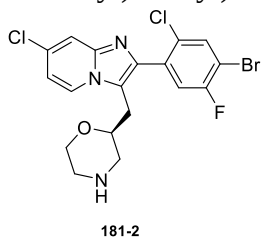
Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



5

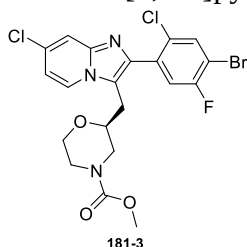
To a 100 mL round-bottom flask were added **intermediate 180-1** (1.5 g, 1.0 eq), **compound 179-1** (0.81 g, 1.0 eq), **compound 1-4** (1.3 g, 1.0 eq), CuCl (0.19 g, 0.3 eq), Cu(OTf)₂ (0.69 g, 0.3 eq), toluene (20 mL) and DMA (2 mL) sequentially. The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material **compound 1-4** was consumed as detected by TLC. Then the reaction system was cooled to room temperature, added with aqueous ammonia (10 mL) and water (15 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with dichloromethane (15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 181-1** (0.35 g, 99% purity) in the form of a light yellow foamy solid. LC-MS: [M+H]⁺ = 558.1.

Step (2) Preparation of (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-chloroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine



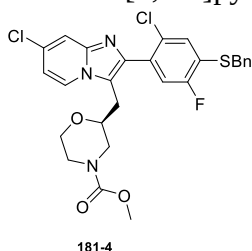
Intermediate 181-1 (0.1 g) was dissolved in methanol (5 mL), and HCl/EtOAc (10 mL, 3 M) was added. The reaction system was stirred at room temperature for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated brine, dried, and concentrated by rotary evaporation to give **intermediate 181-2** (80 mg) in the form of a white solid. The crude product was used directly in the next step. LC-MS: $[M+H]^+ = 458.1$.

Step (3) Preparation of methyl (S)-2-((2-(4-bromo-2-chloro-5-fluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 181-2 (80 mg, 1.0 eq) was dissolved in dichloromethane (2 mL), and triethylamine (36 mg, 2.0 eq) was added, followed by the dropwise addition of methyl chloroacetate (25 mg, 1.5 eq). The reaction system was reacted for 1.0 h. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL), and stirred for 10 min, followed by liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (10 mL × 2). The dichloromethane phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 181-3** (0.1 g, 96% purity) in the form of a white solid. LC-MS: $[M+H]^+ = 515.9$.

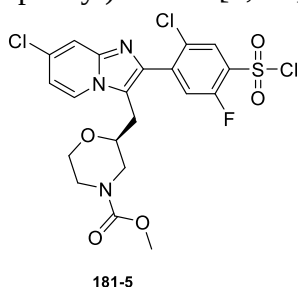
Step (4) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2-chloro-5-fluorophenyl)-7-chloroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 181-3 (0.45 g, 1.0 eq) was dissolved in dioxane (10 mL), and BnSH (0.13 g, 1.1 eq), $Pd_2(dba)_3$ (0.08 g, 0.1 eq), Xantphos (0.05 g, 0.1 eq) and DIPEA (0.23 g, 2.0 eq) were

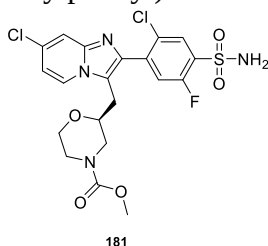
added. The reaction system was purged with N₂ three times and reacted at 85 °C overnight. The starting material was consumed completely as monitored by TLC. After the reaction was completed, the reaction system was added with water (10 mL) and ethyl acetate (20 mL), and stirred for 10 min, followed by liquid separation to give an ethyl acetate phase. The aqueous phase was extracted with ethyl acetate (10 mL × 2). The ethyl acetate phases were combined, washed with saturated sodium chloride, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 181-4** (0.3 g, 95% purity) in the form of a white solid. LC-MS: [M+H]⁺ = 560.2.

Step (5) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-4-(chlorosulfonyl)-5-fluorophenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 181-4 (280 mg) was added to a reaction flask of dichloromethane (5 mL), glacial acetic acid (150 mg, 5.0 eq) was added, and SO₂Cl₂ (270 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at room temperature for 1 h. The starting material was consumed, and **intermediate 181-5** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment.

Step (6) Preparation of methyl (S)-2-((7-chloro-2-(2-chloro-5-fluoro-4-sulfamoylphenyl)imidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate

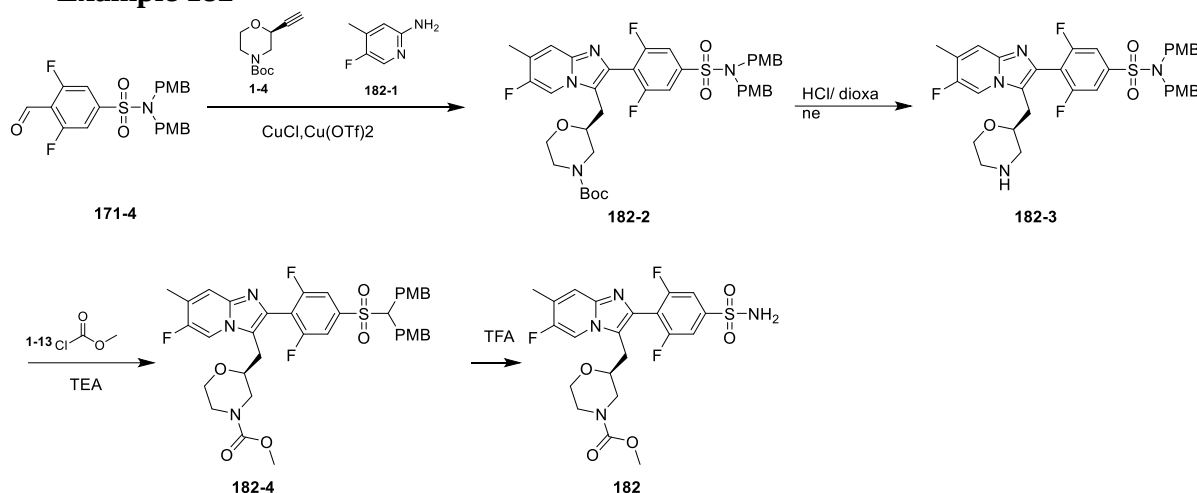


A solution of NH₃ in tetrahydrofuran (0.4 M, 5 mL) was added dropwise slowly to the above reaction system at 0 °C. The reaction system was reacted at room temperature for 2 h. The starting material was consumed completely, and a target product was generated as detected by LCMS. The reaction system was concentrated to dryness. The residue was purified by preparative chromatography to give **compound 181** (30 mg, 99% purity) in the form of a white solid. LC-MS: [M+H]⁺ = 517.1.

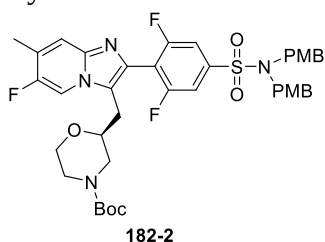
^1H NMR (400 MHz, $\text{DMSO}-d_6$) δ 8.62 (d, J = 7.4 Hz, 1H), 7.94 (s, 2H), 7.90 (d, J = 6.6 Hz, 1H), 7.81 – 7.79 (m, 1H), 7.67 (d, J = 10.1 Hz, 1H), 7.10 (dd, J = 7.4, 2.2 Hz, 1H), 3.84 (s, 1H), 3.65 (s, 2H), 3.57 (s, 3H), 3.52 (t, J = 9.4 Hz, 1H), 3.27 – 3.16 (m, 2H), 3.04 (dd, J = 15.6, 8.5 Hz, 1H), 2.82 (s, 1H), 2.52 (d, J = 1.9 Hz, 1H).

5

Example 182



Step (1) Preparation of *tert*-butyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2,6-difluorophenyl)-6-fluoro-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



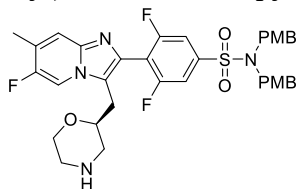
10

To a 100 mL round-bottom flask were added **intermediate 171-4** (1.5 g, 3.25 mmol), 5-fluoro-4-methylpyridin-2-amine (410 mg, 3.25 mmol) and **intermediate 1-4** (686.7 mg, 3.25 mmol), and then toluene (15 mL) and dimethyl acetamide (5 mL) were added, followed by the addition of catalysts copper(I) chloride (96.5 mg, 0.98 mmol) and copper(II) trifluoromethanesulfonate (353 mg, 0.98 mmol). The reaction system was purged with nitrogen three times and reacted at 80 °C overnight. After the reaction was completed, a target product was generated as detected by LC-MS. The reaction system was concentrated by rotary evaporation to remove the solvent, dissolved with dichloromethane (50 mL), added with aqueous ammonia (50 mL), and stirred, followed by liquid separation to give an organic phase. The aqueous phase was extracted with dichloromethane (150 mL $\times$ 3). The organic phase was washed with saturated citric acid solution and saturated brine 3 times, respectively, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column

20

chromatography to give **intermediate 182-2** (920 mg, 36.3% yield) in the form of a light yellow oily liquid.

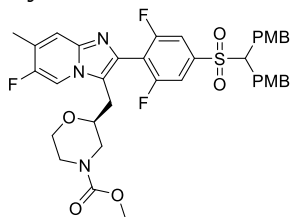
Step (2) Preparation of (S)-3,5-difluoro-4-(6-fluoro-7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)-*N,N*-bis(4-methoxybenzyl)benzenesulfonamide



182-3

Intermediate 182-2 (410 mg, 0.53 mmol) was dissolved in ethyl acetate (2 mL). The reaction system was added with a solution of HCl in 1,4-dioxane solution (2 mL, 4.0 mol/L) in an ice bath, and then reacted at room temperature for 1.0 h. The reaction was completed as detected by TLC. work-up: the reaction system was concentrated by rotary evaporation to remove the solvent, added with water (2 mL) and ethyl acetate (2 mL), and left to stand for liquid separation to give an aqueous phase. The aqueous phase was adjusted to pH 8-9 (weak alkalinity) with saturated aqueous sodium bicarbonate solution, and then extracted with ethyl acetate (3 mL × 3). The organic phases were combined, washed with saturated saline 3 times, and concentrated to give **intermediate 182-3** (330 mg, 92.5% yield) in the form of a light yellow oily liquid, which was used directly in the next step. LC-MS: $[M+H]^+ = 681.2$.

Step (3) Preparation of methyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2,6-difluorophenyl)-6-fluoro-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

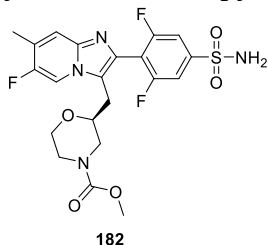


182-4

Intermediate 182-3 (330 mg, 0.49 mmol) was dissolved in dichloromethane (2 mL), and triethylamine (100 mg, 0.98 mmol) was added, followed by the dropwise addition of methyl chloroformate (70 mg, 0.73 mmol) in an ice bath. After the addition was completed, the reaction system was reacted at room temperature for 1 h. After the reaction was completed as detected by TLC, the reaction system was added with water (5 mL) to quench the reaction and left to stand for liquid separation to give a dichloromethane phase. The aqueous phase was extracted with dichloromethane (5 mL × 2). The dichloromethane phases were combined, washed with saturated brine 3 times, dried over anhydrous sodium sulfate, and concentrated to give

intermediate 182-4 (200 mg, 55.1% yield) in the form of a yellow oily liquid. LC-MS: $[M+H]^+ = 739.1$.

Step (4) Preparation of methyl (S)-2-((2-(2,6-difluoro-4-sulfamoylphenyl)-6-fluoro-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



5

Intermediate 182-4 (200 mg, 0.27 mmol) was dissolved in dichloromethane (2 mL), and trifluoroacetic acid (2 mL) was added. The reaction system was reacted at room temperature overnight. The starting material was consumed completely as detected by TLC, and then the work-up was performed. The reaction system was added with water (5 mL) to quench the reaction, then added with dichloromethane (3 mL), and left to stand for liquid separation to give an aqueous phase. The aqueous phase was adjusted to pH 8-9 (weak alkalinity) with saturated aqueous sodium bicarbonate solution, and then extracted with dichloromethane (5 mL $\times$ 3). The dichloromethane phases were combined, washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation to remove the solvent, and purified by preparative chromatography to give **compound 182** (82 mg, 97.8% purity, 60.98% yield) in the form of a white powdery solid. LC-MS: $[M+H]^+ = 499.1$.

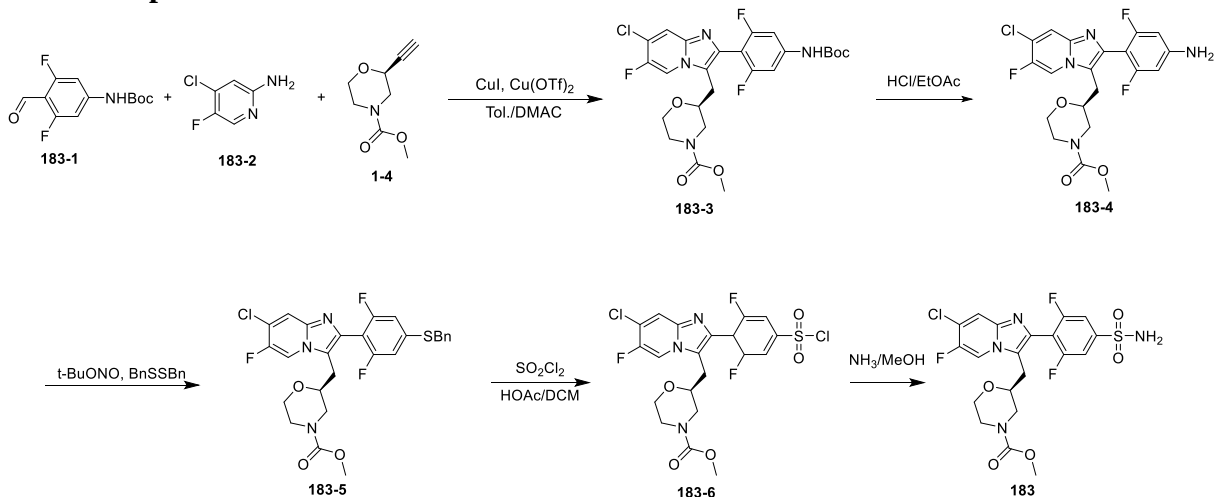
10

15

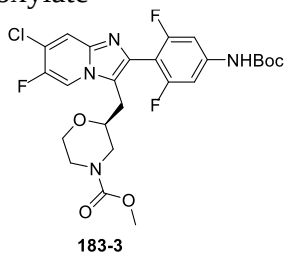
^1H NMR (400 MHz, DMSO- d_6) δ 8.71 (d, $J = 5.5$ Hz, 1H), 7.68 (s, 2H), 7.59 (d, $J = 6.6$ Hz, 2H), 7.50 (d, $J = 7.2$ Hz, 1H), 3.80 (d, $J = 12.3$ Hz, 1H), 3.59 (d, $J = 11.4$ Hz, 2H), 3.53 (s, 3H), 3.44 (s, 1H), 3.26 (s, 1H), 3.18 (dd, $J = 11.8, 9.1$ Hz, 1H), 3.06 (dd, $J = 15.4, 3.8$ Hz, 1H), 2.98 (d, $J = 8.4$ Hz, 1H), 2.76 (s, 1H), 2.32 (s, 3H).

20

Example 183

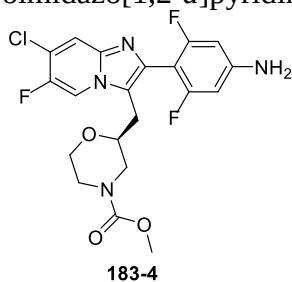


Step (1) Preparation of methyl (S)-2-((2-(4-((*tert*-butoxycarbonyl)amino)-2,6-difluorophenyl)-7-chloro-6-fluoroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



To a 10 mL round-bottom flask were added *tert*-butyl (3,5-difluoro-4-formylphenyl)carboxylate (0.75 g, 1.0 eq, **compound 183-1**), 4-chloro-5-fluoropyridin-2-amine (0.43 g, 1.0 eq, **compound 183-2**), **compound 1-4** (0.5 g, 1.0 eq), CuI (0.17 g, 0.3 eq), Cu(OTf)₂ (0.32 g, 0.3 eq), toluene (10 mL) and DMA (1 mL) sequentially. The reaction system was purged with nitrogen 3 times, heated in an oil bath at 85 °C and reacted overnight (12 h). The starting material **compound 1-4** was consumed as detected by TLC. The reaction system was cooled to room temperature, added with aqueous ammonia (5 mL) and water (15 mL), and stirred for 5 min, followed by liquid separation to give a toluene phase. The aqueous phase was extracted with DCM (15 mL × 2). The organic phase was washed with saturated brine, dried over anhydrous sodium sulfate, and concentrated by rotary evaporation. The residue was purified by column chromatography to give **intermediate 183-3** (0.18 g) in the form of a white solid. LC-MS: [M+H]⁺ = 555.0.

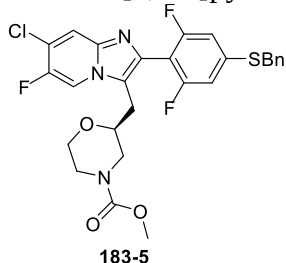
Step (2) Preparation of methyl (S)-2-((2-(4-amino-2,6-difluorophenyl)-7-chloro-6-fluoroimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 183-3 (0.18 g) was dissolved in methanol (2 mL), and HCl/EtOAc (3 M, 5 mL) was added. The reaction system was stirred at room temperature overnight. After the reaction was completed as detected by TLC, the reaction system was added with water (10 mL) and dichloromethane (20 mL). The aqueous phase was adjusted to pH 9-10 (weak alkalinity) with sodium carbonate, followed by liquid separation to give a DCM phase. The aqueous phase was extracted with DCM (10 mL × 2). The DCM phases were combined, washed with saturated

brine, dried, and concentrated by rotary evaporation to give **intermediate 183-4** (0.12 g) in the form of a white solid. LC-MS: $[M+H]^+ = 455.0$.

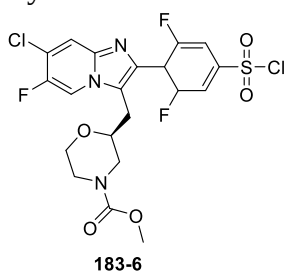
Step (3) Preparation of methyl (S)-2-((2-(4-(benzylthio)-2,6-difluorophenyl)-7-chloro-6-fluoroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



183-5

Intermediate 183-4 (0.2 g, 1.0 eq) and BnSSBn (0.11 g, 1.0 eq) were dissolved in MeCN (3 mL). The reaction system was purged with nitrogen three times. Ascorbic acid (40 mg, 0.5 eq) was dissolved with DMSO (0.5 mL) and added to the reaction system, which was then stirred and reacted for 10 min, and then added with *t*-BuONO (70 mg, 1.5 eq). The starting material was consumed completely as detected by TLC. The reaction system was added with water (2 mL), and then extracted with EtOAc (10 mL $\times$ 3). The EtOAc phase was washed with saturated brine, dried over Na₂SO₄, and concentrated. The residue was purified by column chromatography to give **intermediate 183-5** (0.11 g) in the form of a yellow solid. LC-MS: $[M+H]^+ = 562.1$.

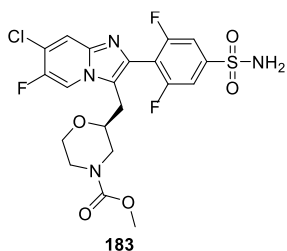
Step (4) Preparation of methyl (2S)-2-((7-chloro-2-(4-(chlorosulfonyl)-2,6-difluorocyclohexa-2,4-dien-1-yl)-6-fluoroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



183-6

Intermediate 183-5 (100 mg) was added to a reaction flask of DCM (1 mL), glacial acetic acid (53.4 mg, 5.0 eq) was added, and SO₂Cl₂ (96.1 mg, 4.0 eq) was added dropwise in an ice bath. The reaction system was reacted at room temperature for 2 h. The starting material was consumed, and **intermediate 183-6** was generated as detected by LCMS. The reaction system was used directly in the next step without treatment. LC-MS: $[M+H]^+ = 540.2$.

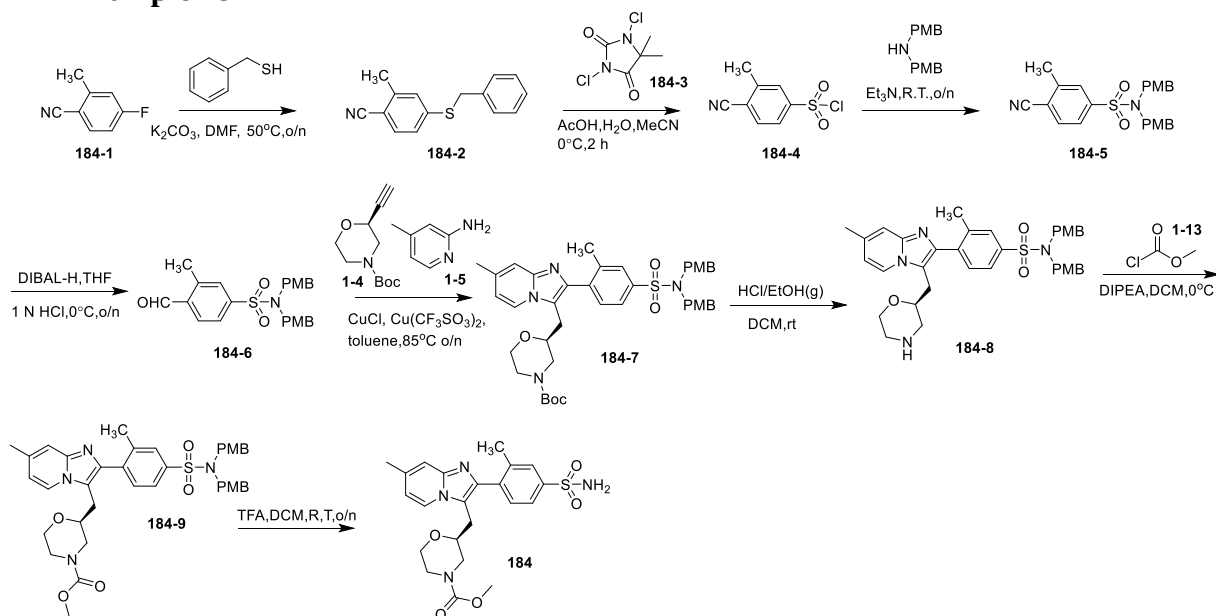
Step (5) Preparation of methyl (S)-2-((7-chloro-2-(2,6-difluoro-4-sulfamoylphenyl)-6-fluoroimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



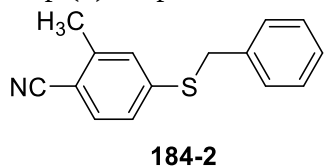
NH₃/MeOH (2 mL, 7 M) was added dropwise slowly to the above reaction system at 0 °C and reacted at room temperature for 2 h. The starting material was consumed completely and a target product was generated as detected by LC-MS. The reaction system was concentrated to dryness. The residue was purified by preparative chromatography to give **compound 183** (3.0 mg) in the form of a white solid. LC-MS: [M+H]⁺ = 519.1.

¹H NMR (400 MHz, DMSO-*d*₆) δ 9.05 (d, *J* = 5.3 Hz, 1H), 8.04 (d, *J* = 7.1 Hz, 1H), 7.73 (s, 2H), 7.64 (d, *J* = 6.7 Hz, 2H), 3.87 (d, *J* = 12.7 Hz, 1H), 3.61 (d, *J* = 8.1 Hz, 3H), 3.57 (s, 4H), 3.50 (d, *J* = 9.9 Hz, 2H), 3.23 – 3.13 (m, 3H), 3.05 – 2.80 (m, 1H).

Example 184



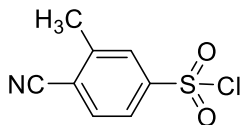
Step (1) Preparation of 4-(benzylthio)-2-methylbenzonitrile



4-fluoro-2-methylbenzonitrile (5.00 g, 37.00 mmol) was dissolved in *N,N*-dimethylformamide (30 mL), and potassium carbonate (7.50 g, 54.35 mmol) and benzylthiol (3.5 mL) were added. The reaction system was stirred at 50 °C overnight. Then the reaction system was cooled to room temperature, added with water (80 mL) to quench the reaction and extracted with ethyl acetate (30 mL × 3). The organic phases were combined, dried over anhydrous sodium

sulfate, filtered and concentrated to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 184-2** (4.88 g, 52.4% yield) in the form of a yellow solid, LC-MS: $[M + H]^+ = 239.8$.

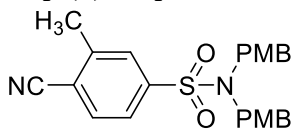
Step (2) Preparation of 4-cyano-3-methylbenzenesulfonyl chloride



184-4

Under nitrogen atmosphere, **intermediate 184-2** (4.88 g, 20.39 mmol) was dissolved in a prepared solvent (75 mL of acetonitrile, 3.5 mL of acetic acid, 1.5 mL of water). The reaction system was cooled to 0 °C, added with 1,3-dichloro-5,5-dimethylimidazolidine-2,4-dione (8.04 g, 40.78 mmol) in portions, and reacted for 2 h. Then the reaction system was added with water (100 mL) to quench the reaction and extracted with ethyl acetate (50 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 184-4** (2.8 g, 60.3% yield) in the form of a white solid. ^1H NMR (301 MHz, CDCl_3) δ 8.02 (s, 1H), 7.96 (d, $J = 8.2$, 1H), 7.87 (d, $J = 8.2$ Hz, 1H), 2.71 (s, 3H).

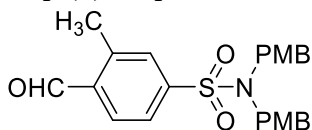
Step (3) Preparation of 4-cyano-*N,N*-bis(4-methoxybenzyl)-3-methylbenzenesulfonamide



184-5

Bis(4-methoxybenzyl)amine (2.80 g, 11.13 mmol) and triethylamine (3.40 g, 33.39 mmol) were dissolved in dichloromethane (50 mL). The reaction system was added with **intermediate 184-4** (2.40 g, 11.13 mmol) in an ice water bath, and stirred at room temperature overnight. Then the reaction system was added with water (50 mL) to quench the reaction and extracted with dichloromethane (50 mL × 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 184-5** (3.30 g, 64.9% yield) in the form of a light yellow solid. LC-MS $[M + H]^+ = 458.6$.

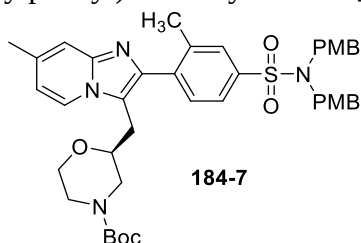
Step (4) Preparation of 4-formyl-*N,N*-bis(4-methoxybenzyl)-3-methylbenzenesulfonamide



184-6

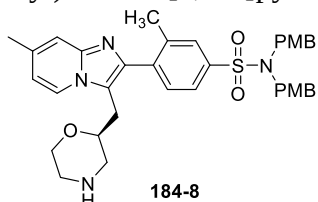
Intermediate 184-5 (3.00 g, 6.87 mmol) was dissolved in anhydrous tetrahydrofuran (60 mL). The reaction system was cooled to 0 °C, then added dropwise with a solution of diisobutylaluminum hydride in *n*-hexane (1 N, 17.2 mL), and stirred at 0 °C overnight. Then the reaction system was added with diluted hydrochloric acid (60 mL, 1 N) to quench the reaction and filtered. The filtrate was extracted with ethyl acetate (50 mL × 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 184-6** (1.68 g, 55.7% yield) in the form of a white solid. LC-MS $[M+H]^+ = 461.6$.

Step (5) Preparation of *tert*-butyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2-methylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Under nitrogen atmosphere, **intermediate 184-6** (500 mg, 1.14 mmol), 4-methylpyridin-2-amine (123 mg, 1.14 mmol), *tert*-butyl (S)-2-ethynylmorpholin-4-carboxylate (242 mg, 1.14 mmol), copper(I) chloride (35 mg, 0.34 mmol) and copper(II) trifluoromethanesulfonate (124 mg, 0.34 mmol) were dissolved in anhydrous toluene (5 mL). The reaction system was heated to 85 °C, then added with *N,N*-dimethylacetamide (0.1 mL), and stirred at 85 °C overnight. After the reaction was completed, the reaction system was filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 184-7** (380 mg, 42.7% yield) in the form of a white solid. LC-MS $[M+H]^+ = 740.5$.

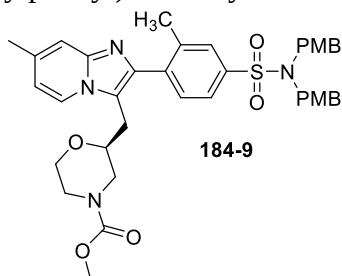
Step (6) Preparation of (S)-*N,N*-(4-methoxybenzyl)-3-methyl-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)benzenesulfonamide



Intermediate 184-7 (380 mg, 0.51 mmol) was dissolved in anhydrous dichloromethane (5 mL), and a solution of HCl in ethanol (1.5 mL, 4 N) was added. The reaction system was stirred at room temperature for 3 h. After the reaction was completed, the reaction system was

concentrated by rotary evaporation to give **intermediate 184-8** (450 mg, crude product) in the form of a white solid, LC-MS $[M + H]^+ = 640.5$.

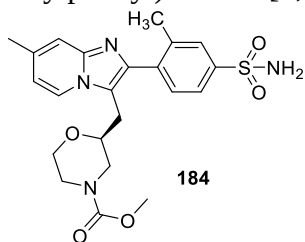
Step (7) Preparation of methyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2-methylphenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



184-9

Intermediate 184-8 (450 mg, crude) and *N,N*-diisopropylethylamine (325 mg, 2.53 mmol) were dissolved in anhydrous dichloromethane (5 mL) under nitrogen atmosphere. The reaction system was cooled in an ice water bath and added dropwise with a solution of methyl chloroformate in dichloromethane (10 mg/mL, 4.5 mL). After the completion of the dropwise addition, the reaction system was reacted at 0 °C for 2 h. After the reaction was completed, the reaction system was added with water (20 mL) and extracted with dichloromethane (15 mL $\times$ 3). The organic phases were combined, washed with saturated brine (20 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 184-9** (150 mg, 35.4% yield) in the form of a white solid, LC-MS $[M + H]^+ = 698.5$.

Step (9) Preparation of methyl (S)-2-((7-methyl-2-(2-methyl-4-sulfamoylphenyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate

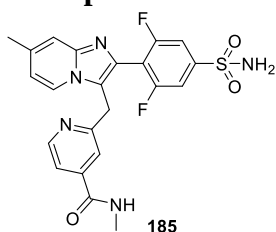


184

Intermediate 184-9 (150 mg, 0.21 mmol) was dissolved in anhydrous dichloromethane (1 mL), and trifluoroacetic acid (2 mL) was added. The reaction system was stirred overnight at room temperature. After the reaction was completed, the reaction system was directly concentrated by rotary evaporation and purified by preparative reverse phase chromatography to give **compound 184** (17 mg, 17.3% yield) in the form of a white solid. LC-MS $[M+H]^+ = 458.5$.

¹H NMR (400 MHz, CDCl₃) δ 8.16 (s, 1H), 7.87 (s, 1H), 7.77 (d, *J* = 7.7 Hz, 1H), 7.56 (s, 1H), 7.42 (d, *J* = 7.8 Hz, 1H), 6.80 (d, *J* = 7.0 Hz, 1H), 5.53 (brs, 2H), 3.98 – 3.76 (m, 3H), 3.69 (s, 3H), 3.58 – 3.47 (m, 1H), 3.43 – 3.31 (m, 1H), 3.05 – 2.85 (m, 3H), 2.60 – 2.50 (m, 1H), 2.49 (s, 3H), 2.35 (s, 3H).

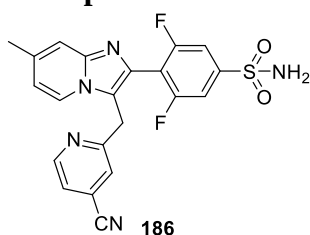
Example 185



The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: $[M+H]^+ = 472.1$.

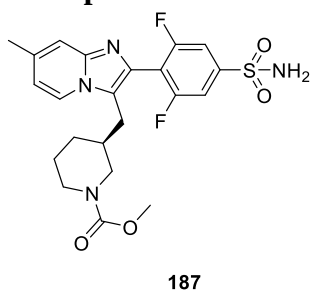
5

Example 186



The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: $[M+H]^+ = 440.2$.

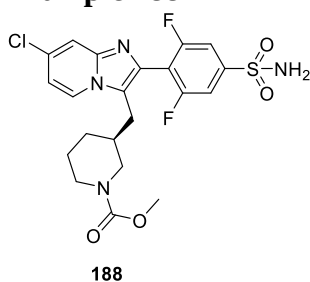
Example 187



10

The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: $[M+H]^+ = 479.1$.

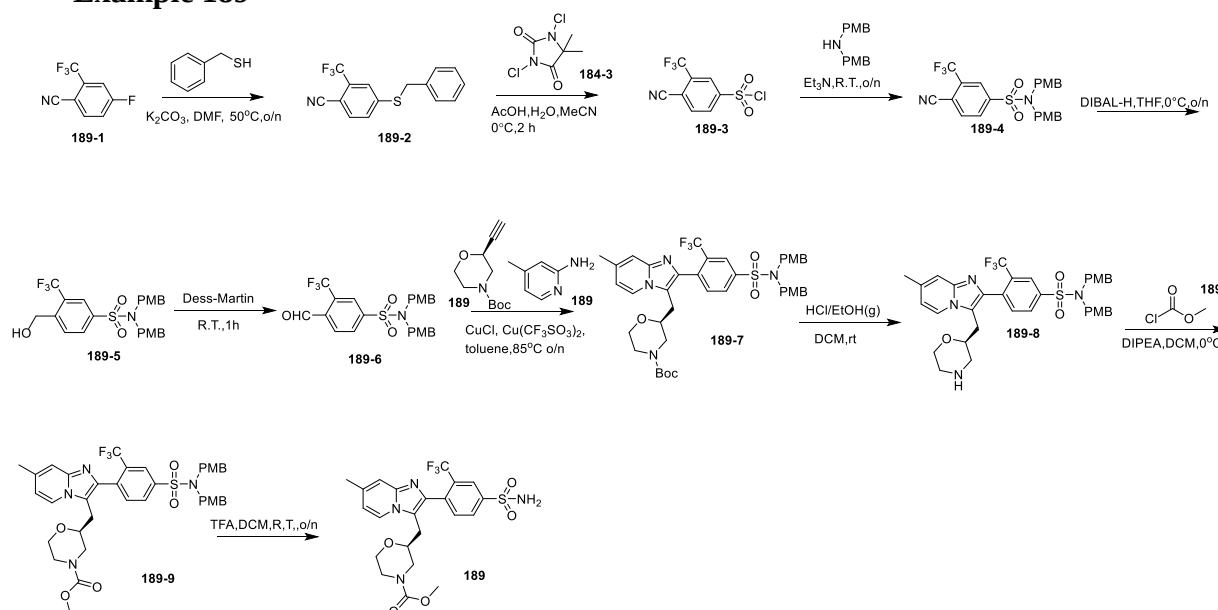
Example 188



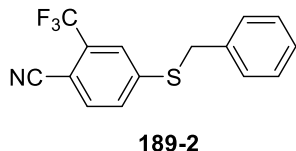
15

The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: $[M+H]^+ = 499.1$.

Example 189

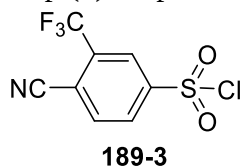


Step (1) Preparation of 4-(benzylthio)-2-(trifluoromethyl)benzonitrile



4-fluoro-2-(trifluoromethyl)benzonitrile (5.00 g, 26.44 mmol) was dissolved in *N,N*-dimethylformamide (30 mL) and potassium carbonate (7.50 g, 54.35 mmol) was added. The reaction system was added with benzylmercaptan (2.80 mL), heated to 50 °C and stirred overnight. Then the reaction system was cooled to room temperature, added with water (80 mL) to quench the reaction, and extracted with ethyl acetate (40 mL $\times$ 3). The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was purified by silica gel column chromatography to give **intermediate 189-2** (6.80 g, 83.3% yield) in the form of a yellow solid. LC-MS: $[M + H]^+ = 293.7$.

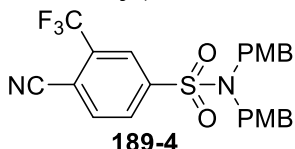
Step (2) Preparation of 4-cyano-3-(trifluoromethyl)benzenesulfonyl chloride



Intermediate 189-2 (6.80 g, 23.18 mmol) was dissolved in a prepared solvent (95 mL of acetonitrile, 4 mL of acetic acid, 2.5 mL of water) under nitrogen atmosphere. The reaction system was cooled to 0 °C, added with 1,3-dichloro-5,5-dimethylimidazolidine-2,4-dione (9.14 g, 46.37 mmol) in portions, and reacted for 2 h. Then the reaction system was added with water (100) to quench the reaction and extracted with ethyl acetate (60 mL $\times$ 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate,

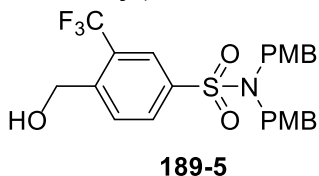
filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 189-3** (3.50 g, 51.5% yield) in the form of a white solid. LC-MS: $[M + H]^+ = 270.2$.

Step (3) Preparation of 4-cyano-*N,N*-bis(4-methoxybenzyl)-3-(trifluoromethyl)benzenesulfonamide



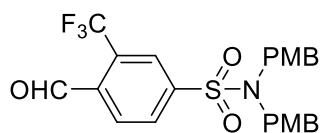
Triethylamine (3.93 g, 38.94 mmol) and bis(4-methoxybenzyl)amine (3.34 g, 12.98 mmol) were dissolved in dichloromethane (85 mL) under nitrogen atmosphere. The reaction system was cooled to 0 °C, added with **intermediate 189-3** (3.50 g, 12.98 mmol) in portions, warmed to room temperature and stirred overnight. Then the reaction system was added with water (50 mL) to quench the reaction. The aqueous phase was extracted with dichloromethane (50 mL $\times$ 2). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 189-4** (5.00 g, 75.1% yield) in the form of a light yellow solid, LC-MS: $[M + H]^+ = 512.7$.

Step (4) Preparation of 4-(hydroxymethyl)-*N,N*-bis(4-methoxybenzyl)-3-(trifluoromethyl)benzenesulfonamide



Intermediate 189-4 (4.40 g, 8.97 mmol) was dissolved in anhydrous tetrahydrofuran (80 mL). The reaction system was cooled to 0 °C, added dropwise with a solution of diisobutylaluminum hydride in *n*-hexane (1 N, 27 mL), slowly warmed to room temperature and stirred overnight. Then the reaction system was added with diluted hydrochloric acid (80 mL, 1 N) to quench the reaction, and then filtered, and the filtrate was extracted with ethyl acetate (50 mL $\times$ 3). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give **intermediate 189-5** (5.00 g, crude product) in the form of a white solid, which was used directly in the next step. LC-MS: $[M + H]^+ = 494.6$.

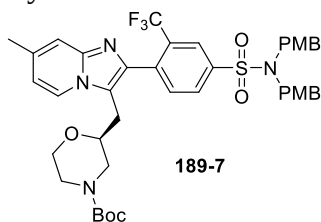
Step (5) Preparation of 4-formyl-*N,N*-bis(4-methoxybenzyl)-3-(trifluoromethyl)benzenesulfonamide



189-6

Intermediate 189-5 (5.00 g, crude product) was dissolved in anhydrous dichloromethane (70 mL), and Dess-Martin periodinane (4.85 g, 11.56 mmol) was added in portions. The reaction system was stirred at room temperature for 1.5 h. Then the reaction system was added with saturated aqueous sodium thiosulfate solution (20 mL) to quench the reaction, and then added with water (30 mL). The aqueous phase was extracted with dichloromethane (40 mL $\times$ 2). The organic phases were combined, washed with saturated brine (50 mL), dried over anhydrous sodium sulfate, filtered and dried by rotary evaporation. The crude product was purified by silica gel column chromatography to give **intermediate 189-6** (2.00 g, 45.2% yield over two steps) in the form of a white solid. LC-MS: $[M + H]^+ = 515.5$.

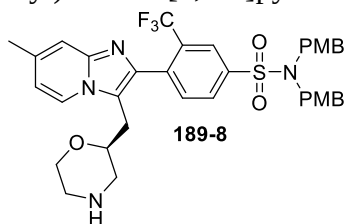
Step (6) Preparation of *tert*-butyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2-(trifluoromethyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



189-7

Intermediate 189-6 (500 mg, 1.01 mmol), 4-methylpyridin-2-amine (110 mg, 1.01 mmol), *tert*-butyl (S)-2-ethynylmorpholine-4-carboxylate (214 mg, 1.01 mmol), copper(I) chloride (30 mg, 0.30 mmol) and copper(II) trifluoromethanesulfonate (110 mg, 0.30 mmol) were dissolved in anhydrous toluene (5 mL) under nitrogen atmosphere. The reaction system was warmed to 85 °C, added with 5 drops of *N,N*-dimethylacetamide, and stirred at 85 °C overnight. The reaction system was cooled to room temperature and filtered, and the filtrate was concentrated. The crude product was purified by silica gel column chromatography to give **intermediate 189-7** (280 mg, 34.8% yield) in the form of a colorless transparent oil, LC-MS: $[M + H]^+ = 794.4$.

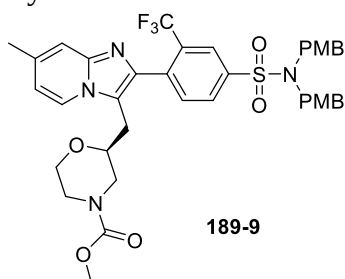
Step (7) Preparation of (S)-*N,N*-bis(4-methoxybenzyl)-4-(7-methyl-3-(morpholin-2-ylmethyl)imidazo[1,2-*a*]pyridin-2-yl)-3-(trifluoromethyl)benzenesulfonamide



189-8

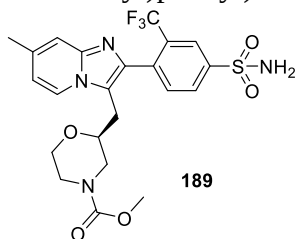
Intermediate 189-7 (280 mg, 0.35 mmol) was dissolved in anhydrous dichloromethane (5 mL), and a solution of HCl in ethanol (0.6 mL, 4 N) was added dropwise. The reaction system was stirred at room temperature for 3 h. Then the reaction system was concentrated under reduced pressure to give **intermediate 189-8** (300 mg, crude product) in the form of a white solid, LC-MS: $[M + H]^+ = 694.5$.

Step (8) Preparation of methyl (S)-2-((2-(4-(*N,N*-bis(4-methoxybenzyl)sulfamoyl)-2-(trifluoromethyl)phenyl)-7-methylimidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 189-8 (300 mg, crude product) was dissolved in anhydrous dichloromethane (4 mL) under nitrogen atmosphere, and *N,N'*-diisopropylethylamine (225 mg, 1.75 mmol) was added under nitrogen atmosphere. The reaction system was cooled in an ice water bath, added dropwise with a solution of methyl chloroformate in dichloromethane (10 mg/mL, 3.3 mL) and then reacted at 0 °C for 4 h. Then the reaction system was added with water (20 mL) to quench the reaction and extracted with dichloromethane (10 mL × 3). The organic phases were combined, washed with saturated brine (10 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation to give a crude product. The crude product was purified by silica gel column chromatography to give **intermediate 189-9** (200 mg, 74.2% yield) in the form of a white solid, LC-MS $[M + H]^+ = 752.4$.

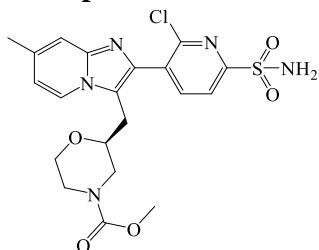
Step (9) Preparation of methyl (S)-2-((7-methyl-2-(4-sulfamoyl)-2-(trifluoromethyl)phenyl)imidazo[1,2-*a*]pyridin-3-yl)methyl)morpholine-4-carboxylate



Intermediate 189-9 (200 mg, 0.26 mmol) was dissolved in dichloromethane (2 mL), and trifluoroacetic acid (2 mL) was added. The reaction system was stirred overnight at room temperature. Then the reaction system was directly concentrated by rotary evaporation and purified by preparative reverse phase chromatography to give **compound 189** (51 mg, 37.5% yield) in the form of a white solid, LC-MS: $[M + H]^+ = 512.6$.

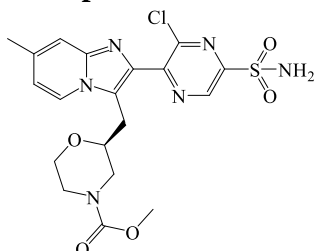
¹H NMR (400 MHz, CDCl₃) δ 8.25 (brs, 1H), 8.13 (brs, 1H), 8.04 (brs, 1H), 7.60 (brs, 1H), 7.47 (brs, 1H), 7.00 – 6.65 (m, 3H), 4.00 – 3.70 (m, 3H), 3.38 (s, 3H), 3.52 – 3.28 (m, 2H), 3.00 – 2.77 (m, 3H), 2.62 – 2.50 (m, 1H), 2.49 (s, 3H).

Example 190



The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: [M+H]⁺ = 480.2.

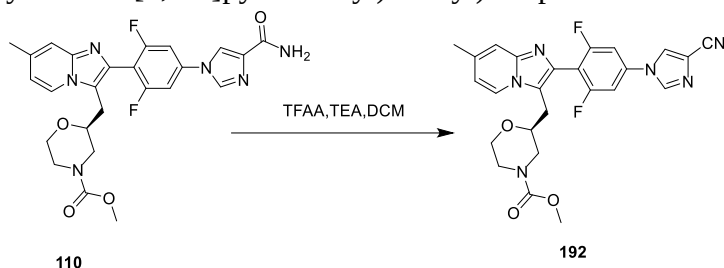
Example 191



The compound was prepared according to the preparation method as described in **Example 181**. LC-MS: [M+H]⁺ = 481.4.

Example 192

Preparation of methyl (S)-2-((2-(4-(4-cyano-1H-imidazol-1-yl)-2,6-difluorophenyl)-7-methylimidazo[1,2-a]pyridin-3-yl)methyl)morpholine-4-carboxylate



Compound 110 (100 mg, 0.2 mmol) and triethylamine (89 mg, 0.88 mmol) were dissolved in dichloromethane (5 mL), and a solution of trifluoroacetic anhydride (82 mg, 0.39 mmol) in dichloromethane (1 mL) was added dropwise under nitrogen atmosphere in an ice water bath. The reaction system was stirred overnight at room temperature. Then the reaction system was diluted with dichloromethane (30 mL), washed successively with saturated sodium bicarbonate solution (15 mL × 2) and saturated brine (15 mL), dried over anhydrous sodium sulfate, filtered and concentrated by rotary evaporation. The crude product was purified by preparative

chromatography to give **compound 192** (30 mg, 30% yield) in the form of a white solid. LC-MS: $[M+H]^+ = 492.8$.

^1H NMR (400 MHz, MeOD) δ 8.83 (d, $J = 7.1$ Hz, 1H), 8.61 (d, $J = 1.2$ Hz, 1H), 8.54 (d, $J = 1.1$ Hz, 1H), 7.80 (d, $J = 8.5$ Hz, 2H), 7.72 (s, 1H), 7.39 (d, $J = 7.1$ Hz, 1H), 4.01 (d, $J = 12.9$ Hz, 1H), 3.83 (d, $J = 13.4$ Hz, 1H), 3.76 (d, $J = 10.8$ Hz, 1H), 3.71 - 3.67 (m, 1H), 3.69 (s, 3H), 3.37 - 3.35 (m, 1H), 3.28 - 3.25 (m, 2H), 2.97–2.85 (m, 1H), 2.82 – 2.67 (m, 1H), 2.64 (s, 3H).

Biological Experiments

Example A: Evaluation of *In Vitro* Biological Activity

The antagonist property of the compounds disclosed herein was determined using the FLIPR (fluorescence imaging plate reader) method, showing that the compounds are inhibitors for the intracellular calcium increase induced by the activation of hP2X3 (human purinergic P2X receptor subtype 3, Accession No. NM_002559.4) expressed in HEK293 cells (human renal epithelial cell line, ATCC).

HEK293 cells that stably express hP2X3 were cultured in DMEM high glucose medium containing 10% FBS (fetal bovine serum, Gibco, 10099-141), 1% penicillin-streptomycin (Gibco, 15140-122) and 1 mg/mL G418 (Invitrogen, 10131027) in a cell incubator at 37 °C, 5% humidity. Cells at 400,000 cells/mL were seeded into a 384-well plate (10,000 cells/well) 18-24 h prior to the FLIPR experiment and then incubated overnight in a cell incubator. On the day of the experiment, the medium was discarded and the cells were washed in an FLIPR buffer (each 30 mL buffer contains 0.3 mL of probenecid (Thermo, P36400), 0.6 mL of 1 M HEPES (Invitrogen, 15630080) and 29.1 mL of HBSS (Invitrogen, 14065056)). Each well was added with 20 μL of 0.5 $\times$ Calcium 6 fluorescent dye (Molecular Devices, R8190) and then was subjected to dye-loading incubation at 37 °C for 1.5 h. Each well was added with 10 μL of test compound (which was dissolved in DMSO at a concentration of 10 mM and serially diluted with buffer) or vehicle, and then was left to equilibrate for 30 min at room temperature. The cell plate was then placed in the FLIPR for baseline fluorescence measurements (excitation at 485 nm and emission at 525-535 nm). An agonist (BZ-ATP (Sigma, B6396) at a final concentration of 2.5 μM) or a vehicle (ultrapure water) was then added at 10 μL /well, fluorescence values were measured for 2 min at 1-second intervals, and finally the output fluorescence counts were analyzed.

The IC₅₀ values obtained by the above method are shown in Table 2.

Table 2. IC₅₀ values of compounds of Examples 1-192 for P2X3 receptor

Compound	P2X3 IC ₅₀ (μM)	Compound	P2X3 IC ₅₀ (μM)
Compound 1	D	Compound 2	A
Compound 3	B	Compound 4	A
Compound 5	A	Compound 6	C
Compound 7	A	Compound 8	D
Compound 9	D	Compound 10	A
Compound 11	B	Compound 12	A
Compound 13	B	Compound 14	D
Compound 15	A	Compound 16	B
Compound 17	B	Compound 18	C
Compound 19	D	Compound 20	D
Compound 21	D	Compound 22	D
Compound 23	D	Compound 24	D
Compound 25	A	Compound 26	C
Compound 27	A	Compound 28	B
Compound 29	A	Compound 30	B
Compound 31	D	Compound 32	D
Compound 33	D	Compound 34	A

Compound 35	D	Compound 36	B
Compound 37	A	Compound 38	A
Compound 39	D	Compound 40	D
Compound 41	D	Compound 42	A
Compound 43	A	Compound 44	A
Compound 45	D	Compound 46	D
Compound 47	D	Compound 48	C
Compound 49	A	Compound 50	A
Compound 51	C	Compound 52	A
Compound 53	A	Compound 54	A
Compound 55	A	Compound 56	A
Compound 57	A	Compound 58	A
Compound 59	A	Compound 60	A
Compound 61	A	Compound 62	D
Compound 63	C	Compound 64	C
Compound 65	C	Compound 66	C
Compound 67	B	Compound 68	B

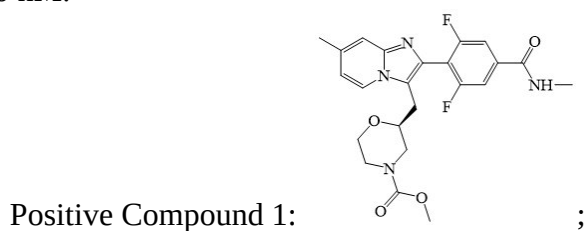
Compound 69	B	Compound 70	B
Compound 71	A	Compound 72	C
Compound 73	C	Compound 74	C
Compound 75	B	Compound 76	C
Compound 77	A	Compound 78	B
Compound 79	B	Compound 80	B
Compound 81	A	Compound 82	B
Compound 83	C	Compound 84	C
Compound 85	C	Compound 86	B
Compound 87	C	Compound 88	B
Compound 89	B	Compound 90	B
Compound 91	A	Compound 92	B
Compound 93	B	Compound 94	C
Compound 95	B	Compound 96	B
Compound 97	C	Compound 98	B
Compound 97	C	Compound 98	B
Compound 99	B	Compound 100	B

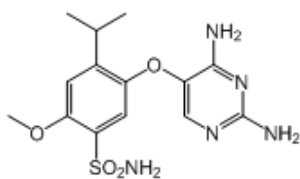
Compound 101	B	Compound 102	A
Compound 103	C	Compound 104	C
Compound 105	A	Compound 106	B
Compound 107	B	Compound 108	D
Compound 109	C	Compound 110	B
Compound 111	B	Compound 112	D
Compound 113	D	Compound 114	B
Compound 115	D	Compound 116	B
Compound 117	B	Compound 118	D
Compound 119	B	Compound 120	A
Compound 121	D	Compound 122	D
Compound 123	C	Compound 124	A
Compound 124-2	A	Compound 125	A
Compound 126	B	Compound 127	A
Compound 128	D	Compound 129	B
Compound 130	D	Compound 131	B
Compound 132	A	Compound 133	B

Compound 134	B	Compound 135	A
Compound 136	C	Compound 137	B
Compound 138	A	Compound 139	A
Compound 140	C	Compound 141	B
Compound 142	A	Compound 143	A
Compound 144	B	Compound 145	C
Compound 146	C	Compound 147	B
Compound 148	A	Compound 149	A
Compound 150	A	Compound 151	A
Compound 152	A	Compound 153	C
Compound 154	A	Compound 155	A
Compound 156	A	Compound 157	A
Compound 158	A	Compound 159	A
Compound 160	A	Compound 161	B
Compound 162	A	Compound 163	A
Compound 164	B	Compound 165	A
Compound 166	A	Compound 167	A

Compound 168	B	Compound 169	A
Compound 170	C	Compound 171	0.0021
Compound 172	0.0007	Compound 173	0.0018
Compound 174	0.0016	Compound 175	>10
Compound 176	0.0075	Compound 177	0.0011
Compound 178	0.0001	Compound 179	0.0004
Compound 180	0.0002	Compound 181	0.0018
Compound 182	0.0035	Compound 183	0.0055
Compound 184	0.0056	Compound 185	0.013
Compound 186	0.065	Compound 187	0.050
Compound 188	0.009	Compound 189	0.0100
Compound 190	A	Compound 191	A
Compound 192	B		
Positive compound 1	0.0115	Positive compound 2	0.0522

A: $IC_{50} \leq 10$ nM, B: $10 < IC_{50} \leq 50$ nM, C: $50 < IC_{50} \leq 200$ nM, D: $200 < IC_{50} \leq 5000$ nM.





Positive Compound 2:

As can be seen from the data in Table 2, the compounds disclosed herein exhibit good inhibitory activity against P2X3, wherein most of the compounds show stronger inhibitory activity than that of positive compound 2, and some of the compounds show stronger inhibitory activity than that of positive compound 1. Compounds with stronger inhibitory activity than that of positive compound 2 are preferred, and compounds with stronger inhibitory activity than that of positive compound 1 are more preferred.

Example B: Evaluation of *In Vitro* P2X2/3 Receptor Selectivity

The selectivity of the compounds disclosed herein for the P2X2/3 receptor was determined using the FLIPR (fluorescence imaging plate reader) method, showing that the compounds were inhibitors for the intracellular calcium increase induced by the activation of hP2X2/3 (heterodimeric receptors formed by human purinergic P₂X receptors subtype 2 and subtype 3, Accession No. NM_002559.4 for P2X2 and Accession No. NM_002559.4 for P2X3) expressed in HEK293 cells (human renal epithelial cell line, ATCC).

HEK293 cells that stably express hP2X2/3 were cultured in DMEM high glucose medium containing 10% FBS (fetal bovine serum, Gibco, 10099-141), 1% penicillin-streptomycin (Gibco, 15140-122) and 1 mg/mL G418 (Invitrogen, 10131027) in a cell incubator at 37 °C, 5% humidity. Cells at 250000 cells/mL were seeded into a 96-well plate (25000 cells/well) 18-24 h prior to the FLIPR experiment and then incubated overnight in a cell incubator. On the day of the experiment, the medium was discarded and the cells were washed in an FLIPR buffer (0.3 mL of probenecid (Thermo, P36400), 0.6 mL of 1 M HEPES (Invitrogen, 15630080) and 29.1 mL of HBSS (Invitrogen, 14065056) per 30 mL of buffer). Each well was added with 75 µL of 1 mM Fluo-4 AM fluorescent dye (Thermo, F14202) and then was subjected to dye-loading incubation at 37 °C for 1.0 h. The 96-well plate was then washed once with buffer, added with a buffer containing a test compound or a vehicle at 50 µL per well and then incubated for 30 min at room temperature. The cell plate was then placed in the FLIPR for baseline fluorescence measurements (excitation at 485 nm and emission at 525-535 nm). An agonist (BZ-ATP (Sigma, B6396) at a final concentration of 5. µM) or a vehicle (ultrapure water) was then added at 50 µL/well, fluorescence values were measured for 2 min at 1-second intervals, and finally the output fluorescence counts were analyzed.

The IC₅₀ values obtained by the above method are shown in Table 2.

Table 3. IC₅₀ values and folds of selectivity of some of the compounds for P2X_{2/3} receptors

Compound	IC ₅₀	Folds of selectivity	Compound	IC ₅₀	Folds of selectivity
Compound 2	>60	A	Compound 3	>60	A
Compound 4	14.31	A	Compound 5	>60	A
Compound 7	>60	A	Compound 10	37.39	A
Compound 11	17.56	B	Compound 12	12.13	A
Compound 13	14.44	B	Compound 15	21.56	B
Compound 17	29.81	B	Compound 25	>60	A
Compound 54	>60	A	Compound 58	>60	A
Compound 61	>60	A	Compound 138	>60	A
Compound 149	>60	A	Compound 151	>60	A
Compound 158	>60	A	Compound 159	>60	A
Compound 160	>60	A	Compound 161	>60	A
Compound 162	>60	A	Compound 163	>60	A
Compound 164	>60	A	Compound 165	>60	A
Compound 166	>60	A	Compound 167	>60	A
Compound 168	>60	A	Compound 169	>60	A
Compound 171	36.78	45975	Compound 172	>60	>85714
Compound 173	33.86	18811	Compound 191	>60	A
Positive compound 1	>60	>5217	Positive compound 2	0.36	6.9

A: folds of selectivity > 1000, B: 200 < folds of selectivity ≤ 1000, C: folds of selectivity ≤ 200.

As can be seen from the data in Table 3, the compounds disclosed herein exhibit poor inhibitory activity against P2X2/3 heterodimeric receptor and good selectivity for P2X3 and P2X2/3. Folds of selectivity = (IC₅₀ for P2X2/3)/(IC₅₀ for P2X3). Compounds with the folds of selectivity of more than 200 are preferred, and compounds with the folds of selectivity of more than 1000 are more preferred.

Example C: Activity Test in Citric Acid Only Cough Models

Male Dunkin Hartley guinea pigs (300-350 g) were placed in an animal atomization box, and then the door of the atomization box was closed while the ultrasonic atomizer (Guangdong Yuehua) was turned on. 17.5% citric acid gas was introduced into the atomization box at a maximum atomization rate (about 2 mL/min) for 20 s, and coughs of the animals within 10 min were continuously observed from the time of the start of atomization. During the 10-min observation, coughs of the animals were counted manually, and the number of coughs were judged according to cough postures of the guinea pigs, such as abdominal twitching, mouth opening, and head inclining downward abruptly. The number of coughs during the first 5 min and that during 10 min were recorded, and the cough latency of the guinea pigs, namely the time from the initiation of citric acid induction to the appearance of the first cough, was also recorded.

Table 4. The number of coughs, cough inhibition rate and cough latency obtained *in vivo* for some of the compounds

Group		Mean number of coughs	Cough inhibition rate Vs Vehicle (%)	Cough inhibition rate Vs Baseline (%)	Mean cough latency
Vehicle group	Before administration	17.22	/	/	108.44
	After administration	18.33			105.78
Positive compound 1 30 mg/kg	Before administration	18.63	42.6	47.0	80.13
	After administration	9.88			226.75
Compound 2 30 mg/kg	Before administration	18.50	34.2	38.7	94.50

	After administrati on	11.33			194.83
Compound 7 30 mg/kg	Before administrati on	18.50	38.3	42.6	87.13
	After administrati on	10.63			177.00
Compound 53 30 mg/kg	Before administrati on	18.63	47.7	51.7	138.00
	After administrati on	9.00			250.13
Compound 149 30 mg/kg	Before administrati on	19.33	62.7	66.8	75.92
	After administrati on	6.42			229.00
Compound 161 30 mg/kg	Before administrati on	18.57	63.5	66.2	130.71
	After administrati on	6.29			302.43
Compound 162 30 mg/kg	Before administrati on	18.29	42.8	46.1	144.43
	After administrati on	9.86			278.43
Compound 164 30 mg/kg	Before administrati on	18.29	65.2	67.2	144.86
	After administrati on	6.00			275.14
Compound 165 30 mg/kg	Before administrati on	17.57	53.6	54.5	135.71
	After administrati on	8.00			190.71
Compound 172 30 mg/kg	Before administrati on	19.56	43.9	50.6	87.56

	After administrati on	9.67			118.89
--	-----------------------------	------	--	--	--------

The above experiment was conducted on the following representative compounds: compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164, compound 165 and compound 172. The results as shown in Table 4 and FIGs. 1, 2, 3 and 4 demonstrate that compared with the blank vehicle or the condition before administration, compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164, compound 165 and compound 172 at 30 mg/kg can effectively reduce the number of coughs and prolong the cough latency, and the efficacy of these compounds is not significantly different from that of positive compound 1, indicating that compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164, compound 165 and compound 172 can reduce the number of coughs and improve the cough latency and their efficacy is equivalent to that of the positive compound.

Example D: Activity Test in ATP-Citric Acid Cough Models

Male Dunkin Hartley guinea pigs (300-400 g) were placed into a whole-body plethysmography box to adapt for 3-5 min, then ATP atomization is carried out for 2 min, and 3 min later, 5 min of citric acid atomization was followed, with all atomization rates being about 300 uL/min. The number of coughs and the cough latency of the animals within 10 min from the start of the citric acid atomization were recorded. During the 10-min observation, coughs of the animals were counted manually, and the coughs were judged according to cough postures of the guinea pigs, such as abdominal twitching, mouth opening, head inclining downward abruptly and the like. The number of coughs during 10 min were recorded, and the cough latency of the guinea pigs was also recorded.

The animals were divided into a vehicle group, a dextromethorphan (63 mg/kg) positive group, an AF-219 (30 mg/kg) positive group and Example 173 compound (3, 10, 30 mg/kg) administration groups, and all the compounds were orally administered. Except that dextromethorphan was administered 40 min before ATP-citric acid exposure, the remaining compounds were administered 2 h before ATP-citric acid exposure.

Table 5. Ratios of quinine/water intake obtained for the compounds

Compound		Number of coughs during 10 min	Cough inhibition ratio vs. vehicle (%)
Vehicle		20.83	
Dextromethorphan group (63 mg/kg)		8.00	61.60
AF-219	30 mg/kg	11.42	45.20
Compound 172	3 mg/kg	14.42	30.80
	10 mg/kg	10.17	51.20
	30 mg/kg	9.83	52.80

As shown in Table 5 and FIGs. 5 and 6, all the compound treatment groups show a tendency of reduce in the number of coughs as compared with the vehicle group, both the dextromethorphan and compound 172 can significantly reduce the number of coughs, and compound 172 shows inhibition in a dose-dependent manner. Compound 172 exhibits stronger cough inhibition effect than positive compound 2.

Example E: *In Vitro* Cytotoxicity Assay

In vitro cytotoxicity assay for the compounds disclosed herein was performed in HepG2 cells using the CCK-8 method. HepG2 cells (Beina Bio) in the logarithmic growth phase were collected, the concentration of cell suspension was adjusted, and then the cells were plated on a 96-well cell culture plate at 50,000 cells/well. The cells were then incubated overnight in a cell incubator of 5% humidity at 37 °C, and after 80-90% cell confluence was achieved, test compounds or vehicle (DMSO) at various concentration gradients were added after medium change. The plate was incubated in a cell incubator of 5% humidity at 37 °C for 48 h. After the treatment, the medium in the plate was discarded. The plate was washed twice with PBS, added with CCK-8 working solution (Beyotime) at 100 µL per well, and then incubated at 37 °C for 1.5 h away from light. Absorbance at OD_{450nm} was measured for each well on a microplate reader, and CC₅₀ value of each compound was analyzed and calculated.

The CC₅₀ values obtained by the above method are shown in Table 6.

Table 6. CC₅₀ values obtained for some of the compounds

Compound	HepG2 CC ₅₀ (µM)	Compound	HepG2 CC ₅₀ (µM)
Compound 2	123.5	Compound 3	59.30
Compound 4	13.21	Compound 5	72.02
Compound 7	194.6	Compound 10	15.77
Compound 11	79.44	Compound 12	52.73

Compound 13	66.48	Compound 15	52.63
Compound 17	50.95	Compound 25	>100
Compound 26	>100	Compound 149	>100
Compound 158	>100	Compound 159	>100
Compound 160	>100	Compound 161	>100
Compound 162	>100	Compound 163	>100
Compound 164	>100	Compound 165	>100
Compound 166	>100	Compound 167	>100
Compound 168	>100	Compound 169	>100
Compound 171	200	Compound 172	57.05
Compound 173	>200	Compound 174	>200
Compound 176	>200	Compound 181	138.14
Compound 182	>200	Compound 184	>200
Positive compound 1	128.4	Positive compound 2	>200

As can be seen from the data in Table 4, most of the compounds disclosed herein exhibit relatively good safety, and the CC₅₀ ranges are all more than 10 µM, which satisfies the requirement for *in vitro* cytotoxicity of general compounds. Compounds with a CC₅₀ value of more than 30 µM are preferred, and compounds with a CC₅₀ value of more than 50 µM are more preferred.

Example F: Test of *In Vitro* Metabolic Stability

The *in vitro* metabolic stability of the compounds disclosed herein was determined through incubation of liver microsomes of various species. A proper amount of test compound was added into a liver microsome reaction system (1 mg/mL liver microsome protein, 25 U/mL glucose-6 phosphate dehydrogenase, 1 mM NADP, 6 mM D-glucose 6-phosphate and 5 mM MgCl₂), and then the mixture was incubated in a water bath kettle at 37 °C to start the reaction. At each time point, 100 µL of the reaction mixture was added into a centrifuge tube containing 400 µL of internal standard working solution (containing a 200 ng/mL solution of dexamethasone, diclofenac, tolbutamide and labetalol in acetonitrile) precooled at 0 °C so as to terminate the reaction, and the mixture was then centrifuged at 10,000 g for 10 min at 4 °C. The supernatant was collected for LC-MS assay so as to obtain the values of *in vitro* metabolic half-life of the test compounds in liver microsomes of various species.

The T_{1/2} values obtained by the above method are shown in Table 7.

Table 7. T_{1/2} values obtained for some of the compounds

Compound	T_{1/2} in human liver microsome min	T_{1/2} in rat liver microsome min	T_{1/2} in guinea pig liver microsome min
Compound 2	895.35	52.34	95.33
Compound 3	260.22	28.72	50.49
Compound 4	61.05	4.91	4.38
Compound 5	648.70	37.31	47.62
Compound 7	127.80	19.58	20.89
Compound 10	222.61	30.80	13.51
Compound 11	151.19	5.65	4.27
Compound 12	91.92	7.78	7.56
Compound 13	61.51	2.24	2.25
Compound 15	190.59	25.39	16.51
Compound 17	997.99	19.35	8.61
Compound 25	297.97	227.75	NA
Compound 26	253.12	80.94	127.80
Compound 149	206.63	52.50	178.02
Compound 168	522.56	57.59	29.24
Compound 169	827.52	35.47	164.42
Compound 171	523.3	8.48	140.1
Example 172	824.3	223.4	410.3
Example 173	NA	101.2	646.6
Example 174	NA	398.3	1110.5
Example 175	>30	>30	>30
Example 176	235.9	3.61	60.96
Compound 177	324.61	/	/
Compound 178	1233.69	/	/
Compound 181	1400.2	187.65	234.42
Compound 182	217.27	10.95	60.96
Compound 184	24383.0	5.66	173.41

Compound 189	144.57	/	/
Positive compound 1	1454.06	53.42	112.04
Positive compound 2	NA	1090.01	892.21

Note: NA indicates essentially not metabolized; "/" indicates not detected.

As can be seen from the data in Table 5, the compounds disclosed herein exhibit relatively good metabolic stability in human, rat and guinea pig. As compared with positive compound 1, the compounds disclosed herein have similar metabolic half-life in human liver microsome but longer half-life in rat and guinea pig. In addition, as compared with positive compound 1, the metabolism of the compounds disclosed herein in rat and guinea pig are more close to that in human, which is beneficial to the related experimental evaluation of the compounds in rat and guinea pig. Compounds with a $T_{1/2} > 30$ min in human liver microsome are preferred, and compounds with a $T_{1/2} > 90$ min in human liver microsome are more preferred.

Example G: Two-Bottle Dysgeusia Test

Male SD rats (SPF grade, aged 6-8 weeks) were subjected to adaptive drinking training for 3 days after warehousing, and the specific training content was as follows: animals were fed in separate cages, and two bottles of water (both were common drinking water) were placed in each cage; during the adaptive training, drinking water was completely prohibited at night every day (drinking water bottles were removed out) and then resupplied during 8:30 am to 5:30 pm; and the operation was repeated for 3 days, the positions of the two bottles of water were changed every day, and the animals could eat freely during the whole adaptive training period. All drinking water bottles were removed out 20 h before the start of the formal experiment, and all the animals were deprived of water until the start of the experiment. In the experiment, all the animals were randomly grouped and then subjected to a single intraperitoneal injection of a test compound or a vehicle before water was resupplied. The administration time was determined according to the T_{max} of the test compound. Then the animals were placed in separate cages, and two bottles of drinking water (one bottle of normal drinking water and one bottle of 0.3 mM quinine water) were provided. The water intake of the animals was observed within 15 min, statistical analysis was performed based on quinine water intake/normal water intake, and positive compound 2 was used as the dysgeusia positive control.

Table 8. Ratios of quinine/water intake obtained for some of the compounds

Compound	Quinine/tap water*100%
Vehicle	4.54%
Positive compound 2	59.41%
Compound 2	5.49%
Compound 7	6.91%
Compound 53	7.65%
Compound 149	4.54%
Compound 161	5.03%
Compound 162	6.34%
Compound 164	6.69%
Compound 165	3.90%

The above experiment was conducted on the following representative compounds: compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164 and compound 165. The results as shown in Table 8 and FIG. 7 demonstrate that for compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164 and compound 165 at 20 mg/kg, the ratio of quinine/water intake is almost unaffected when compared with the blank vehicle while, and the ratio of quinine/water intake is remarkably reduced when compared with positive compound 2, indicating that the influence of compound 2, compound 7, compound 53, compound 149, compound 161, compound 162, compound 164 and compound 165 on animal taste is smaller.

Table 9. Ratios of quinine/water intake obtained for the compounds

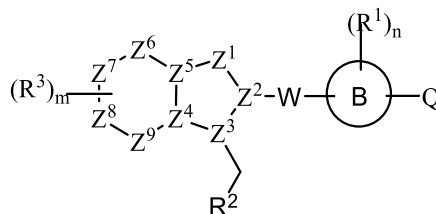
Compound		Quinine/tap water*100%
Vehicle		7.19%
Positive compound 2	10	18.62%
	20	34.99%
Compound 172	5	7.05%
	10	7.41%
	20	8.66%

The above experiment was conducted on the representative compound 172. The results as shown in Table 9 and FIG. 8 demonstrate that for compound 172 at 5 mg/kg, 10 mg/kg and 20 mg/kg, the ratio of quinine /water intake of rats is almost unaffected when compared with the blank vehicle, indicating that compound 172 has small influence on animal taste; as compared with the positive compound 2, the ratio of quinine/water intake is remarkably reduced, showing that the risk of causing dysgeusia by compound 172 is far lower than that by positive compound 2, and compound 172 shows better safety.

Although examples of the present invention are illustrated and described above, it will be appreciated that the above examples are exemplary and not to be construed as limiting the present invention, and that changes, modifications, substitutions and alterations can be made to the above examples by those skilled in the art within the scope of the present invention.

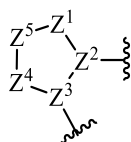
CLAIMS

1. A heterocyclic compound of formula I or a stereoisomer, a geometric isomer, a tautomer, a nitrogen oxide, a hydrate, a solvate, a metabolite, an ester, a pharmaceutically acceptable salt or a prodrug thereof,

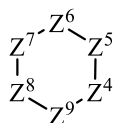


I

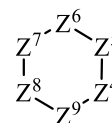
wherein,



is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl, wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



$Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8$ and Z^9 independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z^6 or Z^9 represents a single bond;

m is 1, 2, 3 or 4;

R^3 is H, halogen, -OH, a C_1 - C_6 haloalkyl, -CN or a C_1 - C_6 alkyl; when m is not 1, R^3 is independently the same or different; or when two adjacent R^3 are present, they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkyl, a C_3 - C_6 cycloalkenyl, a C_3 - C_9 heterocycloalkyl, a C_3 - C_9 heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl; wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N;

and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C₃-C₉ heterocycloalkyl; wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

n is 1, 2, 3 or 4;

R¹ is independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); when a plurality of substituents are present, they are the same or different; and when n is not 1, R¹ is independently the same or different;

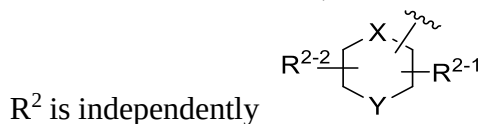
Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₃-C₁₀ cycloalkyl, the substituted C₆-C₁₀ aryl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or

different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₂-C₆ alkynyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₂-C₆ alkynyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl, the substituted phenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -COOCH₃, -NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d, wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different; or

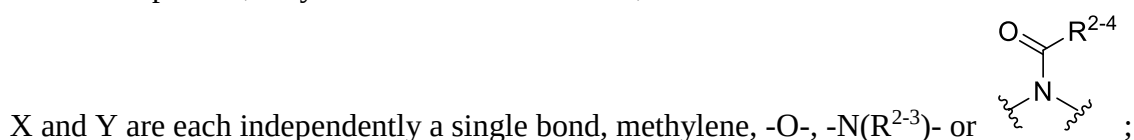
R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; wherein the C₃-C₉ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; the C₃-C₉ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; the 5-10 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e.,

two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b, wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;



5-6 membered heteroaryl; wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CN, an unsubstituted or halogen-substituted C₁-C₆ alkyl, an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-, -C(=O)R²⁻⁷ or -C(=O)N(R²⁻⁸R²⁻⁹), wherein when a plurality of substituents are present, they are the same or different;

R²⁻¹ and R²⁻² are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

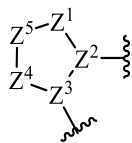


R²⁻³, R²⁻⁴, R²⁻⁵ and R²⁻⁶ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

R²⁻⁷ is independently H, halogen, -OH, -CN, an unsubstituted or halogen-substituted C₁-C₆ alkyl, or an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-; and

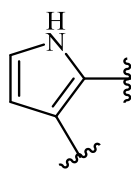
R^{2-8} and R^{2-9} are each independently H, or an unsubstituted or halogen-substituted C_1-C_6 alkyl.

2. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein when

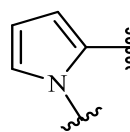


is a 5 membered heteroaryl, the 5 membered heteroaryl is pyrrolyl, furanyl, thienyl,

imidazolyl or pyrazolyl, the pyrrolyl is preferably

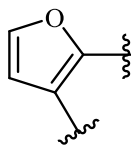


or

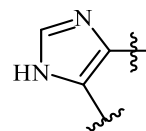


, the furanyl is

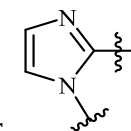
preferably



, the imidazolyl is preferably

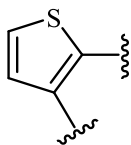


or

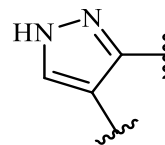


, the thienyl is

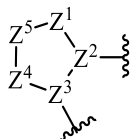
preferably



and the pyrazolyl is preferably

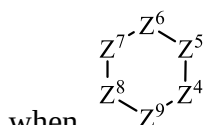
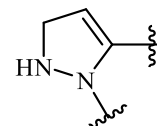


; and/or



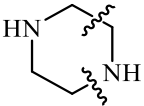
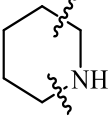
10 when is a 5 membered heterocycloalkenyl, the 5 membered heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered

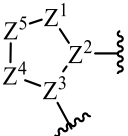
heterocycloalkenyl is preferably 2,3-dihydro-1H-pyrazolyl, and more preferably and/or

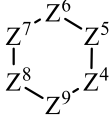


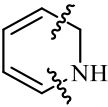
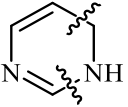
when is a 5-6 membered heterocycloalkyl, the 5-6 membered heterocycloalkyl

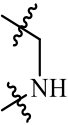
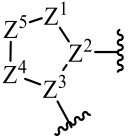
15 is a 6 membered heterocycloalkyl, wherein the 6 membered heterocycloalkyl contains 1-3 N heteroatoms; and the 5-6 membered heterocycloalkyl is preferably piperazinyl or piperidinyl, the

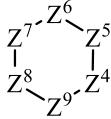
piperazinyl is preferably  ; and the piperidinyl is preferably  ; wherein the

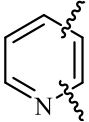
right-side carbon-nitrogen bond  represents the position fusing to  ; and/or

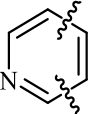
when  is a 5-6 membered heterocycloalkenyl, the 5-6 membered heterocycloalkenyl is a 6 membered heterocycloalkenyl, wherein the 6 membered heterocycloalkenyl contains 1-3 N heteroatoms; and the 5-6 membered heterocycloalkenyl is preferably 1,2-dihydropyridinyl or 1,6-dihydropyrimidinyl, the 1,2-dihydropyridinyl is

preferably , and the 1,6-dihydropyrimidinyl is preferably  ; wherein the

right-side carbon-nitrogen bond  represents the position fusing to  ; and/or

when  is a 5-6 membered heteroaryl, the 5-6 membered heteroaryl is a 6 membered heteroaryl, wherein the 6 membered heteroaryl contains 1-3 N heteroatoms; and the

5-6 membered heteroaryl is preferably pyridinyl, and the pyridinyl is preferably  or

 ; and/or

m is 1 or 2; and/or

Z^7 is substituted with R^3 , Z^8 is unsubstituted or substituted with R^3 , Z^6 and Z^9 are unsubstituted; when a plurality of R^3 are present, they are the same or different; and more preferably, Z^7 is substituted with R^3 , and Z^6 , Z^8 and Z^9 are unsubstituted; and/or

when R^3 is halogen, the halogen is fluorine, chlorine, bromine or iodine; and/or

when R^3 is a C_1 - C_6 alkyl, the C_1 - C_6 alkyl is C_1 - C_4 alkyl, and preferably methyl; and/or

when R^3 is a C_1 - C_6 haloalkyl, the halogen is fluorine, chlorine, bromine or iodine; and/or

when R^3 is a C_1 - C_6 haloalkyl, it contains one or more halogens, and preferably 1, 2 or 3 halogens; and/or

when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkyl, the C_3 - C_6 cycloalkyl is cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl; and/or

when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkenyl, the C_3 - C_6 cycloalkenyl is cyclopropenyl, cyclobutenyl, cyclopentenyl, cyclohexenyl or cyclohexadienyl; and/or

when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a C_3 - C_9 heterocycloalkyl, the C_3 - C_9 heterocycloalkyl is a C_4 - C_5 heterocycloalkyl, wherein the C_4 - C_5 heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a C_3 - C_9 heterocycloalkenyl, the C_3 - C_9 heterocycloalkenyl is a C_4 - C_5 heterocycloalkenyl, wherein the C_4 - C_5 heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when two adjacent R^3 are present and they, together with the ring atom to which they directly link, form a 5-6 membered heteroaryl, the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

n is 1 or 2; and/or

each R^1 is independently located at position α or β of the linkage between  and Z^2 ; and/or

when R^1 is halogen, the halogen is fluorine, chlorine, bromine or iodine; and/or

when R^1 is an unsubstituted or substituted C_1 - C_6 alkyl or an unsubstituted or substituted C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl is independently a C_1 - C_4 alkyl, and preferably methyl; and/or

when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, and the substituents are each independently halogen, the halogen is fluorine, chlorine, bromine or iodine; and/or

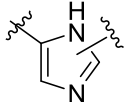
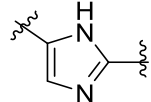
when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, and the substituents are each independently a C_1 - C_6 alkyl or a C_1 - C_6 alkyl-O-, the C_1 - C_6 alkyl in the substituents is independently a C_1 - C_4 alkyl, and preferably methyl; and/or

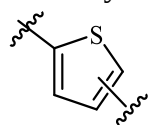
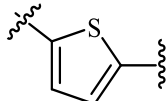
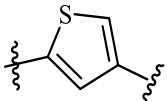
when R^1 is a substituted C_1 - C_6 alkyl or a substituted C_1 - C_6 alkyl-O-, the number of the substituents is independently 1, 2 or 3; and/or

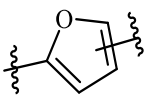
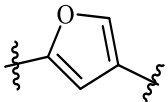
Q is located at position β or γ of the linkage between  and W;

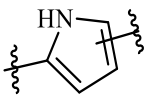
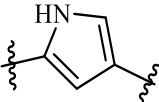
when  is a 5-6 membered heteroaryl, the 5-6 membered heteroaryl is a 5 membered heteroaryl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or

more of N, O and S, and the 5 membered heteroaryl is preferably imidazolyl, thienyl, furanyl or

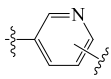
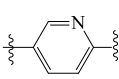
pyrrolyl, the imidazolyl is preferably  and more preferably , the thienyl

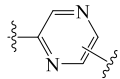
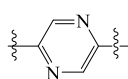
is preferably  and more preferably  or , the furanyl is

preferably  and more preferably , and the pyrrolyl is preferably

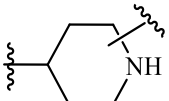
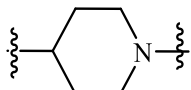
5  and more preferably ; or the 5-6 membered heteroaryl is a 6

membered heteroaryl, wherein the 6 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 6 membered heteroaryl is preferably pyridinyl or

pyrazinyl, the pyridinyl is preferably  and more preferably , and the

pyrazinyl is preferably  and more preferably ; and/or

10 when  is a C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S, the C₄-C₅ heterocycloalkyl is preferably piperidinyl, and the piperidinyl is preferably

15 , and more preferably ; and/or

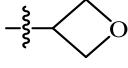
when Q is halogen, the halogen is fluorine, chlorine, bromine or iodine; and/or

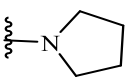
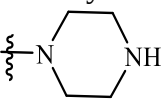
when Q is an unsubstituted or substituted C₁-C₆ alkyl or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently a C₁-C₄ alkyl; and/or

20 when Q is an unsubstituted or substituted C₂-C₆ alkenyl or an unsubstituted or substituted C₂-C₆ alkenyl-O-, the C₂-C₆ alkenyl is a C₂-C₄ alkenyl; and/or

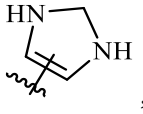
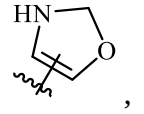
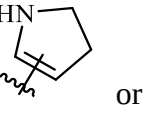
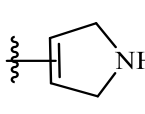
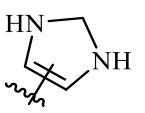
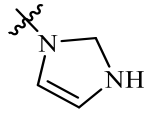
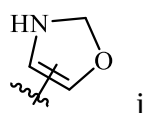
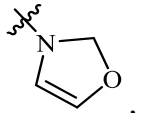
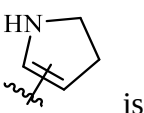
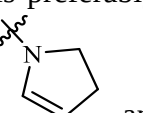
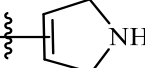
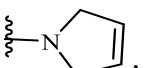
when Q is an unsubstituted or substituted C₃-C₁₀ cycloalkyl, the C₃-C₁₀ cycloalkyl is cyclopropyl, cyclobutyl, cyclopentyl, cyclohexyl, cycloheptyl or cyclooctyl; and/or

25 when Q is an unsubstituted or substituted C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉ heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected

from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl is preferably glycidyl, pyrrolidinyl or piperazinyl, the glycidyl is preferably ,

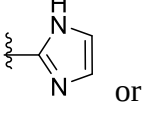
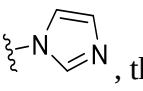
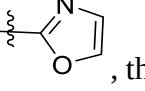
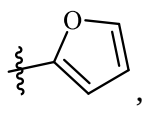
the pyrrolidinyl is preferably  and the piperazinyl is preferably ; and/or

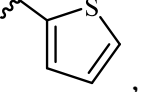
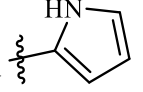
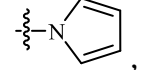
when Q is an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl is a C₃-C₅ heterocycloalkenyl, wherein the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the C₃-C₉ heterocycloalkenyl is preferably a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S, and the

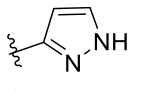
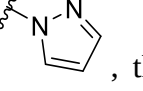
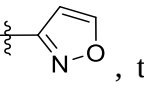
C₄-C₅ heterocycloalkenyl is preferably , ,  or , the  is preferably , the  is preferably , the  is preferably , and the  is preferably ; and/or

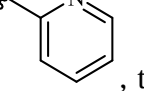
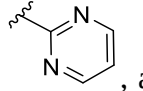
when Q is an unsubstituted or substituted C₆-C₁₀ aryl, the C₆-C₁₀ aryl is phenyl; and/or

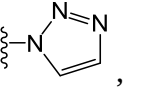
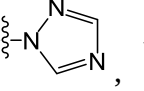
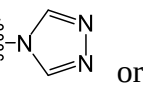
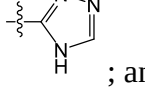
when Q is an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered heteroaryl, and preferably imidazolyl, oxazolyl, furanyl, thienyl, pyrrolyl, pyrazolyl, isoxazolyl, pyridinyl, triazolyl or pyrimidinyl, the imidazolyl is preferably

 or , the oxazolyl is preferably , the furanyl is preferably ,

the thienyl is preferably , the pyrrolyl is preferably  or , the

pyrazolyl is preferably  or , the isoxazolyl is preferably , the

pyridinyl is preferably , the pyrimidinyl is preferably , and the triazolyl is

preferably , ,  or ; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₃-C₁₀ cycloalkyl, a substituted C₆-C₁₀

aryl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of substituents is independently 1, 2, 3 or 4; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₃-C₁₀ cycloalkyl, a substituted C₆-C₁₀ aryl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine, or iodine, and preferably fluorine; and/or

the C₁-C₆ alkyl in the C₁-C₆ alkyl or the C₁-C₆ alkyl-O- that is unsubstituted or substituted with one or more R^a is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl; and/or

the C₃-C₆ cycloalkyl in the C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a is independently cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl; and/or

when R^a is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently a C₁-C₄ alkyl, and preferably methyl, ethyl or *tert*-pentyl; and/or

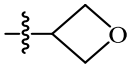
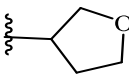
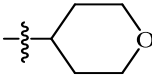
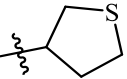
when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₂-C₆ alkenyl, or an unsubstituted or substituted C₂-C₆ alkenyl-O-, the C₂-C₆ alkenyl is independently a C₂-C₄ alkenyl, and preferably ethenyl; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₂-C₆ alkynyl, the C₂-C₆ alkynyl is ethynyl, 1-propynyl, 2-propynyl, 2-butyne, 1-butyne or butadiynyl, and preferably ethynyl; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₁₀ cycloalkyl, the C₃-C₁₀ cycloalkyl is a C₃-C₆ cycloalkyl, and preferably cyclopropyl, cyclobutyl, cyclopentyl and cyclohexyl; and/or

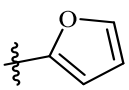
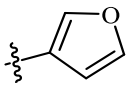
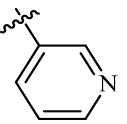
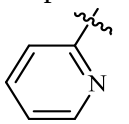
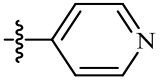
when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl is a C₃-C₅ heterocycloalkyl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N; the C₃-C₉

heterocycloalkyl is preferably a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₃-C₉ heterocycloalkyl is preferably glycidyl, tetrahydrofuranyl, tetrahydropyranyl or

tetrahydrothienyl, the glycidyl is preferably , the tetrahydrofuranyl is preferably , the tetrahydropyranyl is preferably , and the tetrahydrothienyl is preferably ; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl is a C₃-C₅ heterocycloalkenyl, wherein the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N; and the C₃-C₉ heterocycloalkenyl is preferably a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl is a 5-6 membered heteroaryl, wherein the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or two of N, O and S; and the 5-10 membered heteroaryl is preferably furanyl or pyridinyl, the furanyl is preferably

 or , and the pyridinyl is preferably ,  or ; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₂-C₆ alkynyl, a substituted C₃-C₁₀ cycloalkyl, a substituted phenyl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine; and/or

when R^d is independently halogen, the halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine; and/or

the number of R^d is independently 1, 2, 3, or 4, and preferably 1 or 2; and/or

the C₁-C₆ alkyl in the C₁-C₆ alkyl or the C₁-C₆ alkyl-O- that is unsubstituted or substituted with one or more R^d is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl; and/or

the C₃-C₆ cycloalkyl in the C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d is independently cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₂-C₆ alkynyl, a substituted C₃-C₆ cycloalkyl, a substituted phenyl, a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2; and/or

when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkyl, the C₃-C₉ heterocycloalkyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a C₃-C₅ heterocycloalkyl; and/or

when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, the C₃-C₉ heterocycloalkenyl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a C₃-C₅ heterocycloalkenyl; and/or

when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form an unsubstituted or substituted 5-10 membered heteroaryl, the 5-10 membered heteroaryl contains no heteroatom or one heteroatom selected from one of O, S and N in addition to the existing N atom, and is preferably a 5-6 membered heteroaryl; and/or

when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

the C₁-C₆ alkyl in the C₁-C₆ alkyl or the C₁-C₆ alkyl-O- that is unsubstituted or substituted with one or more R^b is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl; and/or

the C₃-C₆ cycloalkyl in the C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b is cyclopropyl, cyclobutyl, cyclopentyl or cyclohexyl; and/or

when R^b is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, R¹⁻¹¹ and R¹⁻¹², R¹⁻¹⁴ and R¹⁻¹⁵, R¹⁻²¹ and R¹⁻²², R¹⁻²³ and R¹⁻²⁴, or R¹⁻²⁵ and R¹⁻²⁶, independently together with the N atom to which they directly link, form a substituted C₃-C₉ heterocycloalkyl, a substituted C₃-C₉ heterocycloalkenyl or a substituted 5-10 membered heteroaryl, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2; and/or

when R²⁻¹ or R²⁻² is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R²⁻¹ or R²⁻² is an unsubstituted or substituted C₁-C₆ alkyl or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently a C₁-C₄ alkyl; and/or

when R²⁻¹ or R²⁻² is a substituted C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R²⁻¹ or R²⁻² is a substituted C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, and the substituent is a C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl in the substituent is independently a C₁-C₄ alkyl; and/or

when R²⁻¹ or R²⁻² is a substituted C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2; and/or

when R²⁻³, R²⁻⁴, R²⁻⁵ or R²⁻⁶ is an unsubstituted or substituted C₁-C₆ alkyl or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently a C₁-C₄ alkyl; and/or

when R²⁻³, R²⁻⁴, R²⁻⁵ or R²⁻⁶ is an unsubstituted or substituted C₃-C₈ cycloalkyl or an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, the C₃-C₈ cycloalkyl is a C₃-C₆ cycloalkyl; and/or

when R²⁻³, R²⁻⁴, R²⁻⁵ or R²⁻⁶ is a substituted C₁-C₆ alkyl, a substituted C₃-C₈ cycloalkyl, a substituted C₃-C₈ cycloalkyl-O- or a substituted C₁-C₆ alkyl-O-, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R²⁻³, R²⁻⁴, R²⁻⁵ or R²⁻⁶ is a substituted C₁-C₆ alkyl, a substituted C₃-C₈ cycloalkyl, a substituted C₃-C₈ cycloalkyl-O- or a substituted C₁-C₆ alkyl-O-, and the substituent is a C₁-C₆

alkyl or a substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl in the substituent is independently a C₁-C₄ alkyl, and preferably methyl or ethyl; and/or

when R²⁻³, R²⁻⁴, R²⁻⁵ or R²⁻⁶ is a substituted C₁-C₆ alkyl, a substituted C₃-C₈ cycloalkyl, a substituted C₃-C₈ cycloalkyl-O- or a substituted C₁-C₆ alkyl-O-, and the substituent is a C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, the number of the substituents is independently 1, 2, 3 or 4, and preferably 1 or 2; and/or

when R²⁻⁷ is independently halogen, R²⁻⁷ is independently a halogen-substituted C₁-C₆ alkyl, or R²⁻⁷ is independently a halogen-substituted C₁-C₆ alkyl-O-, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R²⁻⁷ is independently an unsubstituted or substituted C₁-C₆ alkyl, or R²⁻⁷ is independently an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl; and/or

when R²⁻⁸ or R²⁻⁹ is independently a halogen-substituted C₁-C₆ alkyl, the halogen is independently fluorine, chlorine, bromine or iodine;

when R²⁻⁸ or R²⁻⁹ is independently an unsubstituted or halogen-substituted C₁-C₆ alkyl, the C₁-C₆ alkyl is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl; and/or

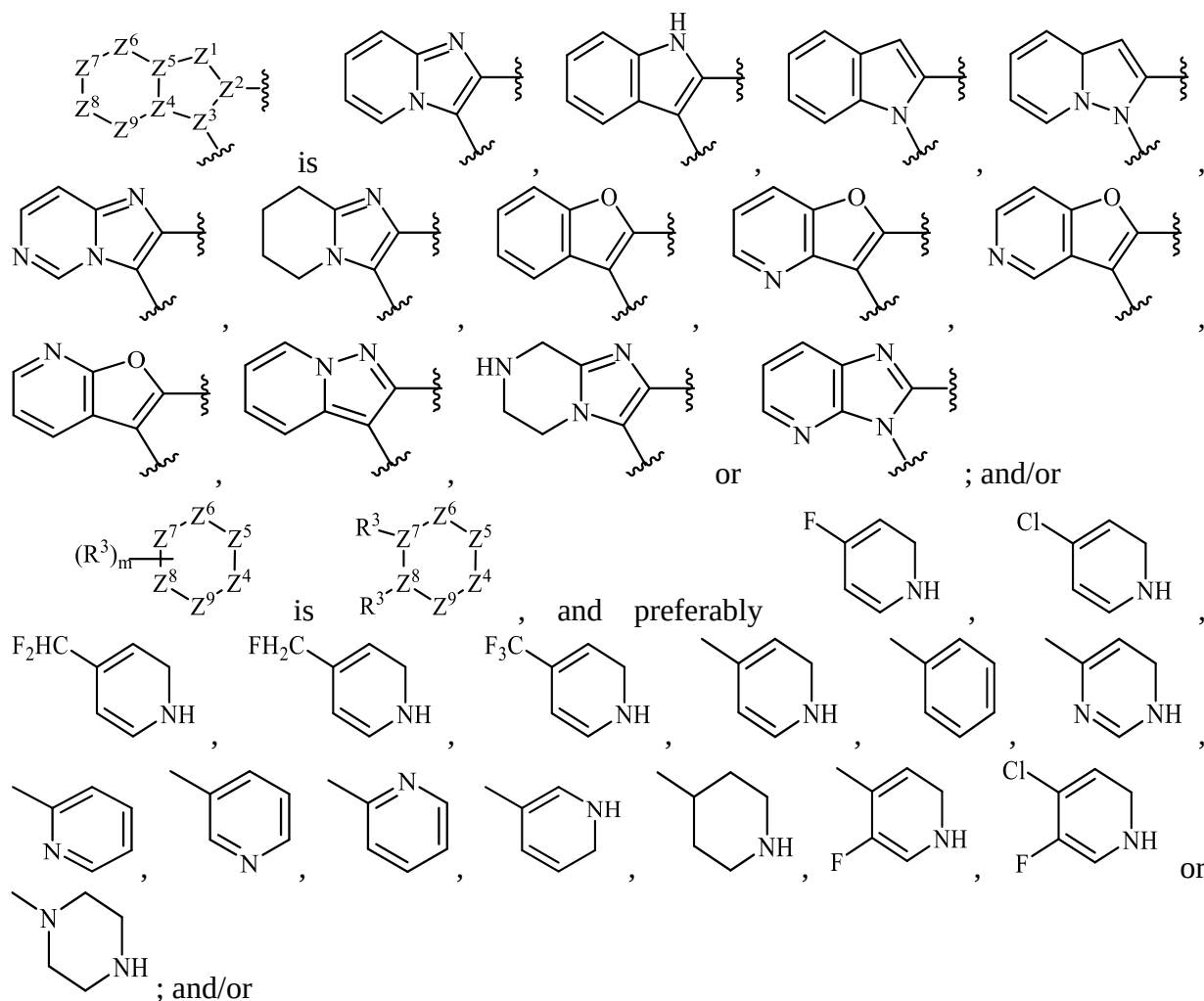
when R² is independently an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl is independently pyridinyl; and/or

when R² is independently a substituted 5-6 membered heteroaryl that may be substituted with halogen, a halogen-substituted C₁-C₆ alkyl or a halogen-substituted C₁-C₆ alkyl-O-, the halogen in the substituents is independently fluorine, chlorine, bromine or iodine; and/or

when R² is independently a substituted 5-6 membered heteroaryl that may be substituted with an unsubstituted or halogen-substituted C₁-C₆ alkyl, or an unsubstituted or halogen-substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl in the substituents is independently methyl, ethyl, *n*-propyl, isopropyl, *n*-butyl, isobutyl, *sec*-butyl or *tert*-butyl.

3. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 2, wherein when R¹ is a substituted C₁-C₆ alkyl or a substituted C₁-C₆ alkyl-O-, and the substituents are each independently halogen, the C₁-C₆ alkyl in the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is independently -CF₃; and/or

when R³ is a C₁-C₆ haloalkyl, the C₁-C₆ haloalkyl is -CF₃, -CH₂F or -CHF₂; and/or



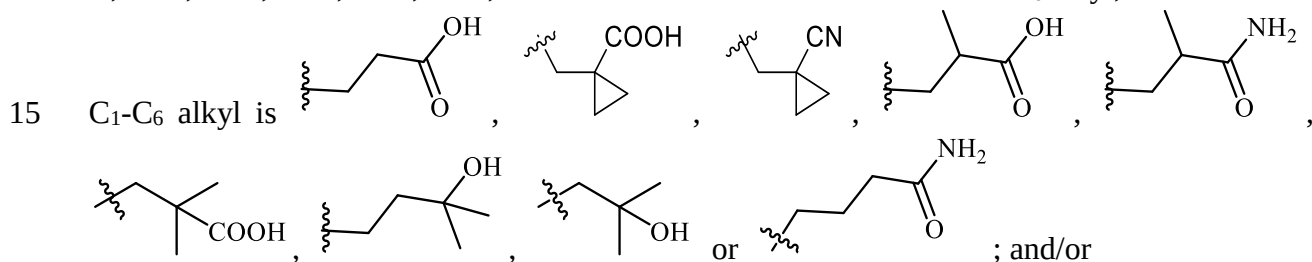
at least one R^1 is located at position α of the linkage between and Z^2 , and preferably,

when n is 2, each R^1 is independently located at position α of the linkage between and

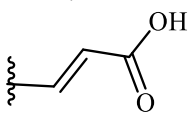
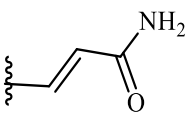
10 Z^2 ; or at least one R^1 is located at position β of the linkage between and Z^2 , and preferably, when n is 2, each R^1 is independently located at position β of the linkage between

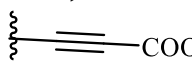
and Z^2 ; and/or

when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} , R^{1-13} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each a substituted C_1 - C_6 alkyl, the substituted

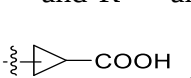
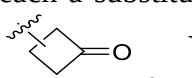
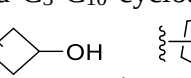
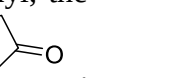
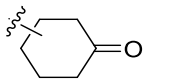
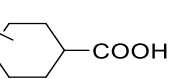
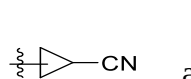
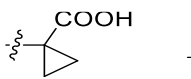
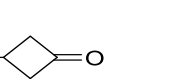
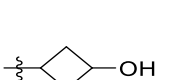
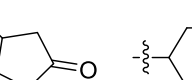
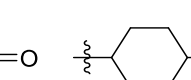
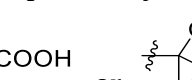


when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each a substituted C_2-C_6 alkenyl, the

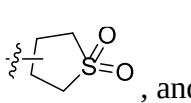
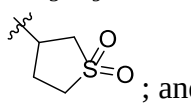
substituted C_2-C_6 alkenyl is  or  ; and/or

5 when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each a substituted C_2-C_6 alkynyl, the substituted C_2-C_6 alkynyl is  ; and/or

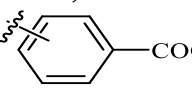
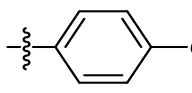
when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each a substituted C_3-C_{10} cycloalkyl, the

10 substituted C_3-C_{10} cycloalkyl is , , , , ,  or , and preferably , , , , ,  or  ; and/or

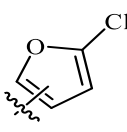
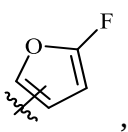
when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each a substituted C_3-C_9 heterocycloalkyl,

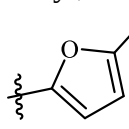
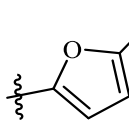
the substituted C_3-C_9 heterocycloalkyl is , and preferably  ; and/or

15 when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each an unsubstituted or substituted phenyl,

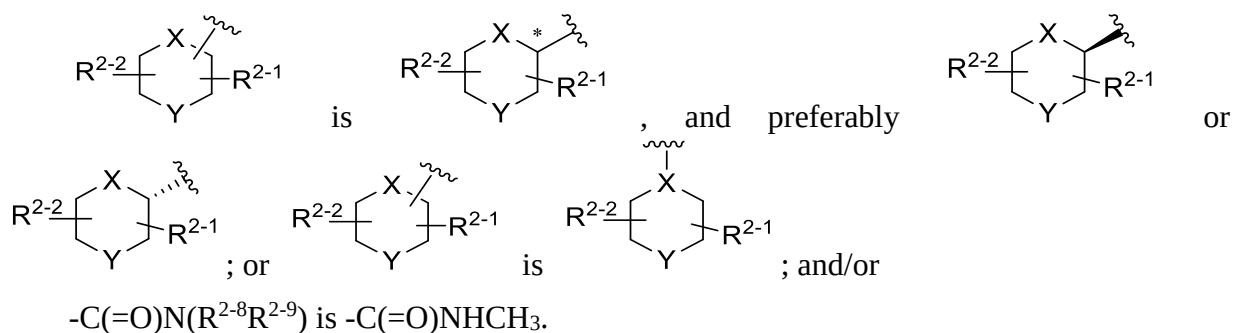
the substituted phenyl is , and preferably  ; and/or

when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}, R^{1-13}, R^{1-14}, R^{1-15}, R^{1-16}, R^{1-21}, R^{1-22}, R^{1-23}, R^{1-24}, R^{1-25}, R^{1-26}$ and R^{1-27} are each an unsubstituted or substituted 5-10

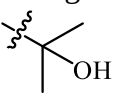
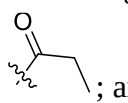
20 membered heteroaryl, the substituted 5-10 membered heteroaryl is  or  ,

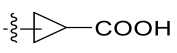
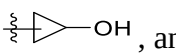
and preferably  or  ; and/or

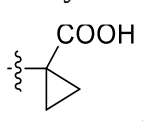
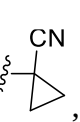
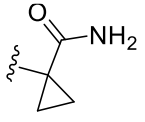
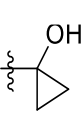
$-C(=O)N(R^{2-5}R^{2-6})$ is $-C(=O)NH_2$; and/or

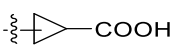
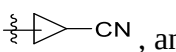


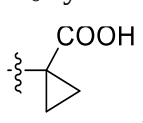
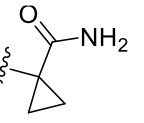
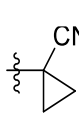
4. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 3, wherein the C₁-C₆

alkyl substituted with one or more R^a is  or  ; and/or

the C₃-C₆ cycloalkyl substituted with one or more R^a is  or  , and

preferably  ,  ,  or  ; and/or

the C₃-C₆ cycloalkyl substituted with one or more R^d is  or  , and

preferably  ,  or  ; and/or

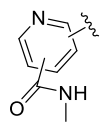
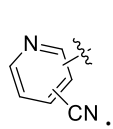
$-\text{N}(\text{R}^{1-21}\text{R}^{1-22})$ is $-\text{NHCH}_3$ or $-\text{NH}_2$; and/or

$-\text{C}(=\text{O})-\text{N}(\text{R}^{1-23}\text{R}^{1-24})$ is $-\text{C}(=\text{O})-\text{NH}_2$ or $-\text{C}(=\text{O})-\text{NHCH}_3$; and/or

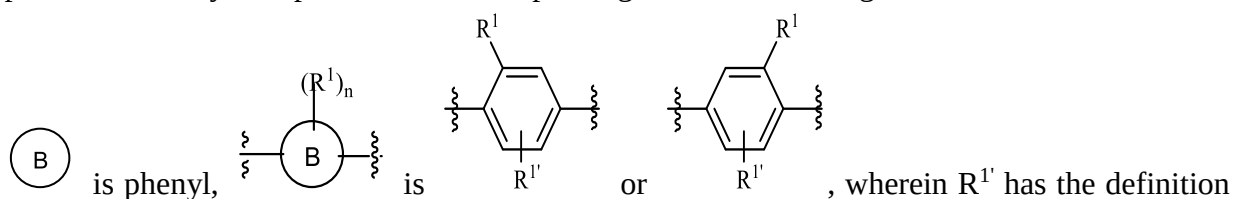
$-\text{S}(=\text{O})_2-\text{N}(\text{R}^{1-25}\text{R}^{1-26})$ is $-\text{S}(=\text{O})_2-\text{NH}_2$; and/or

$-\text{C}(=\text{O})-\text{O}-\text{R}^{1-27}$ is $-\text{COOH}$, $-\text{COOC}_2\text{H}_5$ or $-\text{COOCH}_3$; and/or

when R² is independently an unsubstituted or substituted 5-6 membered heteroaryl, the

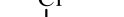
substituted 5-6 membered heteroaryl is independently  or  .

5. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 4, wherein when











when the  is a 5-6 membered heteroaryl, the  is ,

5

 ,  ,  ,  ,  ,  or  , and

preferably











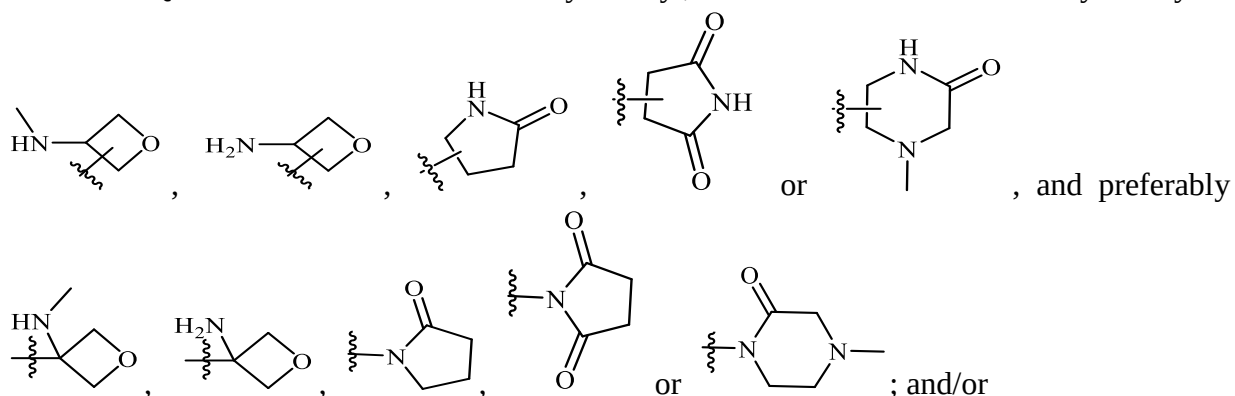



 or
 
 ; and/or

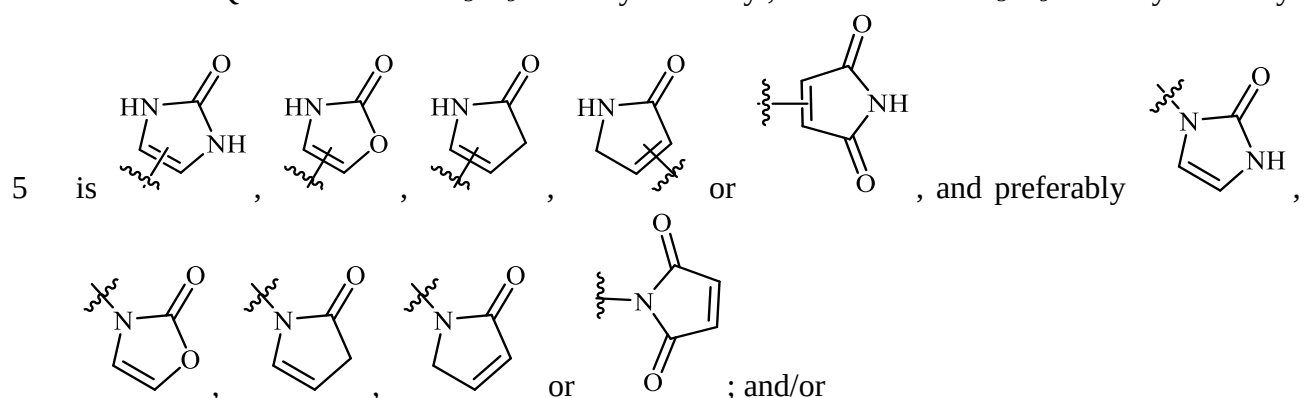
[illegible]



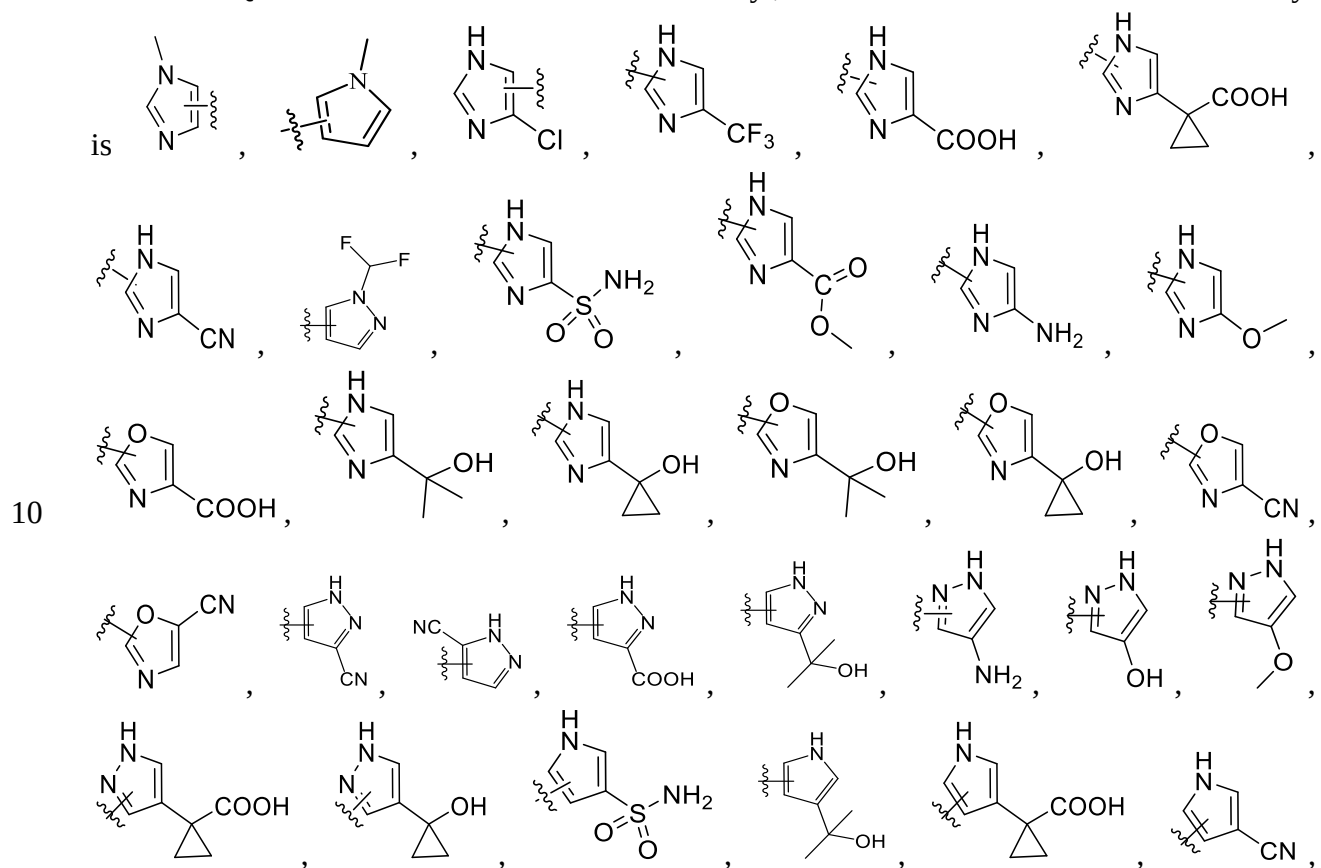
when Q is a substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkyl is

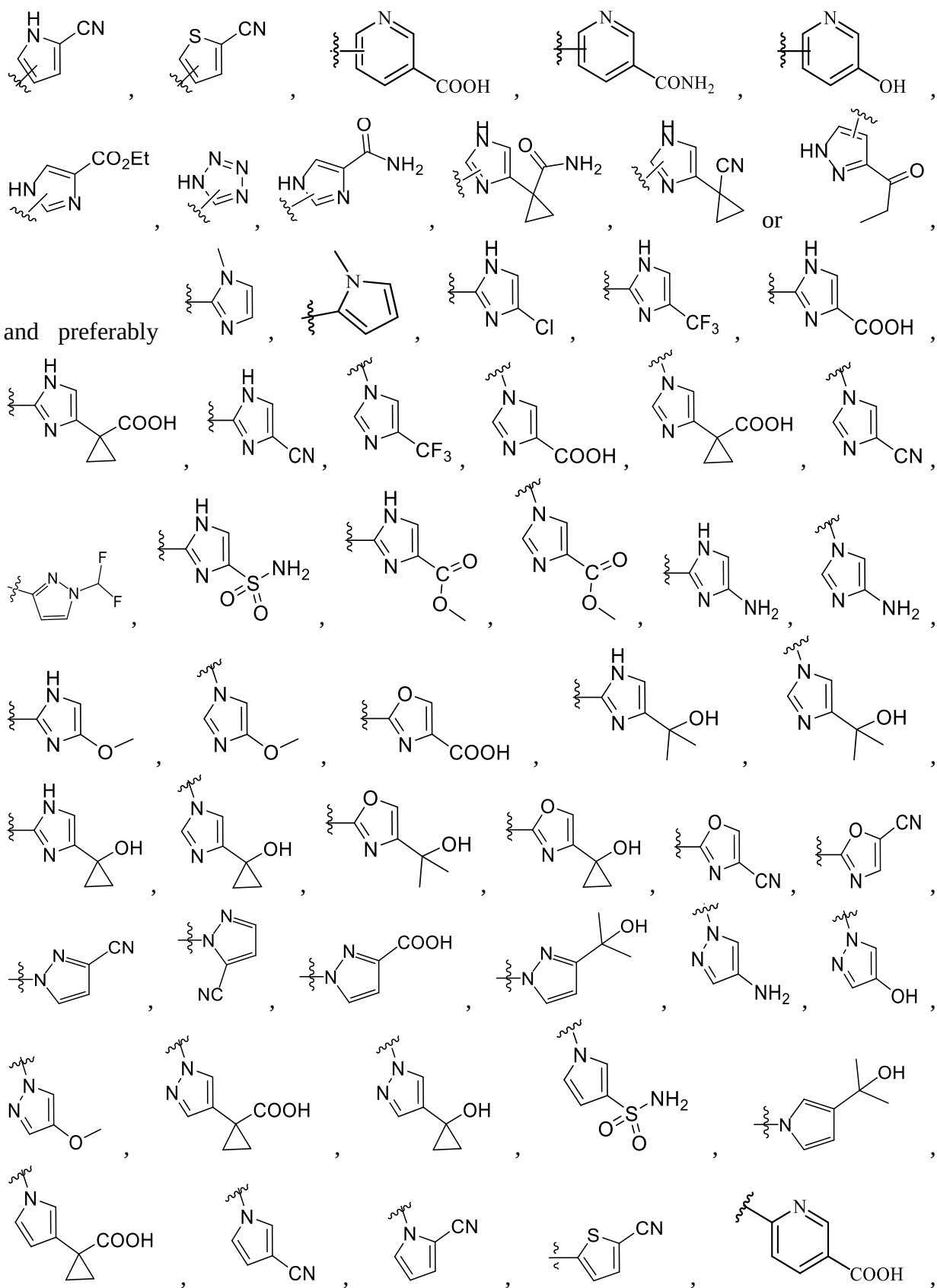


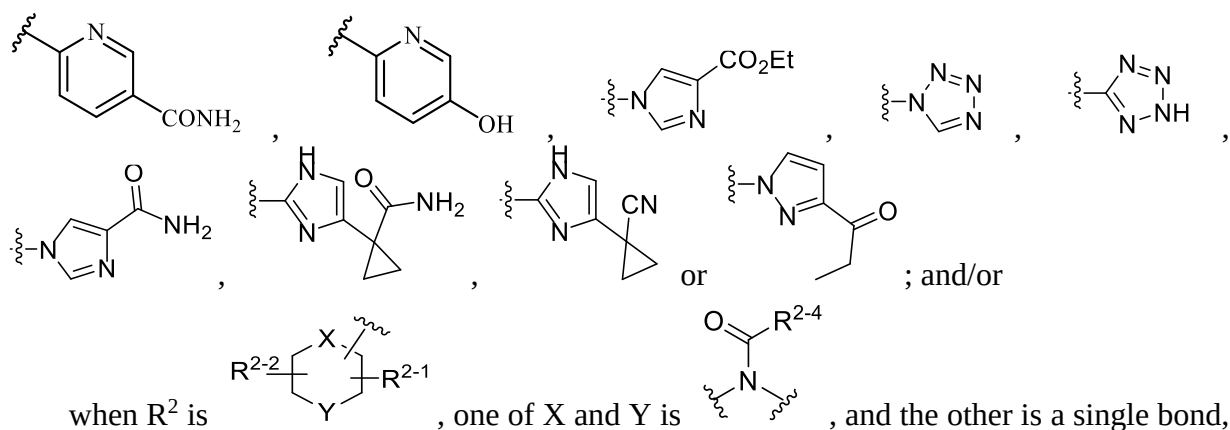
when Q is a substituted C₃-C₉ heterocycloalkenyl, the substituted C₃-C₉ heterocycloalkenyl

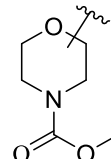
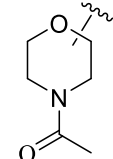


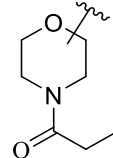
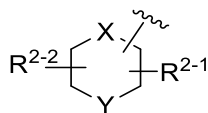
when Q is a substituted 5-6 membered heteroaryl, the substituted 5-10 membered heteroaryl

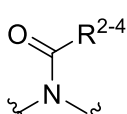
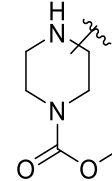


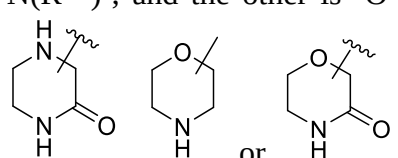




methylene or -O-; preferably X is -O-; and more preferably R² is ,  or

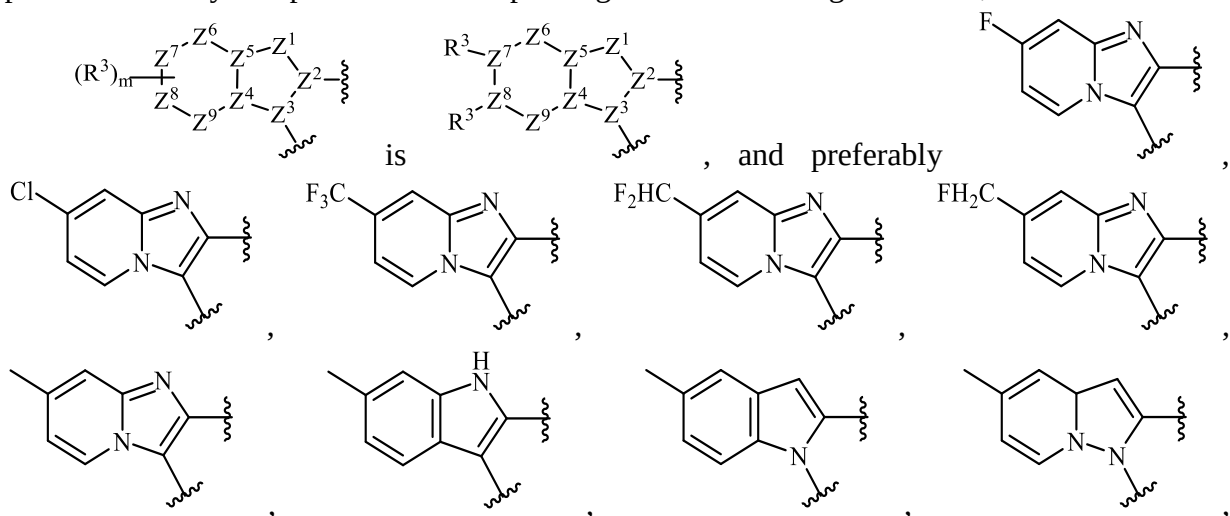
5 ; or when R² is , X or Y is -N(R²⁻³)-; preferably one of X and Y is -

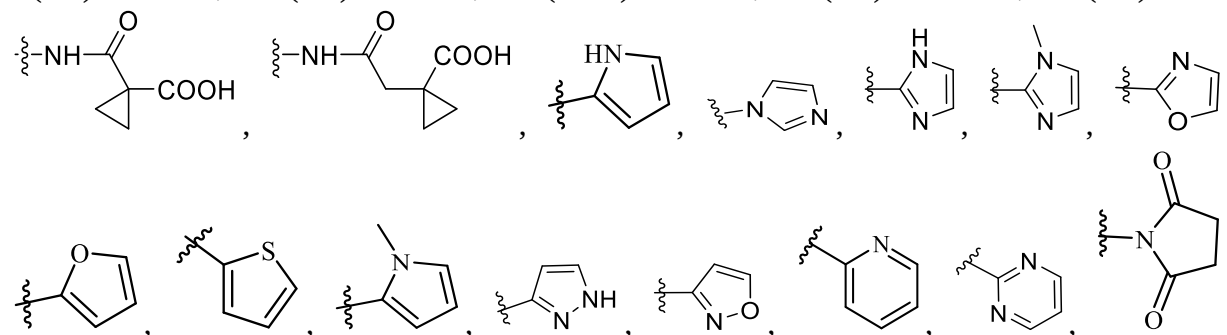
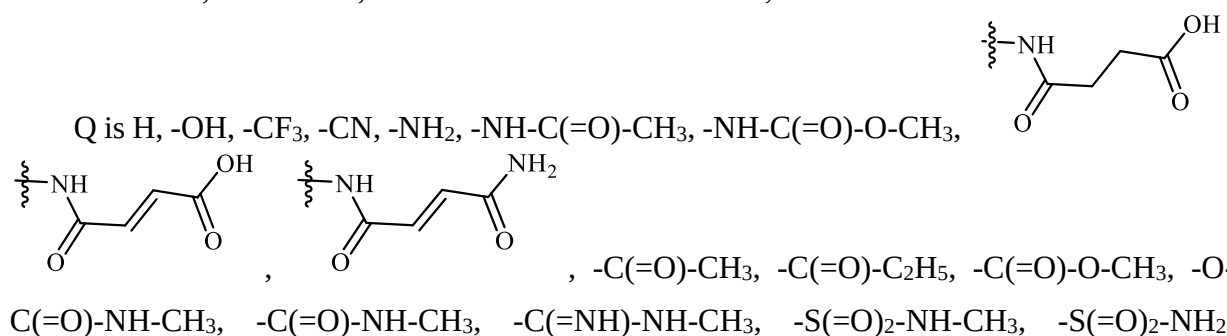
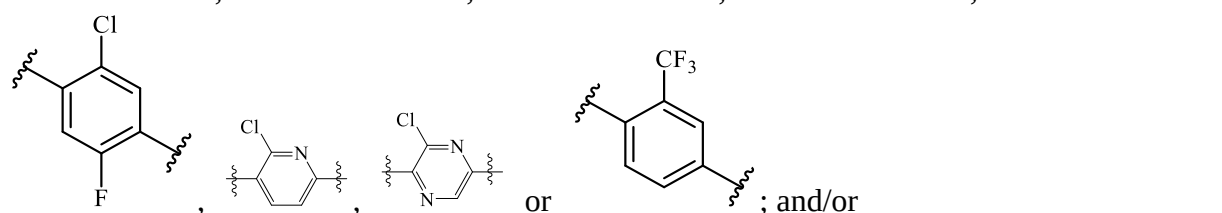
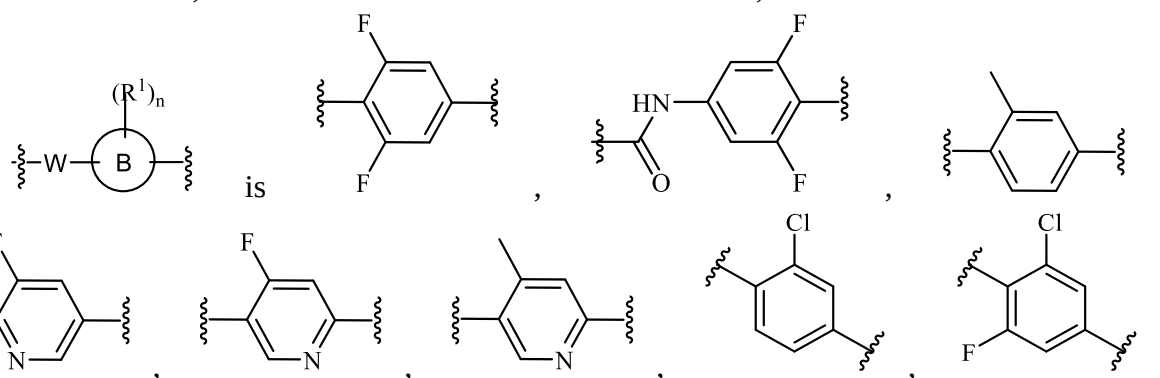
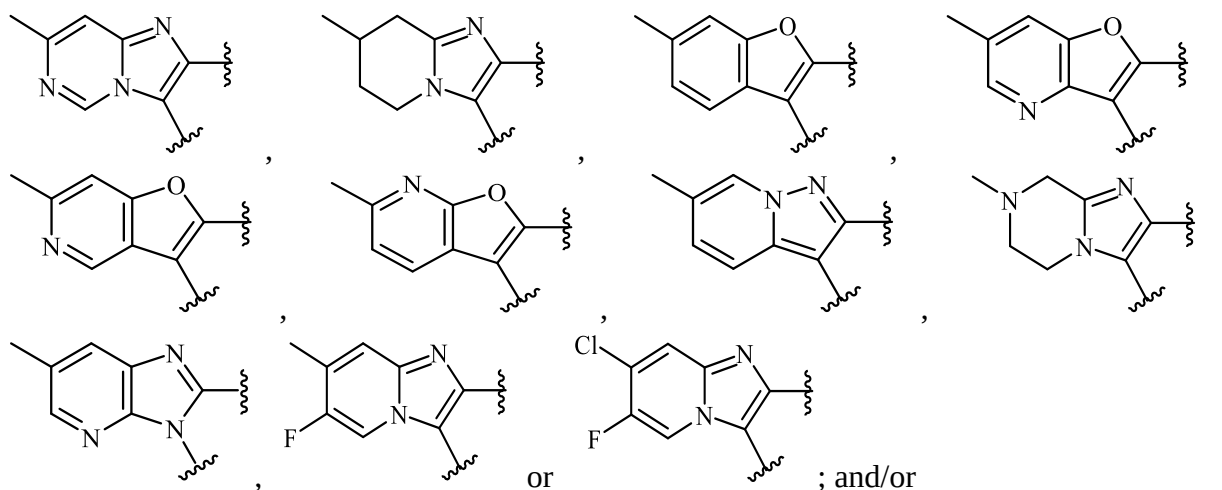
N(R²⁻³)-, and the other is -O-, -N(R²⁻³)- or ; and more preferably R² is ,

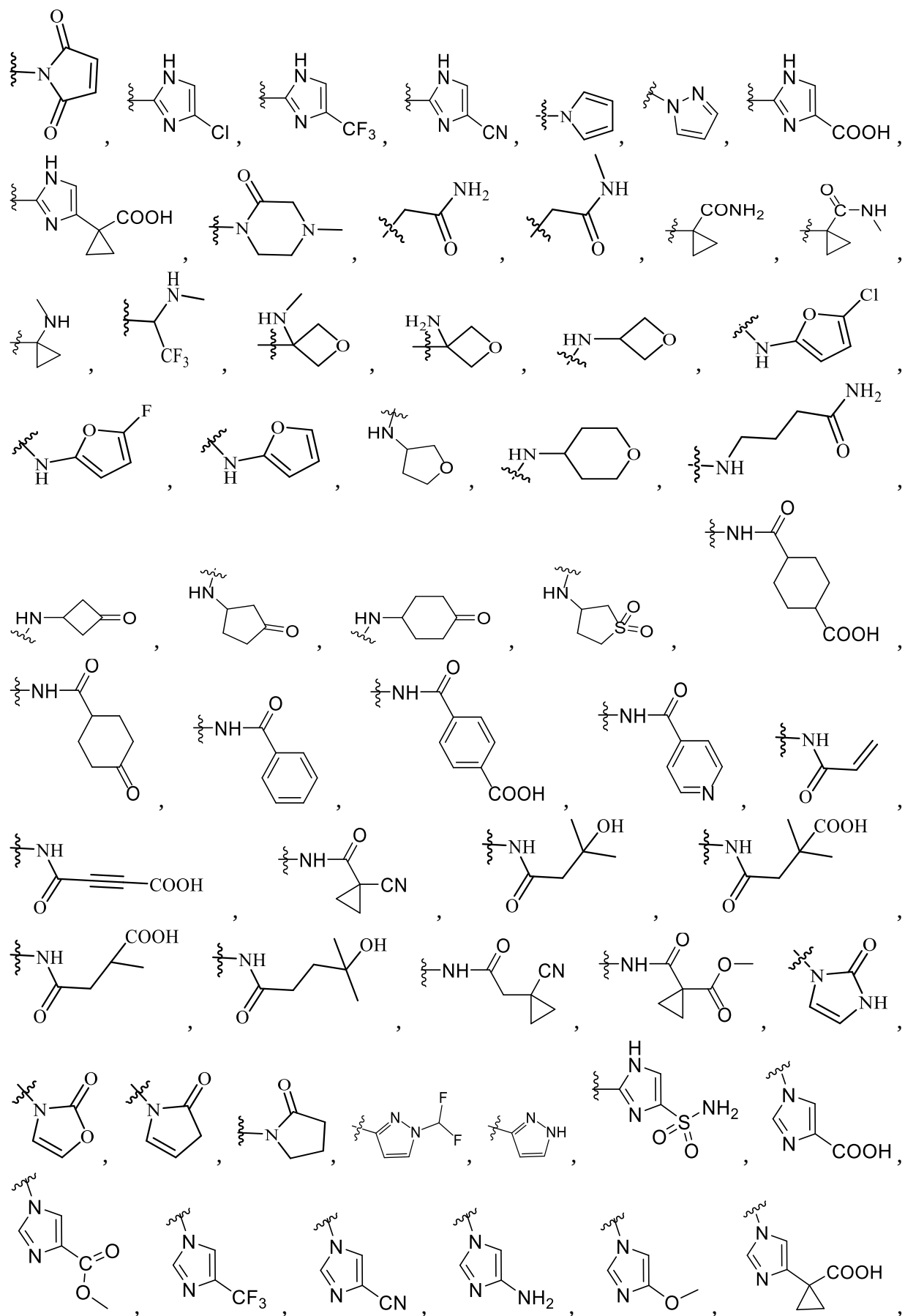


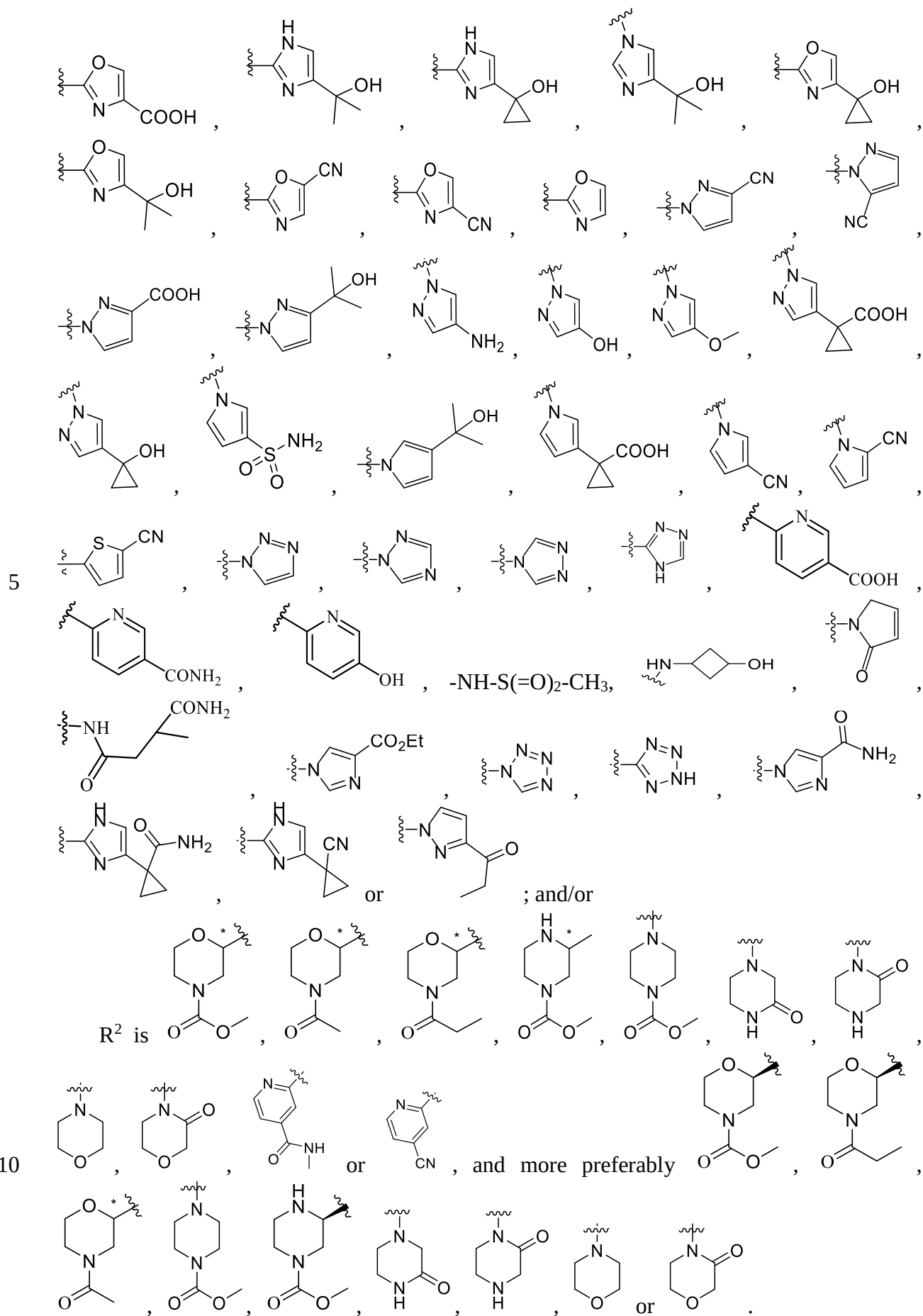
6. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the

10 pharmaceutically acceptable salt or the prodrug thereof according to claim 5, wherein









7. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein R^3 is halogen, $-\text{CHF}_2$, $-\text{CH}_2\text{F}$ or a C_1 - C_6 alkyl;

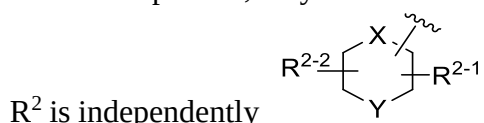
5 Q is $-\text{NH}-\text{C}(=\text{O})-\text{R}^{1-1}$, $-\text{N}(\text{R}^{1-14}\text{R}^{1-15})$, $-\text{NH}-\text{S}(=\text{O})_2-\text{R}^{1-16}$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and
10 the substituted C_1 - C_6 alkyl, the substituted C_3 - C_{10} cycloalkyl, the substituted C_3 - C_9 heterocycloalkyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: $-\text{N}(\text{R}^{1-21}\text{R}^{1-22})$, $-\text{C}(=\text{O})-\text{N}(\text{R}^{1-23}\text{R}^{1-24})$, $-\text{S}(=\text{O})_2-\text{N}(\text{R}^{1-25}\text{R}^{1-26})$ or $-\text{C}(=\text{O})-\text{O}-\text{R}^{1-27}$, wherein when a plurality of substituents are present, they are the same or different;

15 R^{1-1} , R^{1-14} , R^{1-15} , R^{1-16} , R^{1-21} , R^{1-22} , R^{1-23} , R^{1-24} , R^{1-25} , R^{1-26} and R^{1-27} are each independently H, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_2 - C_6 alkynyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and
20 the substituted C_1 - C_6 alkyl, the substituted C_2 - C_6 alkynyl, the substituted C_3 - C_9 heterocycloalkyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, $-\text{OH}$, $-\text{CN}$, $=\text{O}$, $-\text{COOH}$, $-\text{C}(=\text{O})\text{NH}_2$, $-\text{COOCH}_3$, $-\text{NH}_2$, and a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or a C_3 - C_6 cycloalkyl that is unsubstituted
25 or substituted with one or more R^d , wherein when a plurality of R^d are present, they are the same or different, and each R^d is independently H, halogen, $-\text{OH}$, $=\text{O}$, $-\text{CF}_3$, $-\text{CN}$, $-\text{COOH}$ or $-\text{C}(=\text{O})\text{NH}_2$; and when a plurality of substituents are present, they are the same or different; or

R^3 is halogen, a C_1 - C_6 haloalkyl or a C_1 - C_6 alkyl;

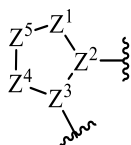
30 Q is $-\text{NH}-\text{C}(=\text{O})-\text{R}^{1-1}$, $-\text{C}(=\text{O})-\text{N}(\text{R}^{1-5}\text{R}^{1-6})$, $-\text{S}(=\text{O})_2-\text{N}(\text{R}^{1-9}\text{R}^{1-10})$, $-\text{N}(\text{R}^{1-14}\text{R}^{1-15})$, $-\text{NH}-\text{S}(=\text{O})_2-\text{R}^{1-16}$, an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one

or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: -OH, -CF₃, -CN, -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH or -COOH; and when a plurality of substituents are present, they are the same or different;

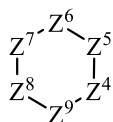


8. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein

the heterocyclic compound of formula I is embodiment I or embodiment II,

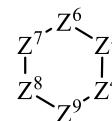


embodiment I: is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl, wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl

contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



$Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8$ and Z^9 independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z^6 or Z^9 represents a single bond;

m is 1, 2, 3 or 4;

R^3 is H, halogen, -OH, a C_1 - C_6 haloalkyl, -CN or a C_1 - C_6 alkyl; when m is not 1, R^3 is independently the same or different; or when two adjacent R^3 are present, they, together with the ring atom to which they directly link, form a C_3 - C_6 cycloalkyl, a C_3 - C_6 cycloalkenyl, a C_3 - C_9 heterocycloalkyl, a C_3 - C_9 heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl, wherein the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C_3 - C_9 heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C_3 - C_9 heterocycloalkyl, wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the C_3 - C_9 heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

n is 1, 2, 3 or 4;

R^1 is independently H, halogen, -OH, -CN, an unsubstituted or substituted C_1 - C_6 alkyl, or an unsubstituted or substituted C_1 - C_6 alkyl-O-; wherein the substituted C_1 - C_6 alkyl or the substituted C_1 - C_6 alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C_1 - C_6 alkyl, a C_1 - C_6 alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); when a plurality of substituents are present, they are the same or different; and when n is not 1, R^1 is independently the same or different;

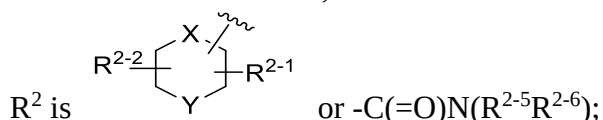
Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)- R^{1-1} , -C(=O)- R^{1-2} , -O-C(=O)-N(R^{1-3} R^{1-4}), -C(=O)-N(R^{1-5} R^{1-6}), -C(=NH)-N(R^{1-7} R^{1-8}), -S(=O)₂-N(R^{1-9} R^{1-10}), -S(=O)-N(R^{1-11} R^{1-12}), -O-C(=O)- R^{1-13} , -N(R^{1-14} R^{1-15}), -NH-S(=O)₂- R^{1-16} , an unsubstituted or substituted C_1 - C_6 alkyl, an unsubstituted or substituted C_1 - C_6 alkyl-O-, an unsubstituted or substituted C_2 - C_6 alkenyl, an unsubstituted or substituted C_2 - C_6 alkenyl-O-, an unsubstituted or substituted C_3 - C_{10} cycloalkyl, an unsubstituted or substituted C_3 - C_9 heterocycloalkyl, an unsubstituted or substituted C_3 - C_9

heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₃-C₁₀ cycloalkyl, the substituted C₆-C₁₀ aryl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), -N(R¹⁻²¹R¹⁻²²), -C(=O)-N(R¹⁻²³R¹⁻²⁴), -S(=O)₂-N(R¹⁻²⁵R¹⁻²⁶), -C(=O)-O-R¹⁻²⁷, and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

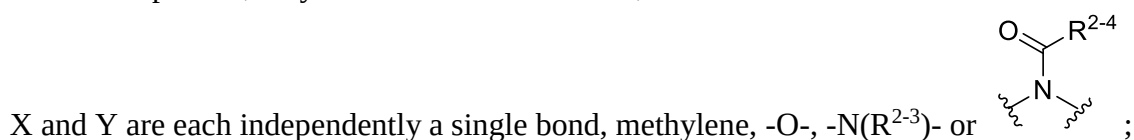
R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹², R¹⁻¹³, R¹⁻¹⁴, R¹⁻¹⁵, R¹⁻¹⁶, R¹⁻²¹, R¹⁻²², R¹⁻²³, R¹⁻²⁴, R¹⁻²⁵, R¹⁻²⁶ and R¹⁻²⁷ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₂-C₆ alkynyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C₃-C₉ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₉ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-10 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₂-C₆ alkynyl, the substituted C₃-C₁₀ cycloalkyl, the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl, the substituted phenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -COOCH₃, -NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^d, wherein when a plurality of R^d are present, they are the same

or different, and each R^d is independently H, halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different; or

R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , R^{1-11} and R^{1-12} , R^{1-14} and R^{1-15} , R^{1-21} and R^{1-22} , R^{1-23} and R^{1-24} , or R^{1-25} and R^{1-26} , independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, wherein the C₃-C₉ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C₃-C₉ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the 5-10 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C₃-C₉ heterocycloalkyl, the substituted C₃-C₉ heterocycloalkenyl or the substituted 5-10 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^b , wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

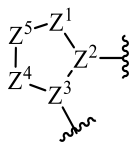


R^{2-1} and R^{2-2} are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;

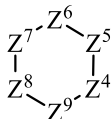


R^{2-3} , R^{2-4} , R^{2-5} and R^{2-6} are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆

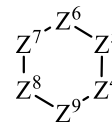
alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and when a plurality of substituents are present, they are the same or different;



embodiment II: is a 5 membered heteroaryl or a 5 membered heterocycloalkenyl, wherein the 5 membered heteroaryl contains 1-2 heteroatoms selected from one or more of N, O and S, and the 5 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S;



is phenyl, a 5-6 membered heterocycloalkyl, a 5-6 membered heterocycloalkenyl, or a 5-6 membered heteroaryl, wherein the 5-6 membered heterocycloalkyl contains 1-3 heteroatoms selected from one or more of N, O and S, the 5-6 membered heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of N, O and S, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S;



Z¹, Z², Z³, Z⁴, Z⁵, Z⁶, Z⁷, Z⁸ and Z⁹ independently represent a ring atom; and when is a 5 membered heterocycloalkyl, a 5 membered heterocycloalkenyl or a 5 membered heteroaryl, Z⁶ or Z⁹ represents a single bond;

m is 1, 2, 3 or 4;

R³ is H, halogen, -OH, -CF₃, -CN or a C₁-C₆ alkyl; when m is not 1, R³ is the same or different; or when two adjacent R³ are present, they, together with the ring atom to which they directly link, form a C₃-C₆ cycloalkyl, a C₃-C₆ cycloalkenyl, a C₃-C₅ heterocycloalkyl, a C₃-C₅ heterocycloalkenyl, phenyl or a 5-6 membered heteroaryl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of O, S and N;

W is a single bond or -C(=O)-NH-;



is phenyl, a 5-6 membered heteroaryl, or a C₃-C₅ heterocycloalkyl, wherein the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S, and the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N;

n is 1, 2, 3 or 4;

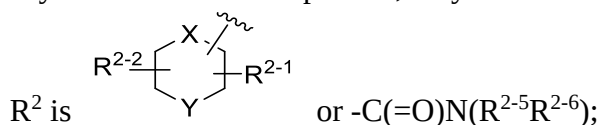
R¹ is independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O; when a plurality of substituents are present, they are the same or different; and when n is not 1, R¹ is the same or different;

Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N, and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₆-C₁₀ aryl, the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;

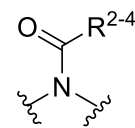
R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² and R¹⁻¹³ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₃-C₆ cycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, an unsubstituted or substituted phenyl, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein the C₃-C₅ heterocycloalkyl contains 1-3 heteroatoms selected from one or more of O, S and N, the C₃-C₅ heterocycloalkenyl contains 1-3 heteroatoms selected from one or more of O, S and N,

and the 5-6 membered heteroaryl contains 1-3 heteroatoms selected from one or more of N, O and S; and the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₃-C₆ cycloalkyl, the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl, the substituted phenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, a C₃-C₆ cycloalkyl, -COOH or =O, wherein when a plurality of substituents are present, they are the same or different; or

R¹⁻³ and R¹⁻⁴, R¹⁻⁵ and R¹⁻⁶, R¹⁻⁷ and R¹⁻⁸, R¹⁻⁹ and R¹⁻¹⁰, or R¹⁻¹¹ and R¹⁻¹², independently together with the N atom to which they directly link, form an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl, wherein the C₃-C₅ heterocycloalkyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, the C₃-C₅ heterocycloalkenyl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom, and the 5-6 membered heteroaryl contains no heteroatom or 1-2 heteroatoms selected from one or more of O, S and N in addition to the existing N atom; and the substituted C₃-C₅ heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O-, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of R^b are present, they are the same or different, and each R^b is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different;



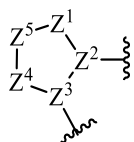
R²⁻¹ and R²⁻² are each independently H, halogen, -OH, -CN, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, or =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O); wherein the substituted C₁-C₆ alkyl or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O, and when a plurality of substituents are present, they are the same or different;



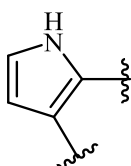
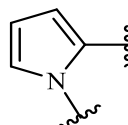
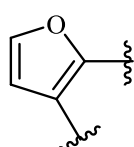
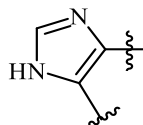
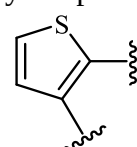
X and Y are each independently a single bond, methylene, -O-, -N(R²⁻³)- or

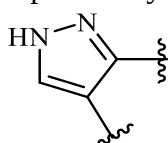
R²⁻³, R²⁻⁴, R²⁻⁵ and R²⁻⁶ are each independently H, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl, an unsubstituted or substituted C₃-C₈ cycloalkyl-O-, or an unsubstituted or substituted C₁-C₆ alkyl-O-; wherein the substituted C₁-C₆ alkyl, the substituted C₃-C₈ cycloalkyl, the substituted C₃-C₈ cycloalkyl-O- or the substituted C₁-C₆ alkyl-O- is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or =O, and when a plurality of substituents are present, they are the same or different.

9. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 8, wherein when



is a 5 membered heteroaryl, the 5 membered heteroaryl is pyrrolyl, furanyl, thienyl

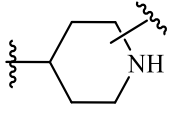
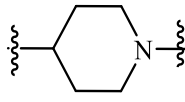
or imidazolyl, the pyrrolyl is preferably  or , the furanyl is preferably , the imidazolyl is preferably , the thienyl is preferably  and

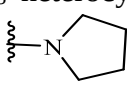
the pyrazolyl is preferably ; and/or

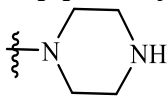
when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

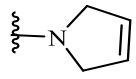
when two adjacent R³ are present and they, together with the ring atom to which they directly link, form a C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, and the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when  is a C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S, and the C₄-C₅ heterocycloalkyl is preferably piperidinyl, the

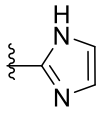
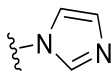
piperidinyl is preferably , and more preferably ; and/or

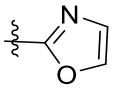
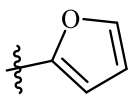
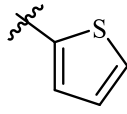
5 when Q is an unsubstituted or substituted C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and the C₃-C₅ heterocycloalkyl is preferably pyrrolidinyl or piperazinyl, the pyrrolidinyl is preferably , and the

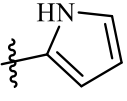
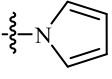
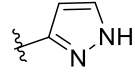
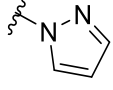
piperazinyl is preferably ; and/or

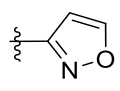
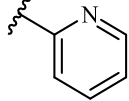
10 when Q is an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, wherein the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S, and the C₄-C₅ heterocycloalkenyl is preferably 2,5-dihydro-1H-pyrrolyl, and more preferably ; and/or

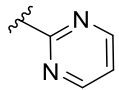
15 when Q is an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl is imidazolyl, oxazolyl, furanyl, thienyl, pyrrolyl, pyrazolyl, isoxazolyl, pyridinyl or

pyrimidinyl, the imidazolyl is preferably  or , the oxazolyl is preferably

, the furanyl is preferably , the thienyl is preferably , the pyrrolyl

is preferably  or , the pyrazolyl is preferably  or , the

20 isoxazolyl is preferably , the pyridinyl is preferably  and the pyrimidinyl is

preferably ; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered

heteroaryl, and the substituent is halogen, the halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₁-C₆ alkyl or a C₁-C₆ alkyl-O- substituted with one or more R^a, the C₁-C₆ alkyl in the substituent is independently a C₁-C₄ alkyl, and preferably methyl; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₃-C₆ cycloalkyl substituted with one or more R^a, the C₃-C₆ cycloalkyl in the substituent is cyclopropyl; and/or

when Q is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted C₆-C₁₀ aryl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, the substituent is a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl substituted with one or more R^a, and R^a is halogen, the halogen is independently fluorine, chlorine, bromine or iodine; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₁-C₆ alkyl, or an unsubstituted or substituted C₁-C₆ alkyl-O-, the C₁-C₆ alkyl is independently a C₁-C₄ alkyl, and preferably methyl or ethyl; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₃-C₅ heterocycloalkyl, the C₃-C₅ heterocycloalkyl is a C₄-C₅ heterocycloalkyl, wherein the C₄-C₅ heterocycloalkyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted C₃-C₅ heterocycloalkenyl, the C₃-C₅ heterocycloalkenyl is a C₄-C₅ heterocycloalkenyl, and the C₄-C₅ heterocycloalkenyl contains 1-2 heteroatoms selected from one or more of N, O and S; and/or

when R¹⁻¹, R¹⁻², R¹⁻³, R¹⁻⁴, R¹⁻⁵, R¹⁻⁶, R¹⁻⁷, R¹⁻⁸, R¹⁻⁹, R¹⁻¹⁰, R¹⁻¹¹, R¹⁻¹² or R¹⁻¹³ is an unsubstituted or substituted 5-6 membered heteroaryl, the 5-6 membered heteroaryl contains 1-2 heteroatoms selected from one or two of N, O and S; and/or

when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted phenyl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is halogen, the
5 halogen is independently fluorine, chlorine, bromine or iodine, and preferably fluorine; and/or

when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted phenyl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₁-C₆ alkyl
10 or a C₁-C₆ alkyl-O-, the C₁-C₆ alkyl in the substituent is independently a C₁-C₄ alkyl; and/or

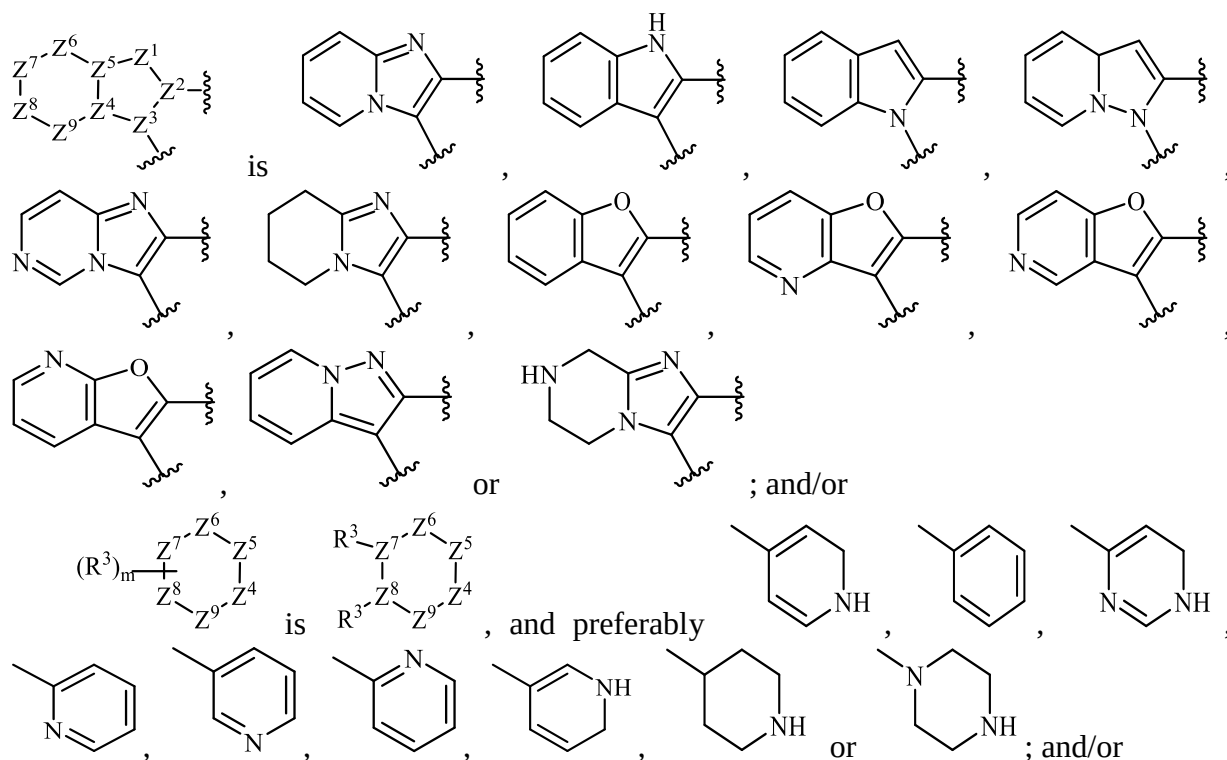
when R^{1-1} , R^{1-2} , R^{1-3} , R^{1-4} , R^{1-5} , R^{1-6} , R^{1-7} , R^{1-8} , R^{1-9} , R^{1-10} , R^{1-11} , R^{1-12} or R^{1-13} is a substituted C₁-C₆ alkyl, a substituted C₁-C₆ alkyl-O-, a substituted C₂-C₆ alkenyl, a substituted C₂-C₆ alkenyl-O-, a substituted phenyl, a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₃-C₆
15 cycloalkyl, the C₃-C₆ cycloalkyl in the substituent is cyclopropyl; and/or

when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₁-C₆ alkyl or a C₁-C₆ alkyl-O- substituted with one or more
20 R^b, the C₁-C₆ alkyl in the substituent is independently a C₁-C₄ alkyl; and/or

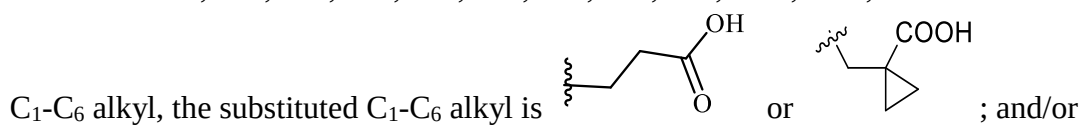
when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, and the substituent is a C₃-C₆ cycloalkyl substituted with one or more R^b, the C₃-C₆
25 cycloalkyl in the substituent is cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl; and/or

when R^{1-3} and R^{1-4} , R^{1-5} and R^{1-6} , R^{1-7} and R^{1-8} , R^{1-9} and R^{1-10} , or R^{1-11} and R^{1-12} , independently together with the N atom to which they directly link, form a substituted C₃-C₅ heterocycloalkyl, a substituted C₃-C₅ heterocycloalkenyl, or a substituted 5-6 membered heteroaryl, the substituent is a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl substituted
30 with one or more R^b, wherein when R^b is halogen, the halogen is independently fluorine, chlorine, bromine or iodine.

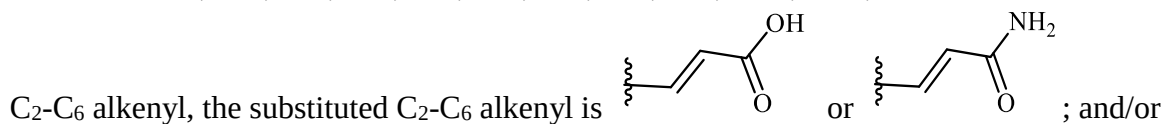
10. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 9, wherein



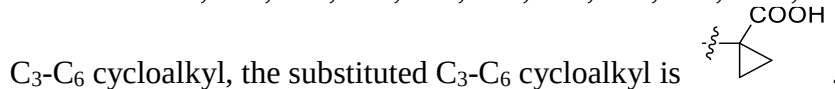
when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}$ or R^{1-13} is a substituted



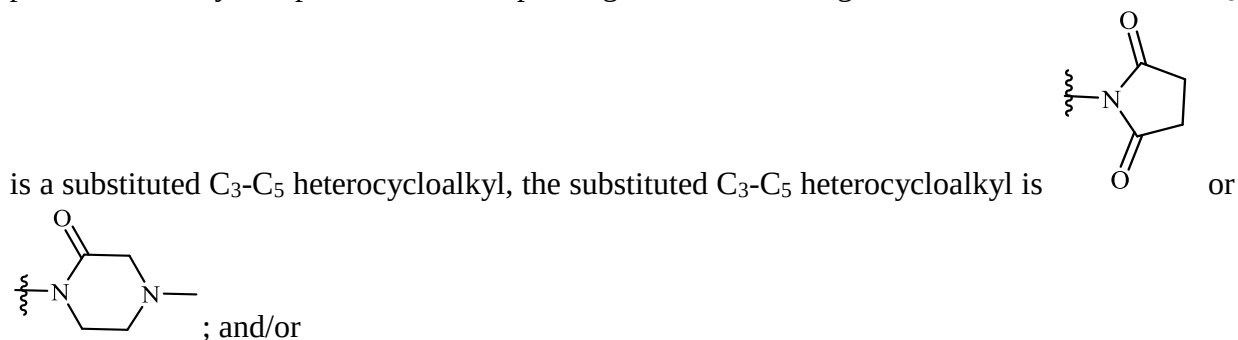
when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}$ or R^{1-13} is a substituted



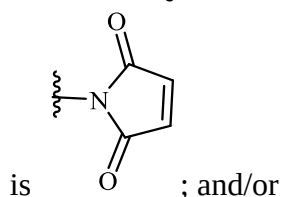
when $R^{1-1}, R^{1-2}, R^{1-3}, R^{1-4}, R^{1-5}, R^{1-6}, R^{1-7}, R^{1-8}, R^{1-9}, R^{1-10}, R^{1-11}, R^{1-12}$ or R^{1-13} is a substituted



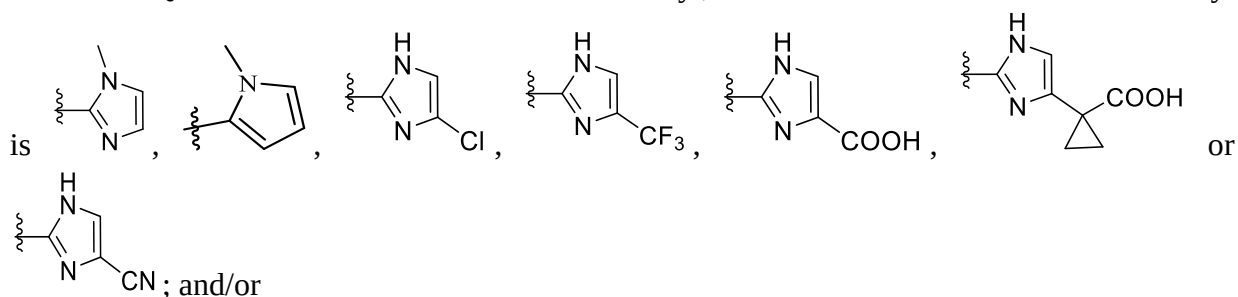
11. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 10, wherein when Q



when Q is a substituted C₃-C₅ heterocycloalkenyl, the substituted C₃-C₅ heterocycloalkenyl

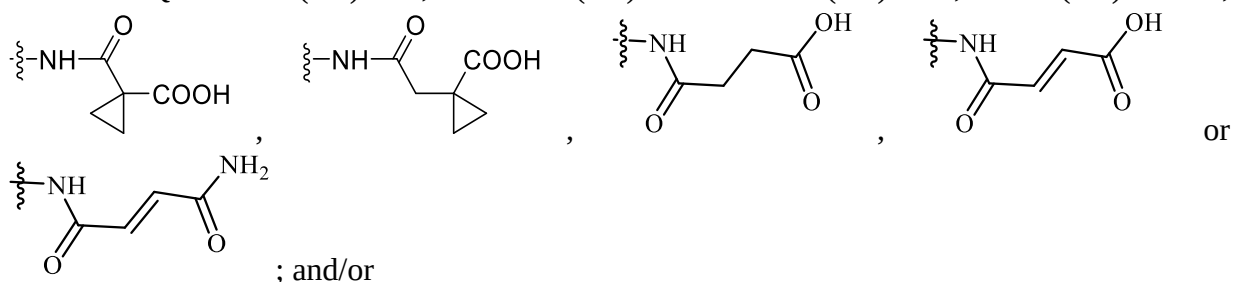


when Q is a substituted 5-6 membered heteroaryl, the substituted 5-6 membered heteroaryl

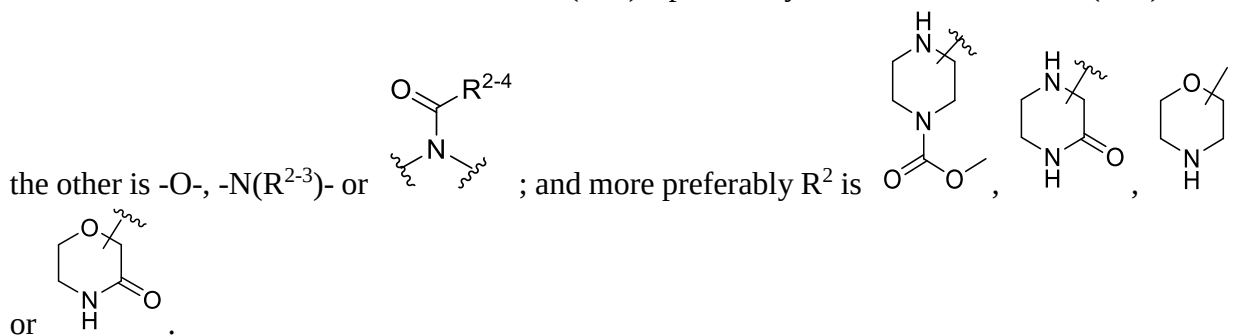
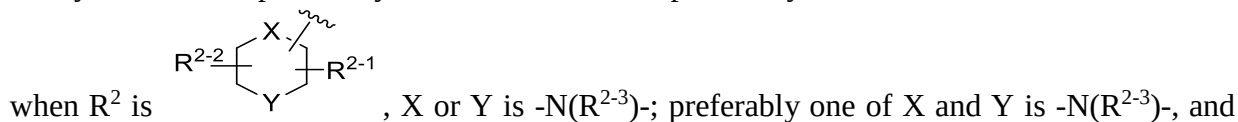
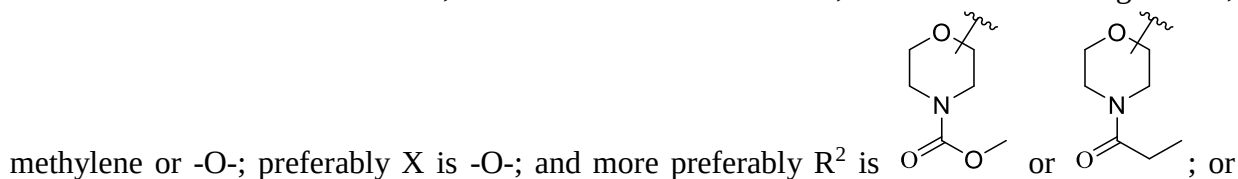
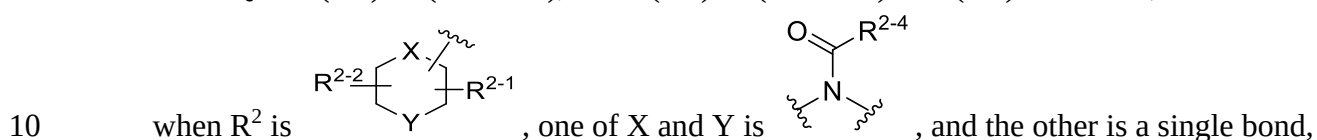


5

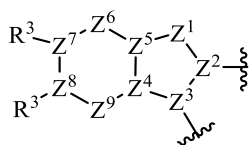
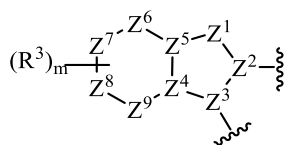
when Q is -NH-C(=O)-R¹⁻¹, the -NH-C(=O)-R¹⁻¹ is -NH-C(=O)-CH₃, -NH-C(=O)-O-CH₃,



when Q is -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), the -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰) is -S(=O)₂-NH-CH₃; and/or

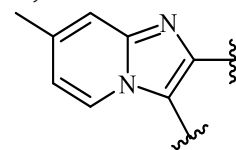


12. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 11, wherein

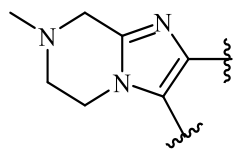
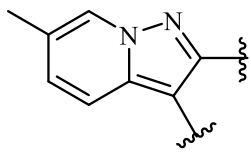
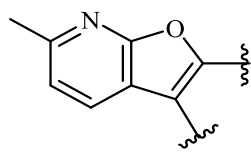
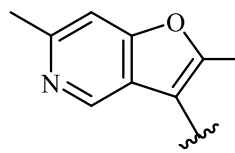
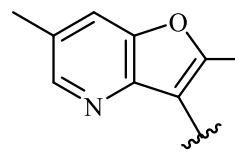
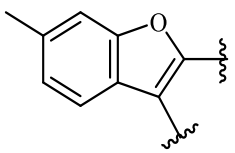
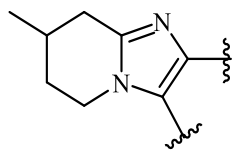
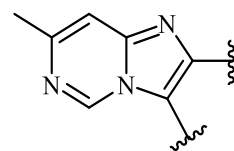
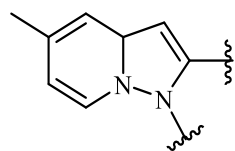
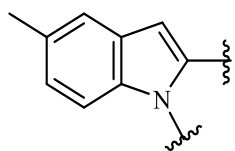
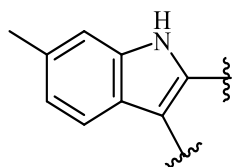


is

, and preferably

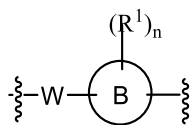


5

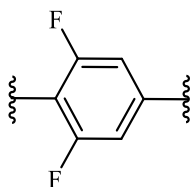


or

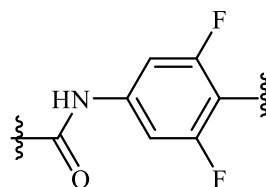
; and/or



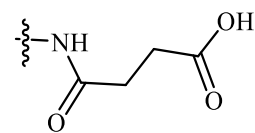
is



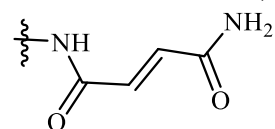
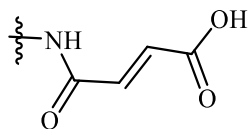
or



; and/or



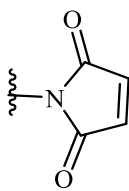
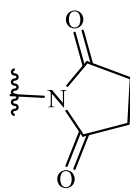
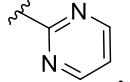
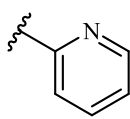
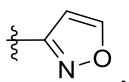
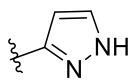
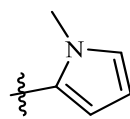
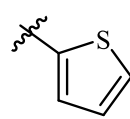
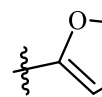
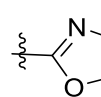
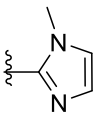
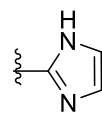
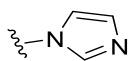
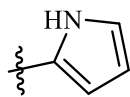
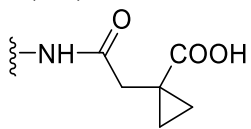
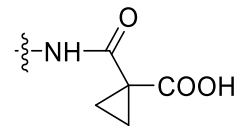
Q is H, -OH, -CF₃, -CN, -NH-C(=O)-CH₃, -NH-C(=O)-O-CH₃,

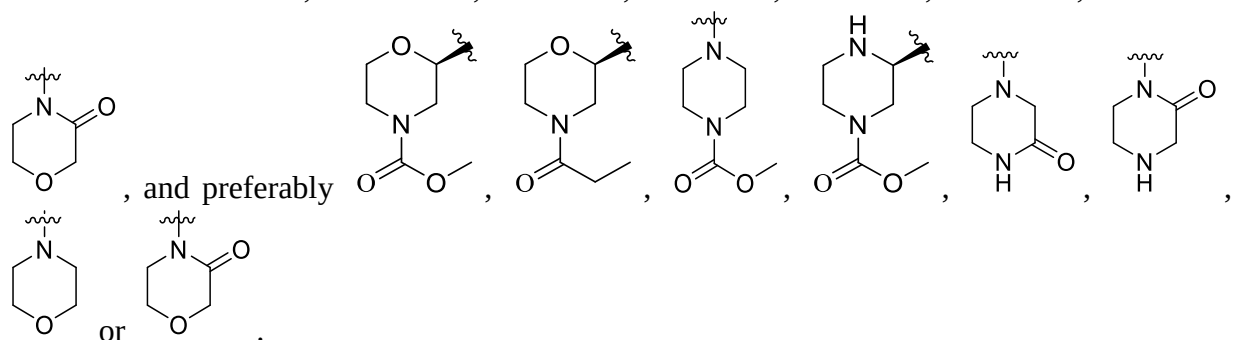
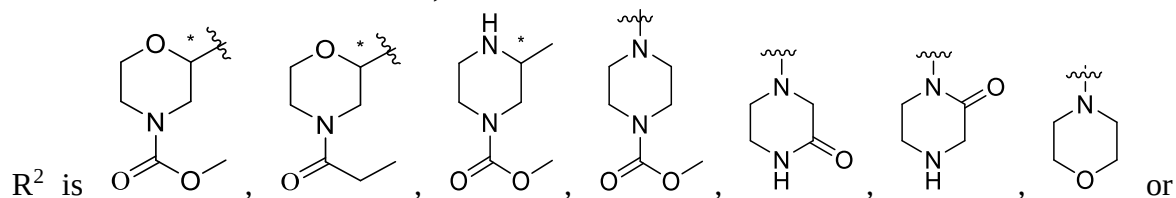
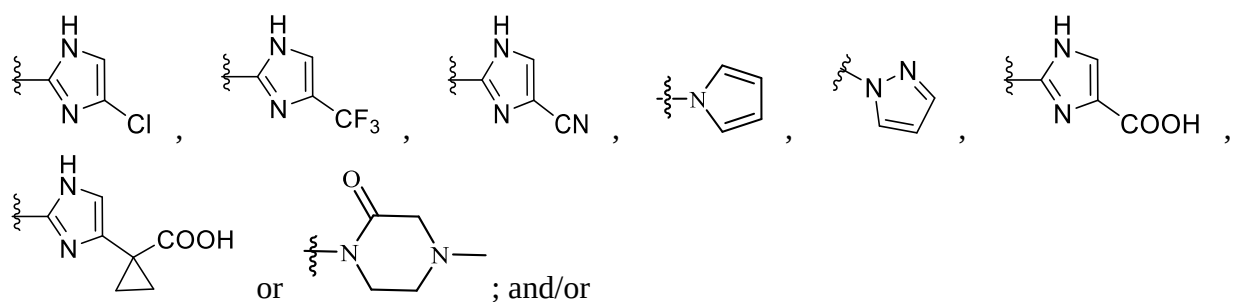


, -C(=O)-CH₃, -C(=O)-C₂H₅, -C(=O)-O-CH₃, -O-

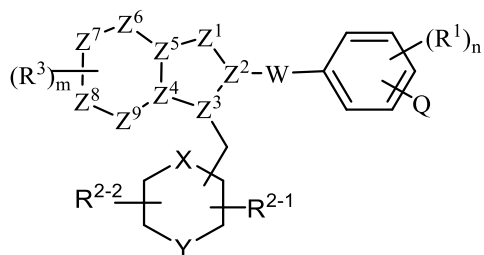
10

C(=O)-NH-CH₃, -C(=O)-NH-CH₃, -C(=NH)-NH-CH₃, -S(=O)₂-NH-CH₃,

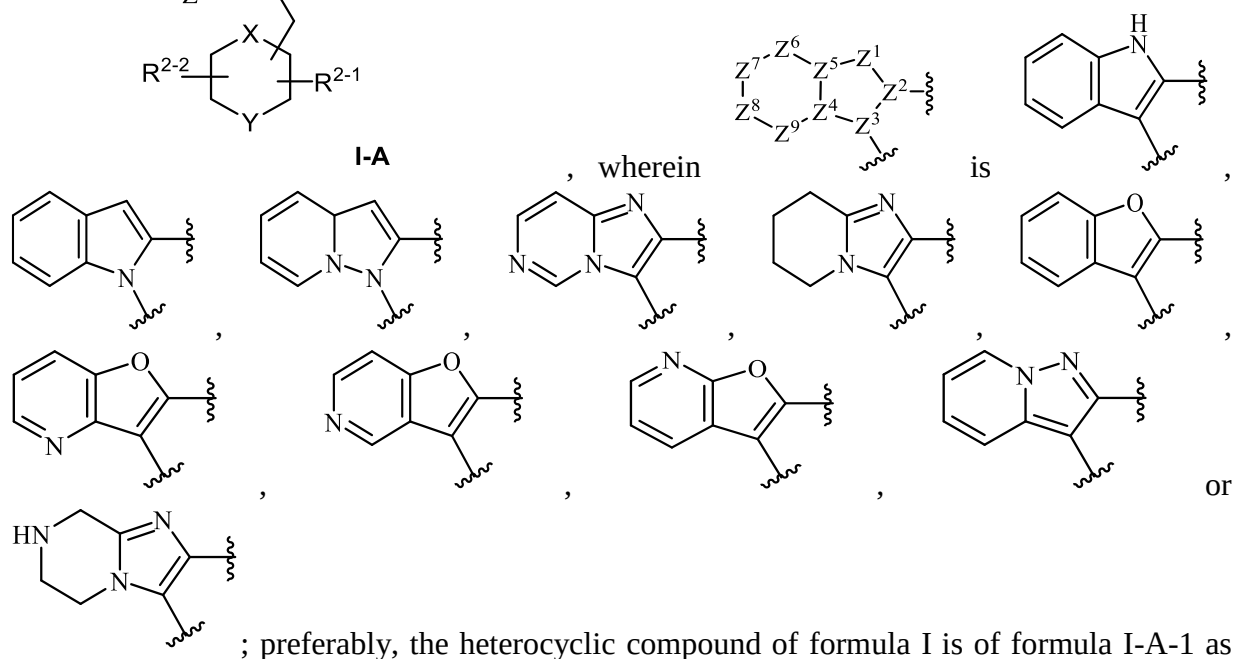


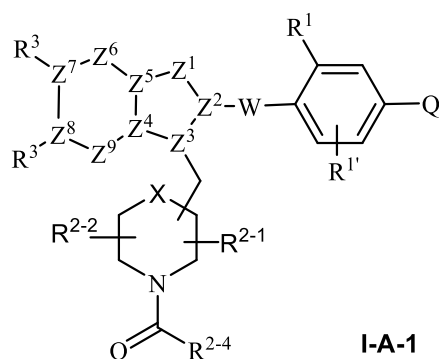


13. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to any one of claims 1, 7 and 8, wherein the heterocyclic compound of formula I is of formula I-A as shown below:

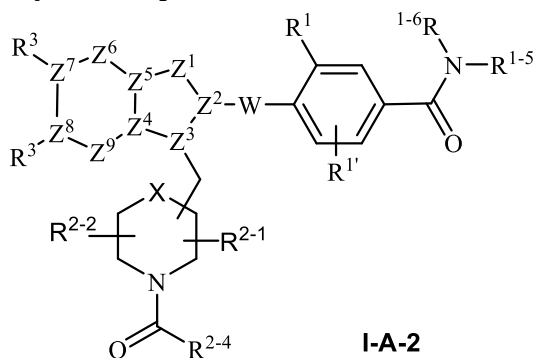


I-A





shown below: **I-A-1**, wherein $R^{1'}$ has the definition as described for R^1 , $R^{1'}$ is the same as or different from R^1 , and W is a single bond; and more preferably, the heterocyclic compound of formula I is of formula I-A-2 as shown below:

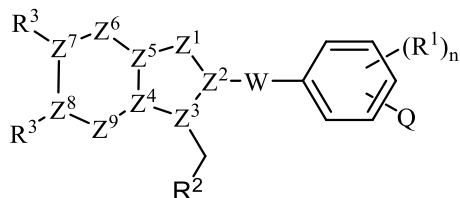


I-A-2

, wherein X is a single bond, methylene or -O-;

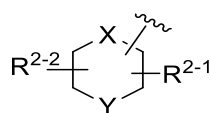
5 or

the heterocyclic compound of formula I is of formula I-B as shown below:



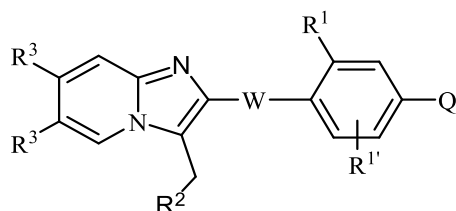
I-B

, wherein R^2 is



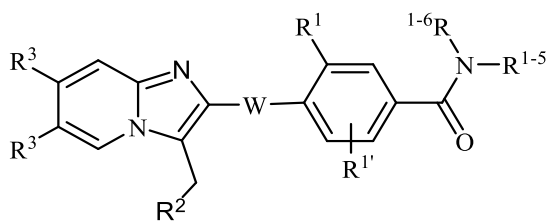
or $-C(=O)N(R^{2-5}R^{2-6})$, and

X or Y is $-N(R^{2-3})$ -; preferably, the heterocyclic compound of formula I is of formula I-B-1 as



I-B-1

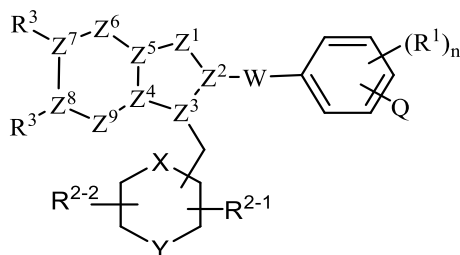
shown below: , wherein $R^{1'}$ has the definition as described for R^1 , $R^{1'}$ is the same as or different from R^1 , and W is a single bond; and more preferably, the heterocyclic compound of formula I is of formula I-B-2 as shown below:



I-B-2

; or

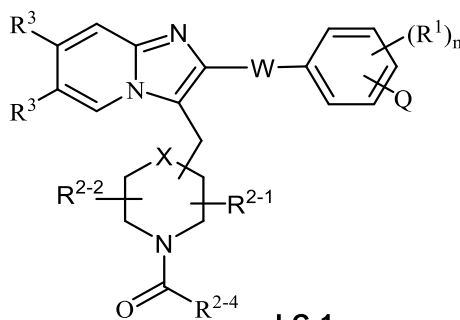
the heterocyclic compound of formula I is of formula I-C as shown below:



I-C

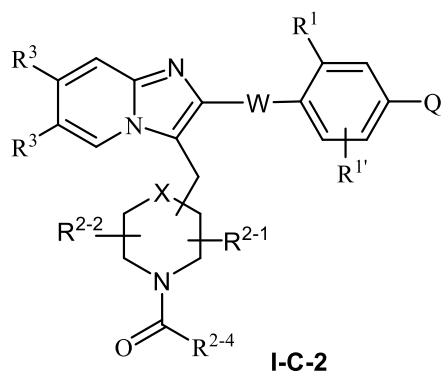
, wherein Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)-R¹⁻¹,
¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-
 5 N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted
 C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-
 C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₃-
 C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or
 substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an
 10 unsubstituted or substituted 5-10 membered heteroaryl;

preferably, the heterocyclic compound of formula I is of formula I-C-1 as shown below:



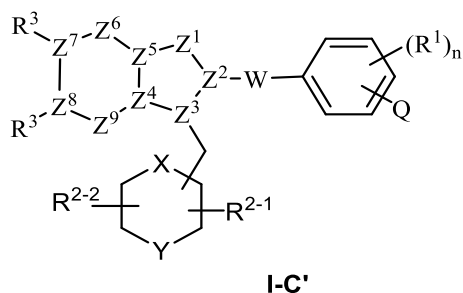
I-C-1

; and more preferably, the heterocyclic compound of
 formula I is of formula I-C-2 as shown below:



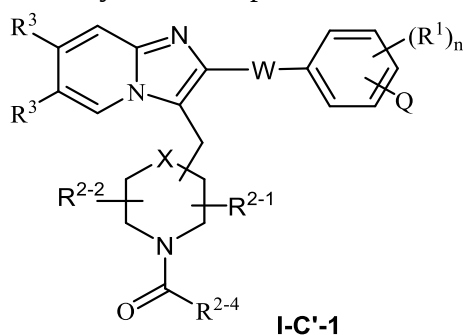
, wherein R^{1'} has the definition as described for R¹, R^{1'} is the same as or different from R¹, and X is a single bond, methylene or -O-; or

the heterocyclic compound of formula I is of formula I-C' as shown below:

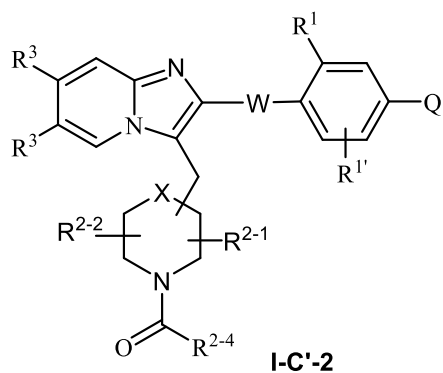


, wherein Q is H, halogen, -OH, -CF₃, -CN, -NH₂, -NH-

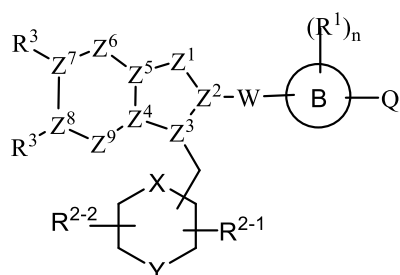
- 5 C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₆-C₁₀ aryl, an unsubstituted or substituted C₃-C₅ heterocycloalkyl, an unsubstituted or substituted C₃-C₅
- 10 heterocycloalkenyl, or an unsubstituted or substituted 5-6 membered heteroaryl; preferably, the heterocyclic compound of formula I is of formula I-C'-1 as shown below:



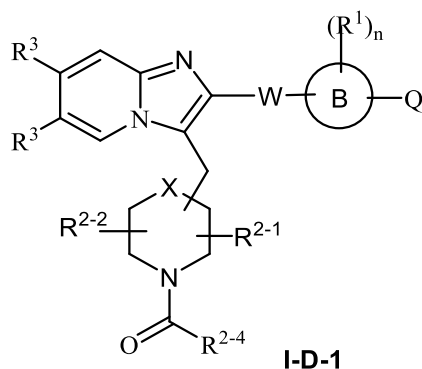
; and more preferably, the heterocyclic compound of formula I is of formula I-C'-2 as shown below:



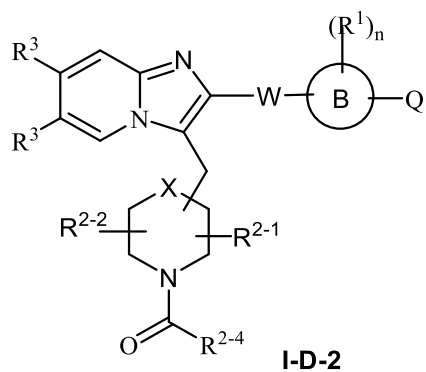
, wherein $R^{1'}$ has the definition as described for R^1 , and $R^{1'}$ is the same as or different from R^1 ; or the heterocyclic compound of formula I is of formula



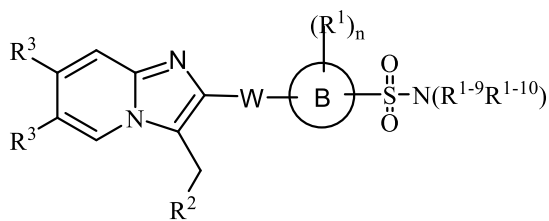
I-D as shown below: , wherein $\textcircled{B}$ is a 5-6 membered heteroaryl or a C_3 - C_5 heterocycloalkyl; preferably, the heterocyclic compound of formula I is of



5 formula I-D-1 as shown below: ; and more preferably, the heterocyclic compound of formula I is of formula I-D-2 as shown below:

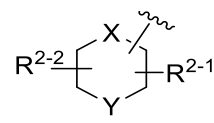


, wherein X is a single bond, methylene or -O-, and W is a single bond; or the heterocyclic compound of formula I is of formula I-E as shown below:

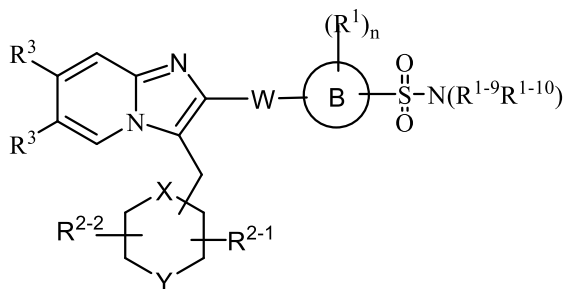


I-E

; wherein R² is independently



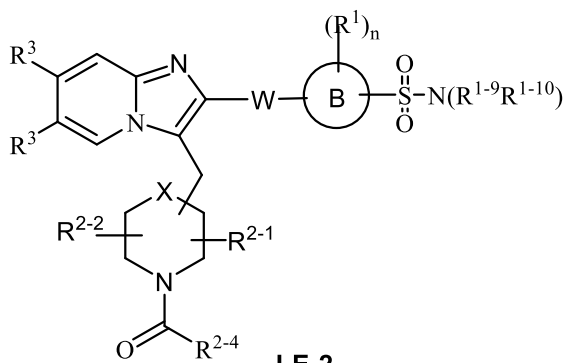
or an unsubstituted or substituted 5-6 membered heteroaryl; for example, W is a single bond; preferably, the heterocyclic compound of formula I is of formula I-E-1 as shown below:



I-E-1

; for example,  is phenyl or 6 membered

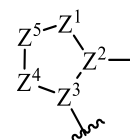
5 heteroaryl; and more preferably, the heterocyclic compound of formula I is of formula I-E-2 as shown below:



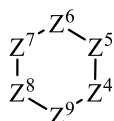
I-E-2

; for example,  is phenyl.

14. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the



10 pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein is a 5 membered heteroaryl; and/or



is phenyl, a 6 membered heterocycloalkyl, a 6 membered heterocycloalkenyl, or a 6 membered heteroaryl; and/or

m is 1 or 2; and/or

R³ is H, halogen, -OH, a C₁-C₆ haloalkyl or a C₁-C₆ alkyl; and/or

W is a single bond; and/or

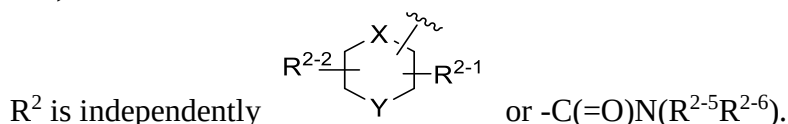
n is 1 or 2; and/or

- 5 R¹ is independently H, halogen, -OH, -CN or an unsubstituted or substituted C₁-C₆ alkyl; and/or

the substituent in R¹ is independently one or more selected from the following: halogen, -OH, -CN or -C(=O)NH₂; and/or

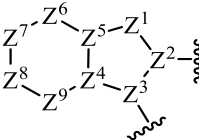
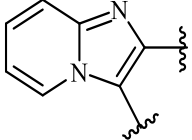
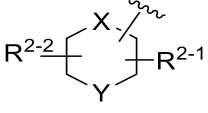
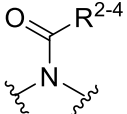
- Q is H, halogen, -OH, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an
10 unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl, and preferably H, halogen, -OH, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=O)-N(R¹⁻⁵R¹⁻⁶), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), an
15 unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, or an unsubstituted or substituted 5-10 membered heteroaryl; and/or

- the substituted C₁-C₆ alkyl, the substituted C₁-C₆ alkyl-O-, the substituted C₂-C₆ alkenyl, the substituted C₂-C₆ alkenyl-O-, the substituted C₆-C₁₀ aryl, the substituted C₃-C₅
20 heterocycloalkyl, the substituted C₃-C₅ heterocycloalkenyl or the substituted 5-6 membered heteroaryl is substituted with one or more substituents independently selected from the following: halogen, -OH, -CF₃, -CN, -COOH, -C(=O)NH₂, =O (i.e., two geminal hydrogens on a carbon atom are substituted with the group O), and a C₁-C₆ alkyl, a C₁-C₆ alkyl-O- or a C₃-C₆ cycloalkyl that is unsubstituted or substituted with one or more R^a, wherein when a plurality of
25 R^a are present, they are the same or different, and each R^a is halogen, -OH, =O, -CF₃, -CN, -COOH or -C(=O)NH₂; and when a plurality of substituents are present, they are the same or different; and/or



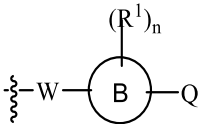
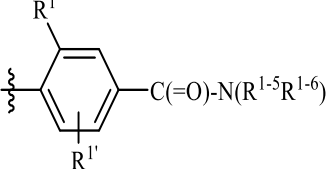
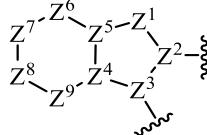
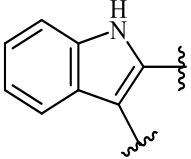
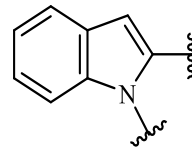
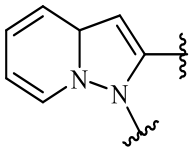
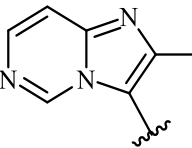
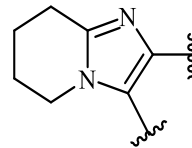
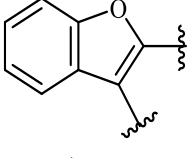
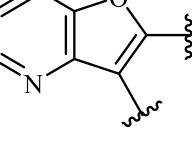
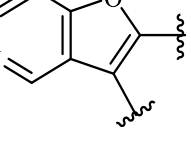
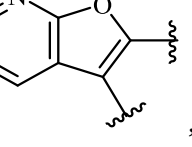
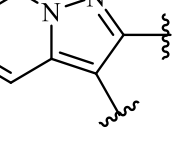
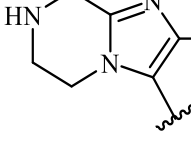
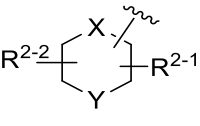
15. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the
30 tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein the heterocyclic compound of formula I is the following embodiments:

embodiment I: when Q is H, halogen, -OH, -CF₃, -CN, -NH-C(=O)-R¹⁻¹, -C(=O)-R¹⁻², -O-C(=O)-N(R¹⁻³R¹⁻⁴), -C(=NH)-N(R¹⁻⁷R¹⁻⁸), -S(=O)₂-N(R¹⁻⁹R¹⁻¹⁰), -S(=O)-N(R¹⁻¹¹R¹⁻¹²), -O-C(=O)-R¹⁻¹³, -N(R¹⁻¹⁴R¹⁻¹⁵), -NH-S(=O)₂-R¹⁻¹⁶, an unsubstituted or substituted C₁-C₆ alkyl, an unsubstituted or substituted C₁-C₆ alkyl-O-, an unsubstituted or substituted C₂-C₆ alkenyl, an unsubstituted or substituted C₂-C₆ alkenyl-O-, an unsubstituted or substituted C₃-C₁₀ cycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkyl, an unsubstituted or substituted C₃-C₉ heterocycloalkenyl, an unsubstituted or substituted C₆-C₁₀ aryl, or an unsubstituted or substituted

5-10 membered heteroaryl,  is , or R² is , ; or

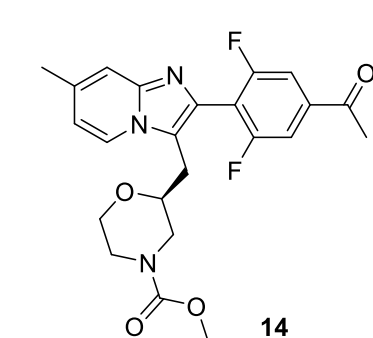
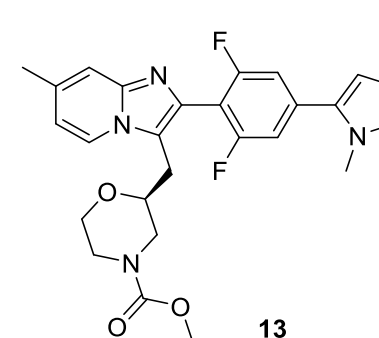
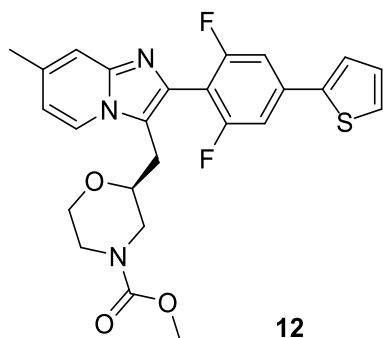
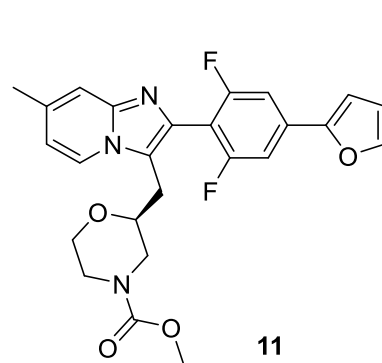
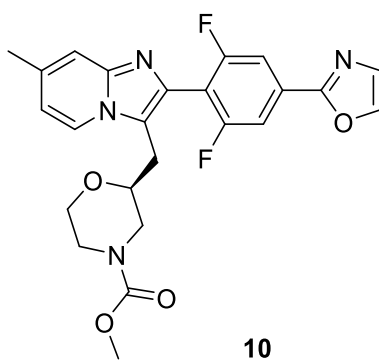
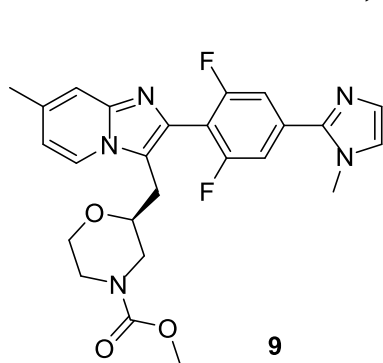
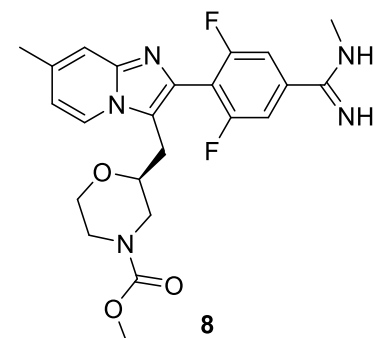
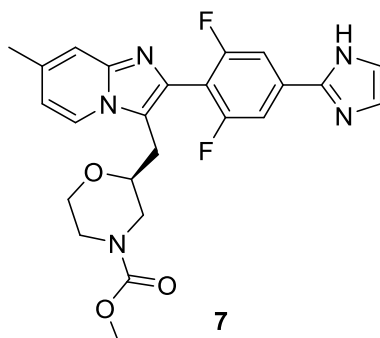
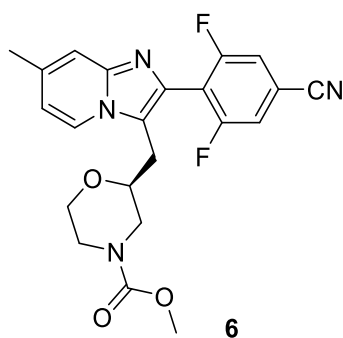
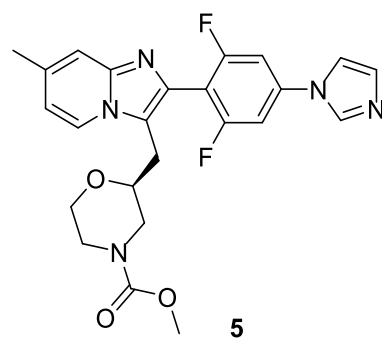
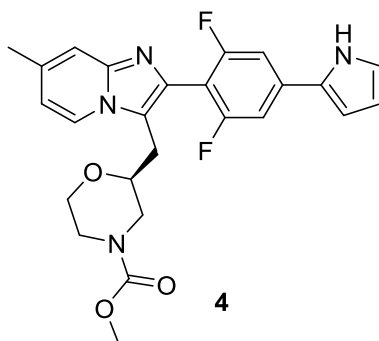
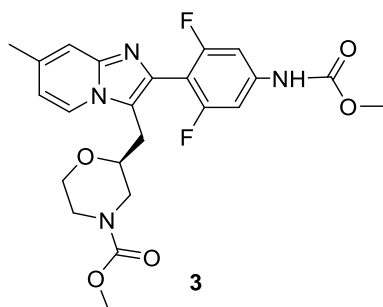
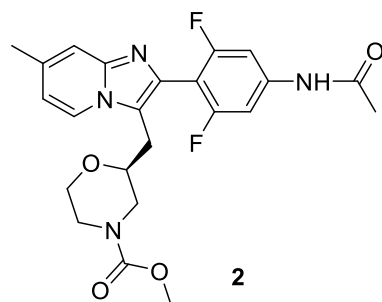
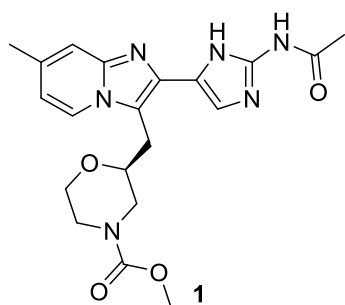
wherein X and Y are each independently a single bond, methylene, -O- or

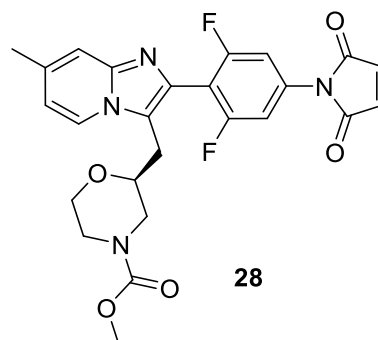
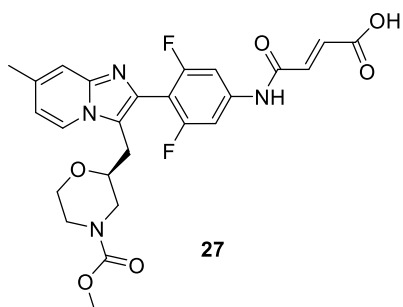
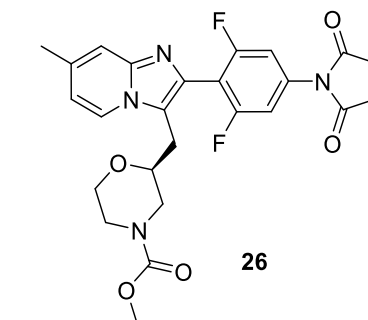
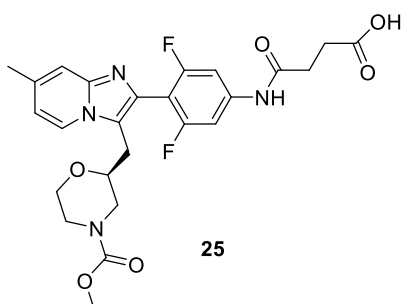
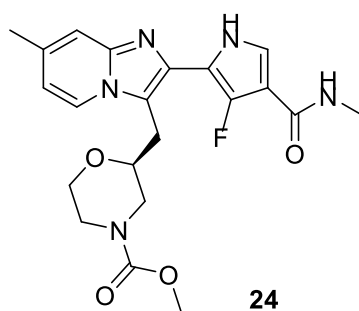
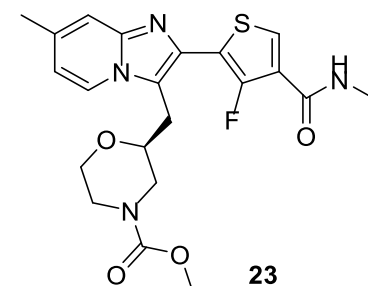
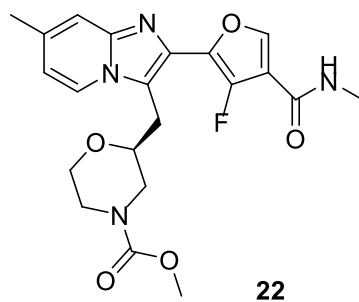
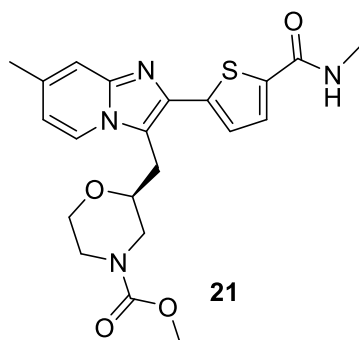
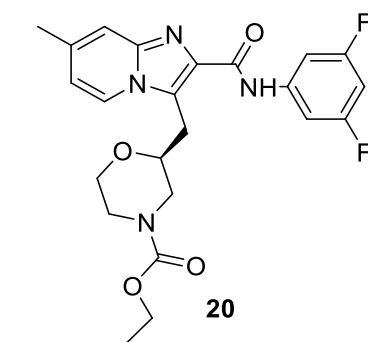
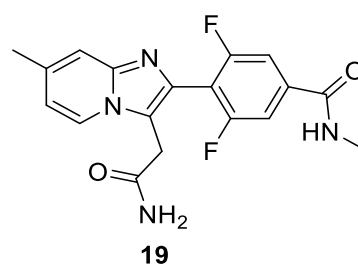
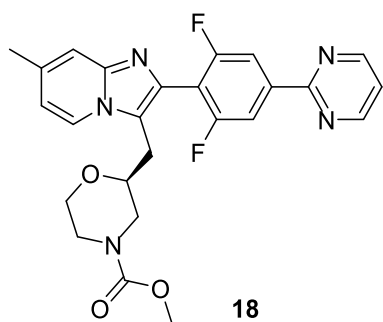
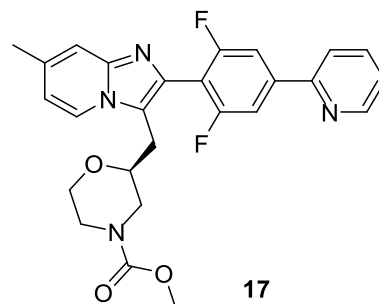
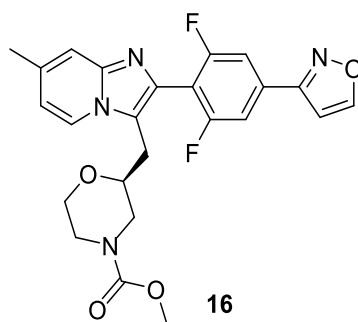
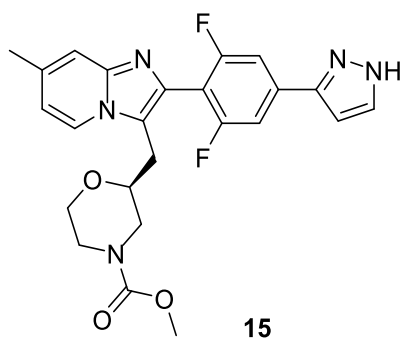
10 embodiment II: when

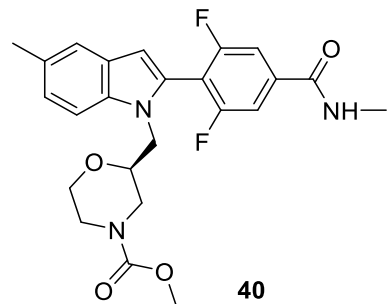
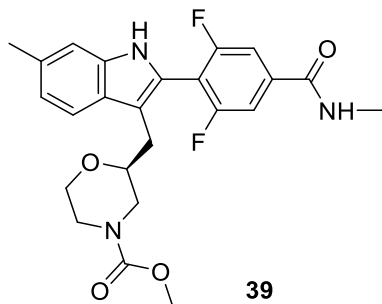
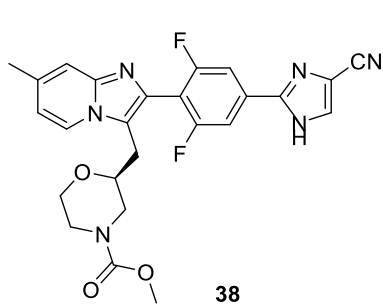
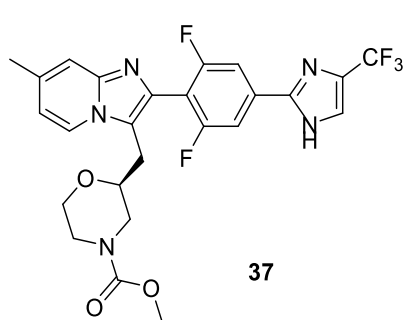
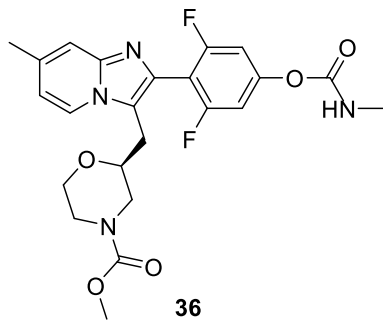
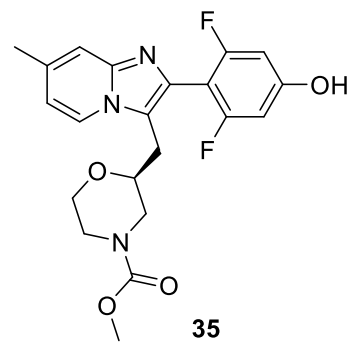
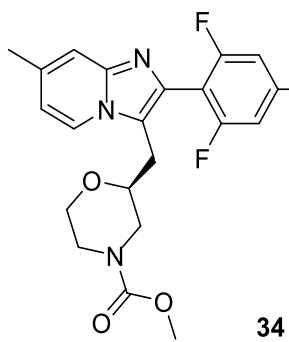
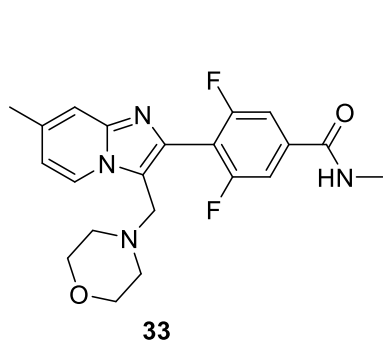
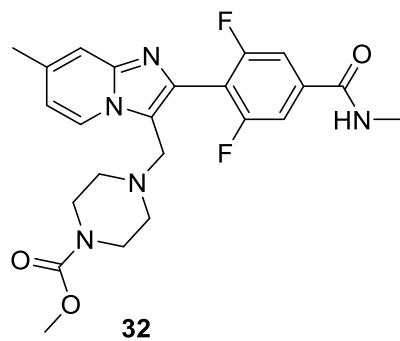
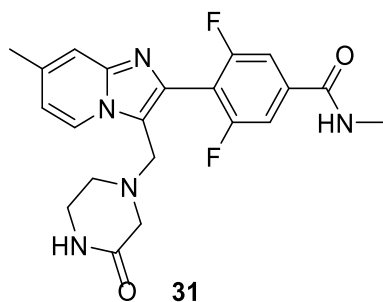
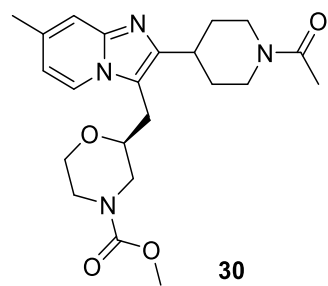
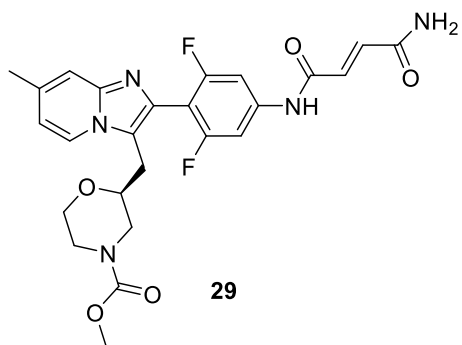
 is ,  is , , , , , , , , , , or , or R² is independently , -C(=O)N(R²⁻⁵R²⁻⁶) or an

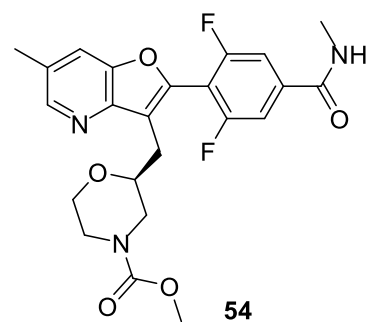
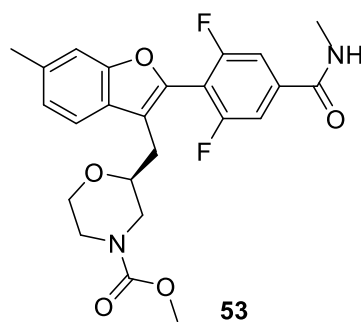
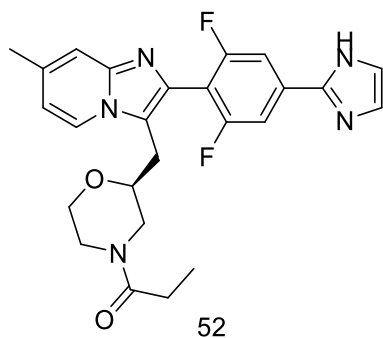
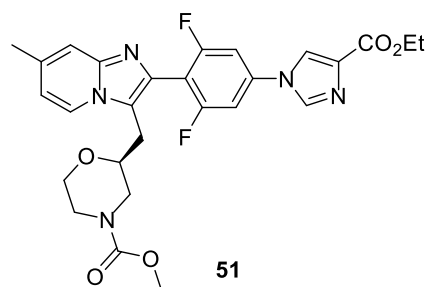
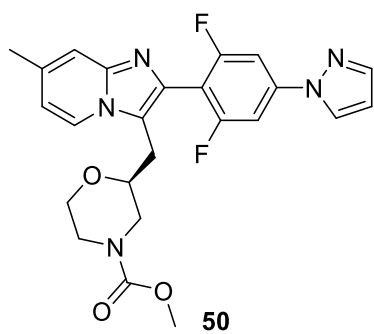
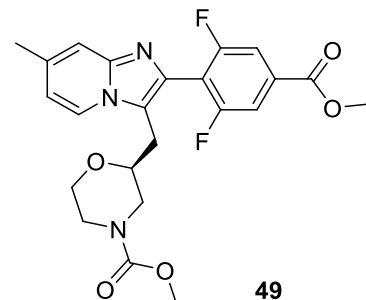
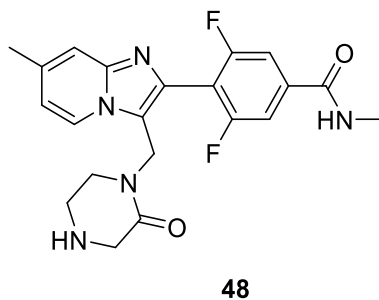
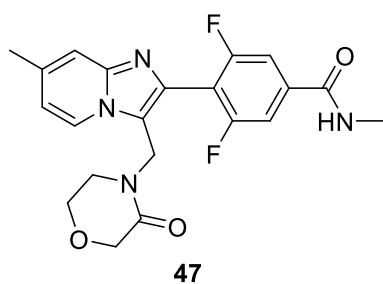
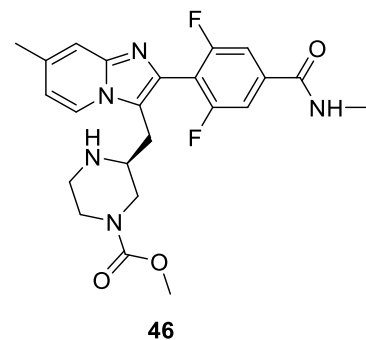
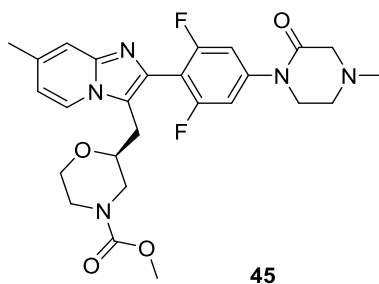
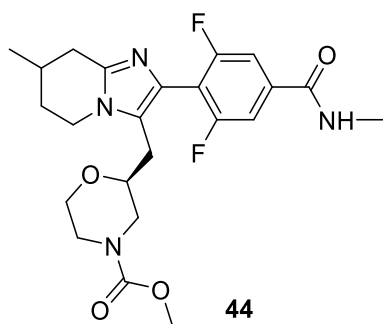
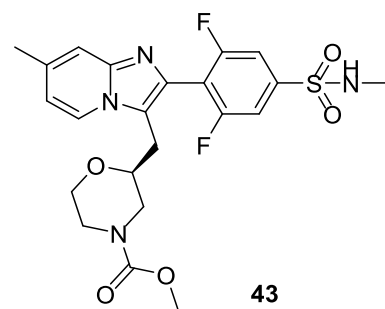
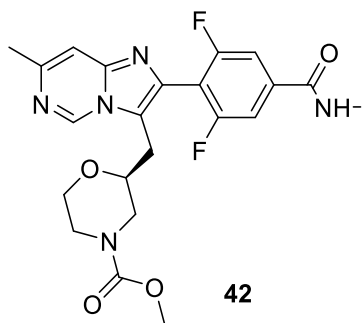
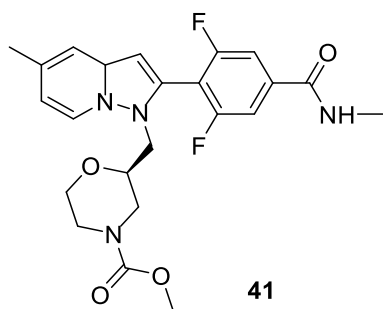
unsubstituted or substituted 5-6 membered heteroaryl, wherein X or Y is -N(R²⁻³)-.

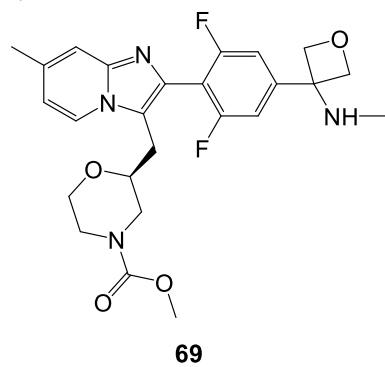
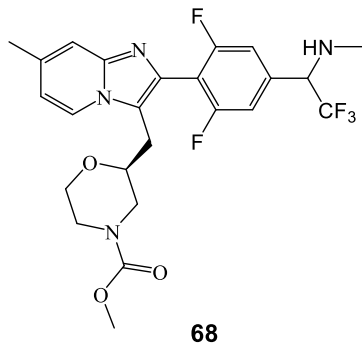
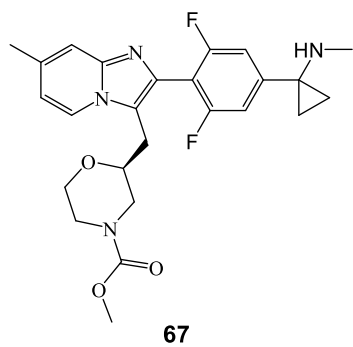
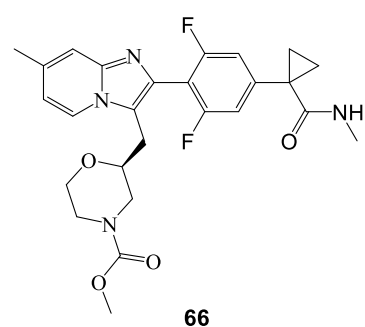
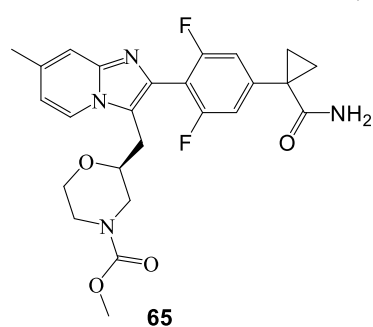
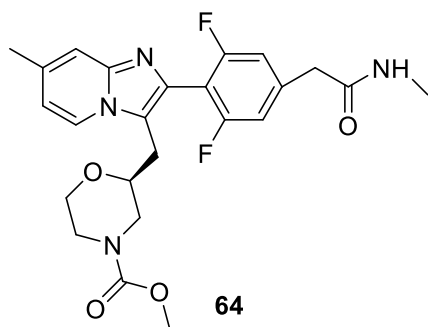
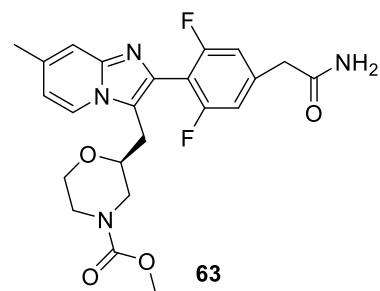
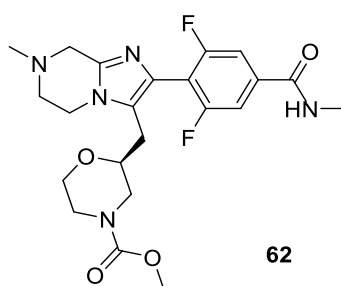
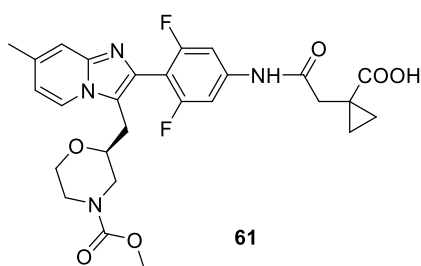
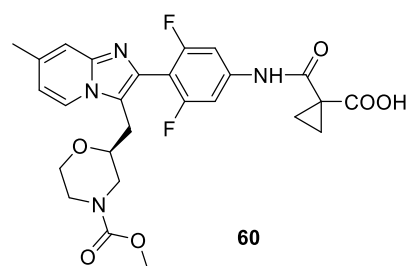
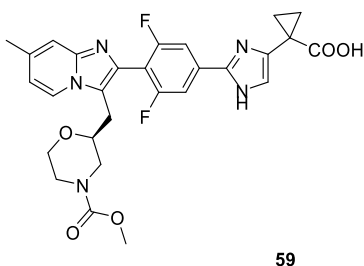
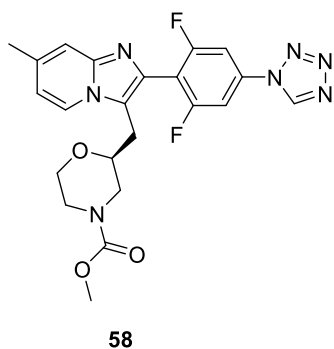
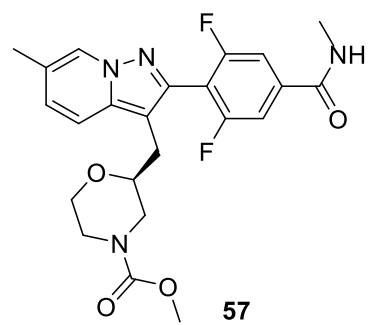
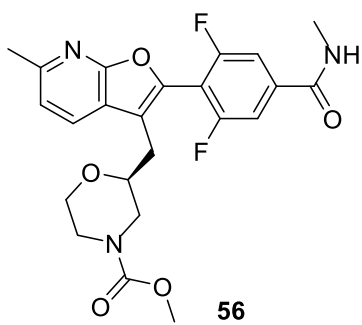
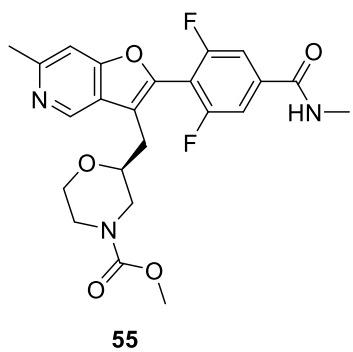
15 16. The heterocyclic compound of formula I or the stereoisomer, the geometric isomer, the tautomer, the nitrogen oxide, the hydrate, the solvate, the metabolite, the ester, the pharmaceutically acceptable salt or the prodrug thereof according to claim 1, wherein the heterocyclic compound of formula I is preferably any one of the following compounds:

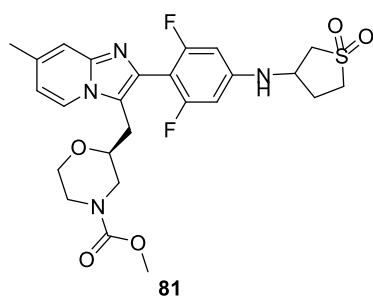
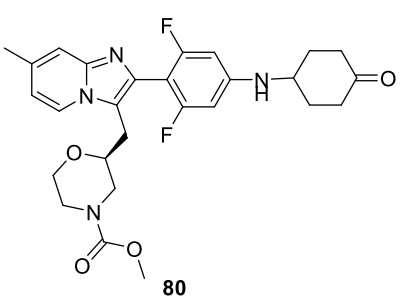
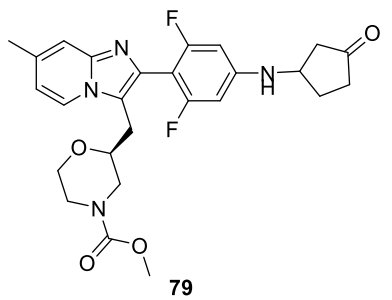
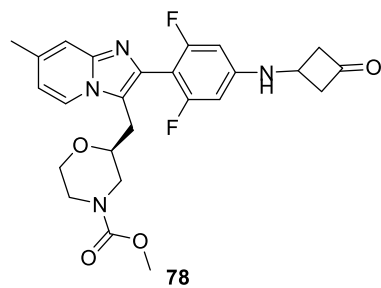
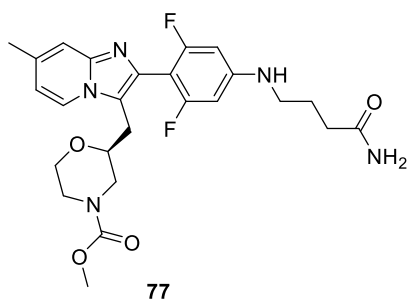
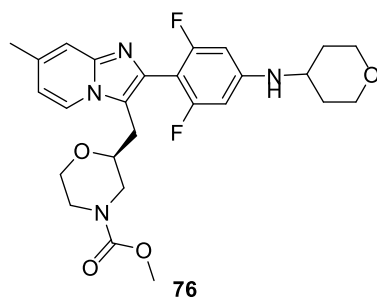
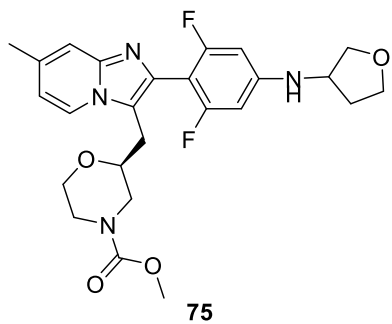
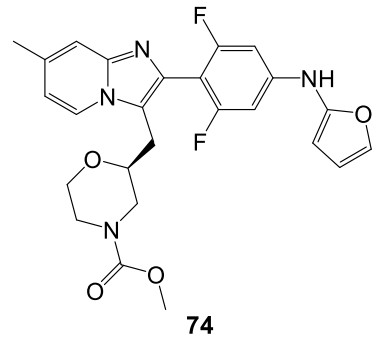
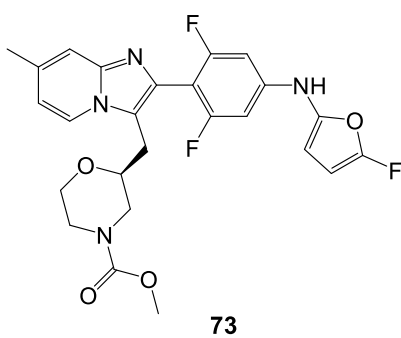
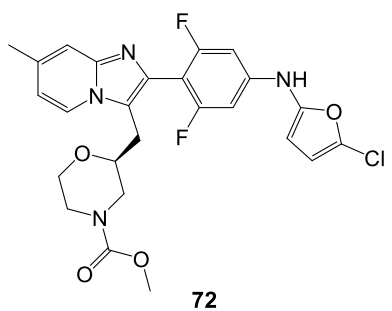
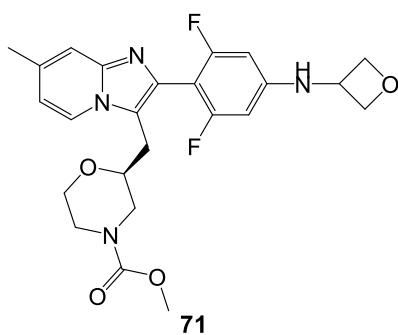
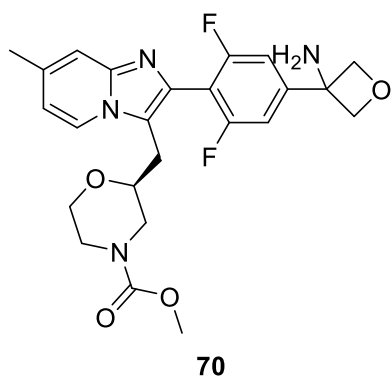


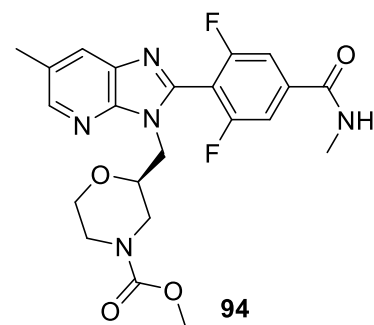
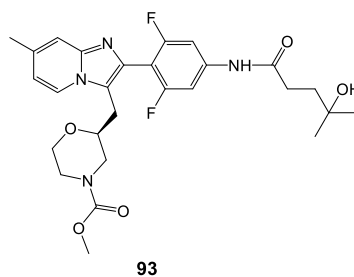
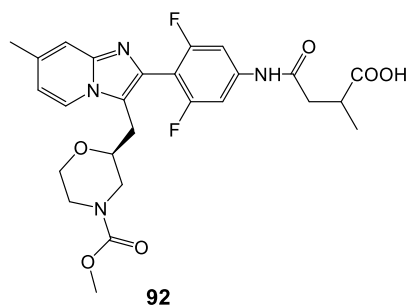
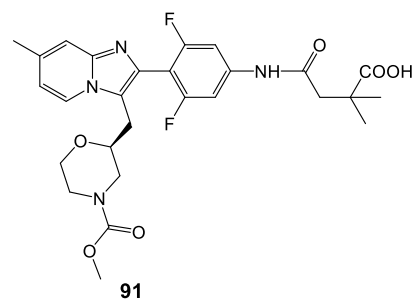
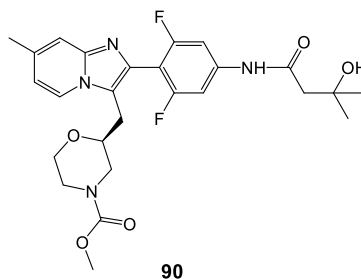
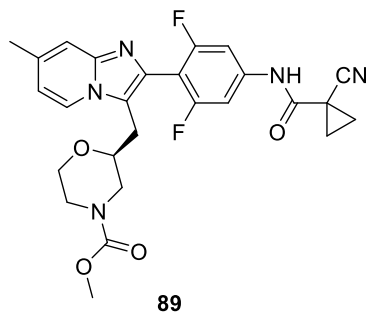
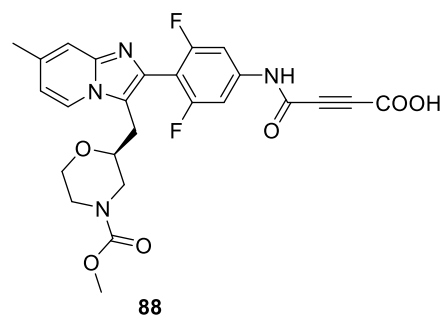
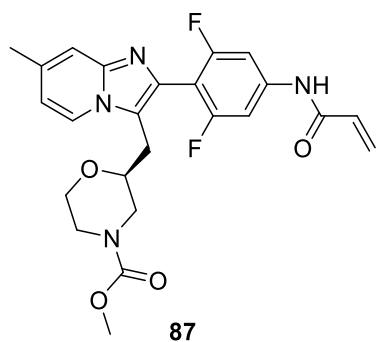
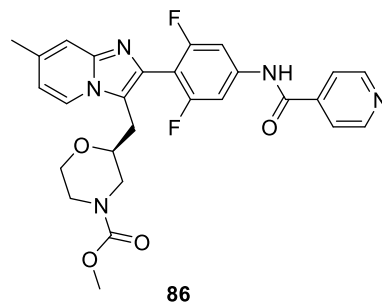
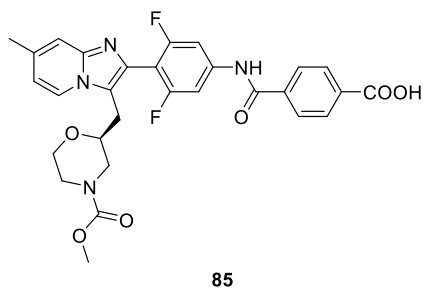
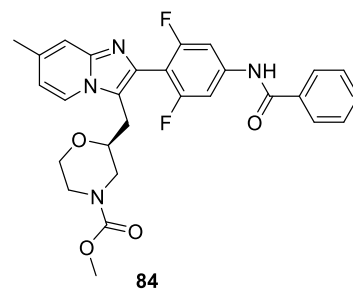
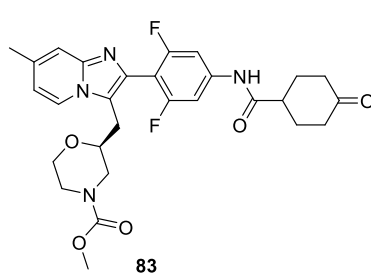
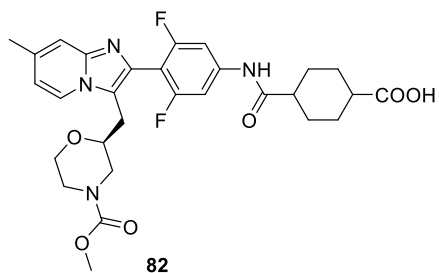


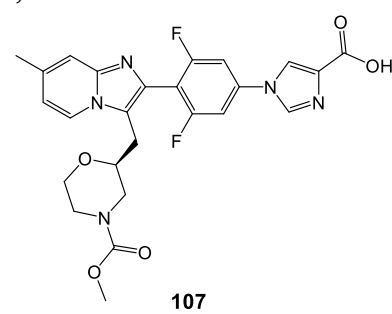
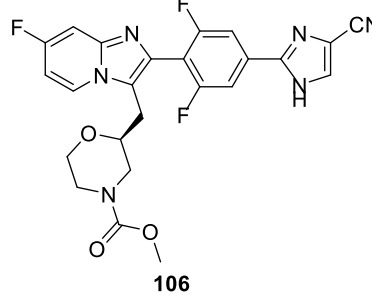
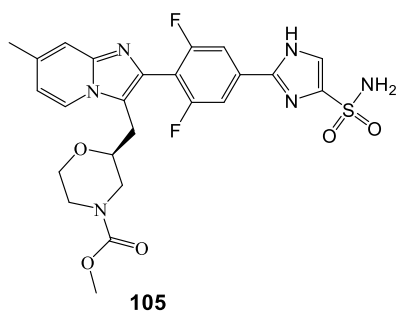
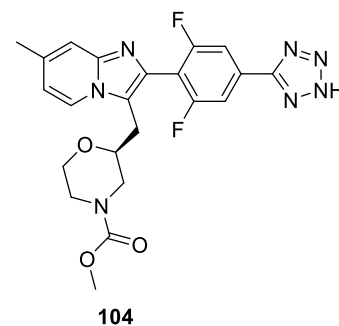
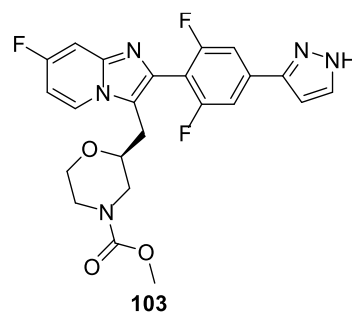
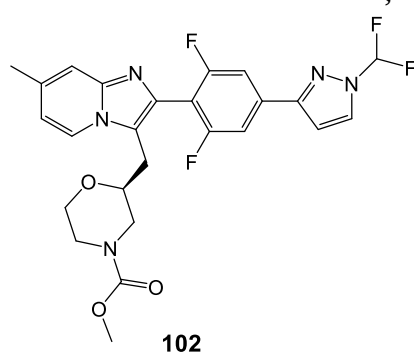
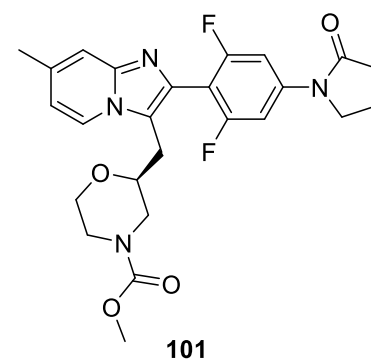
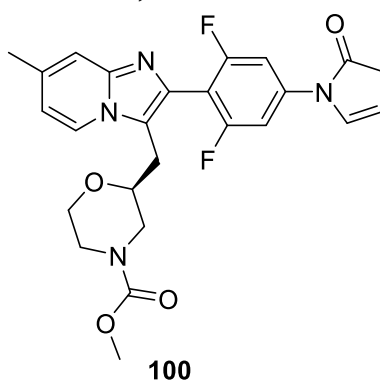
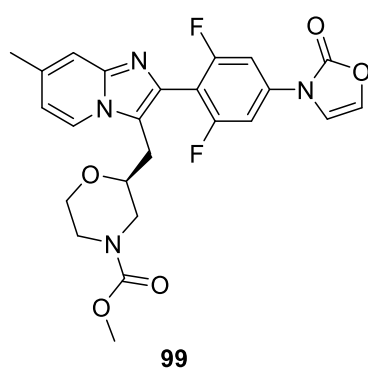
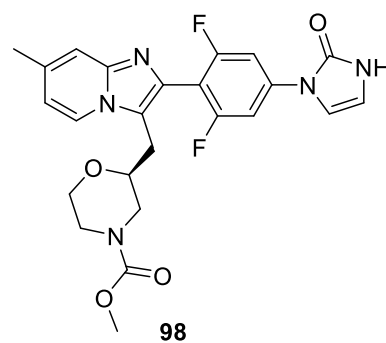
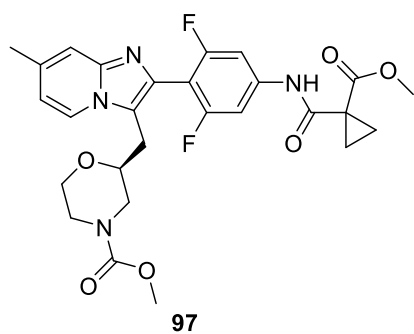
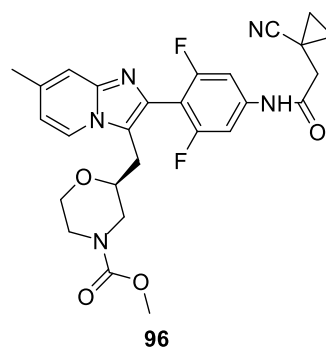
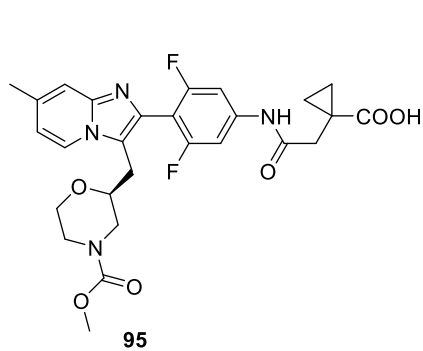


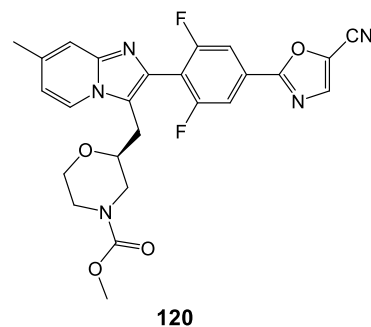
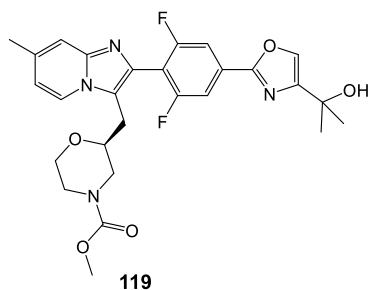
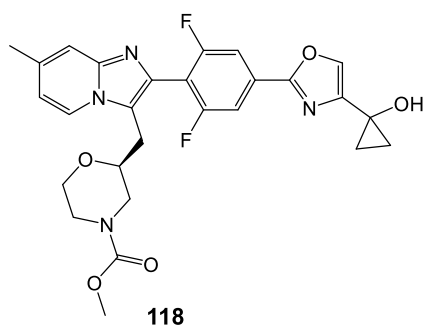
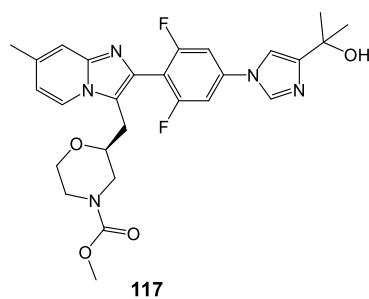
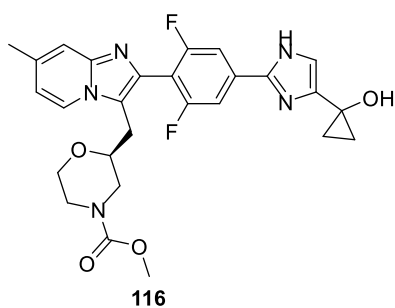
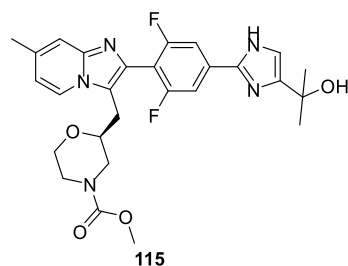
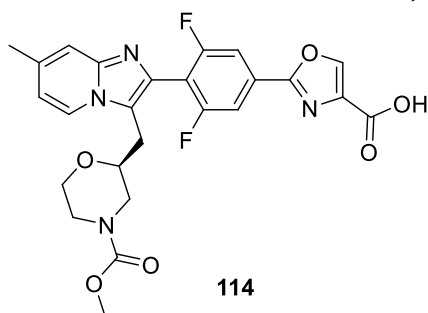
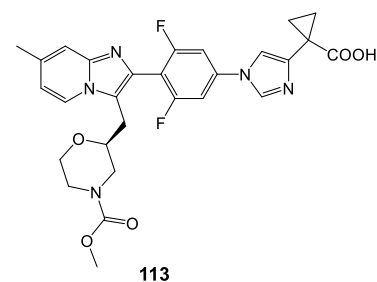
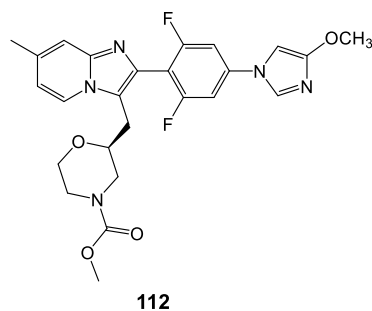
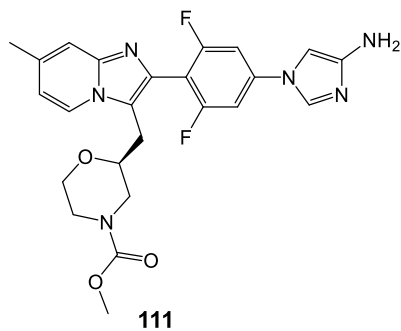
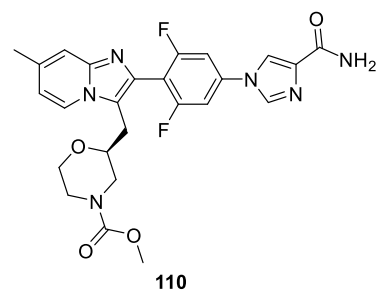
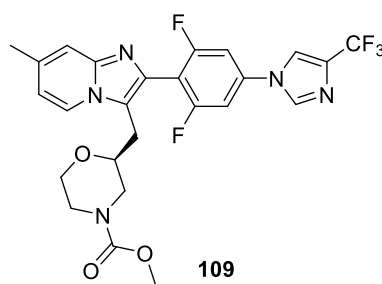
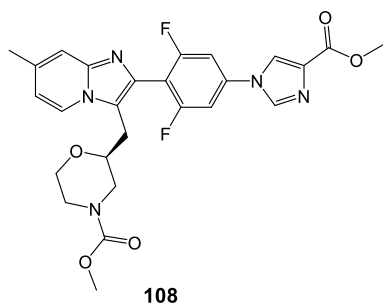


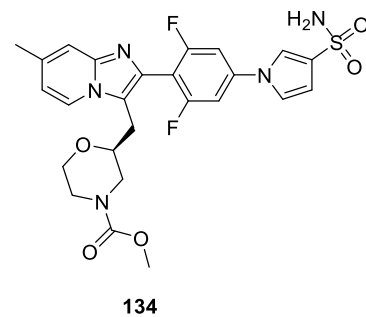
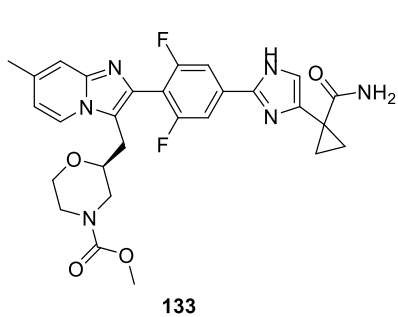
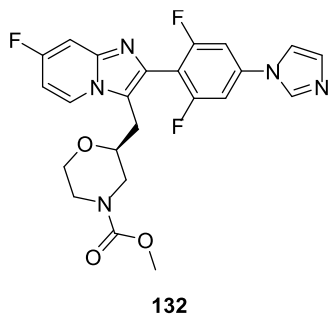
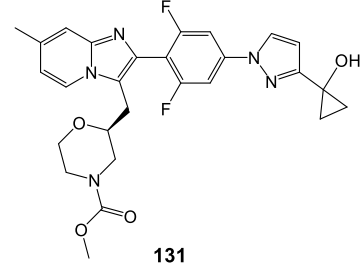
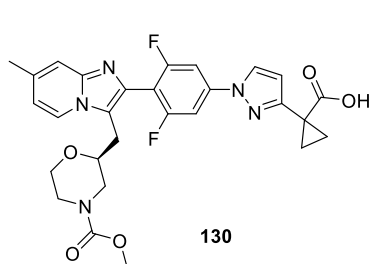
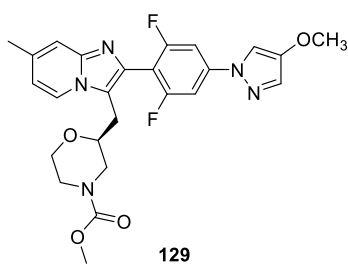
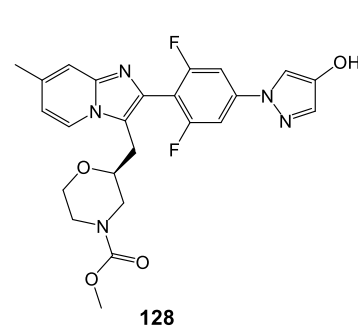
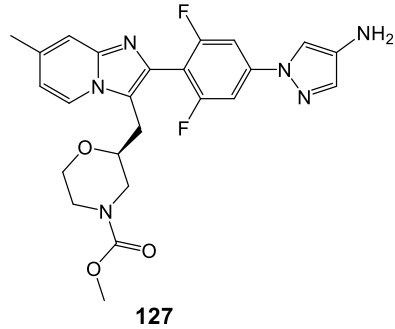
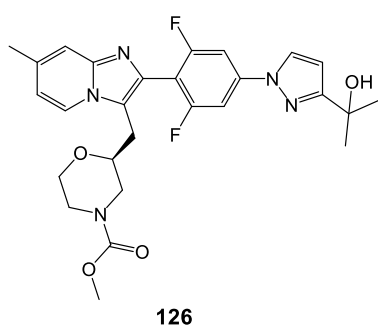
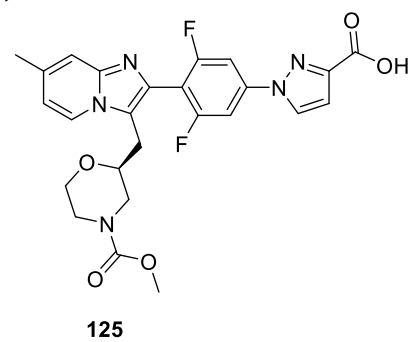
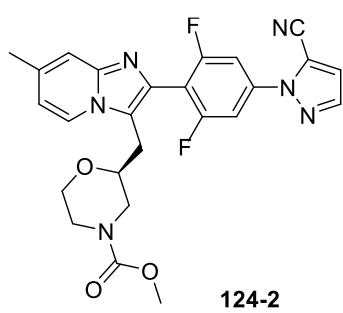
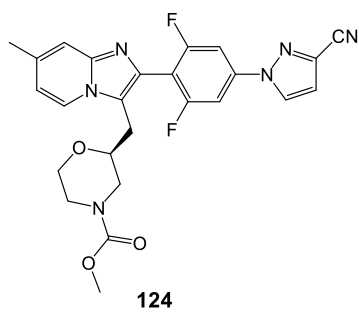
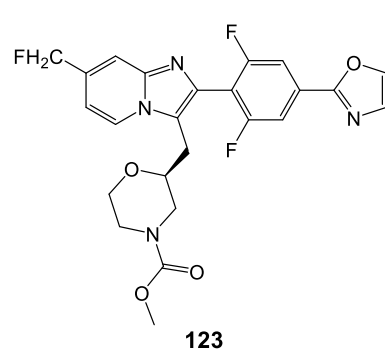
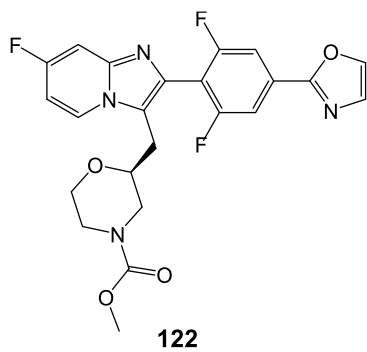
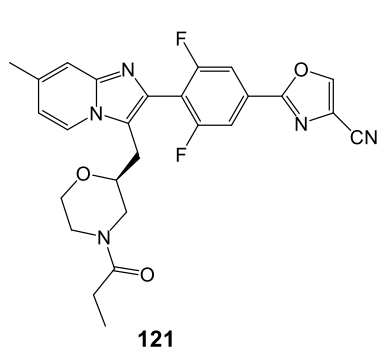


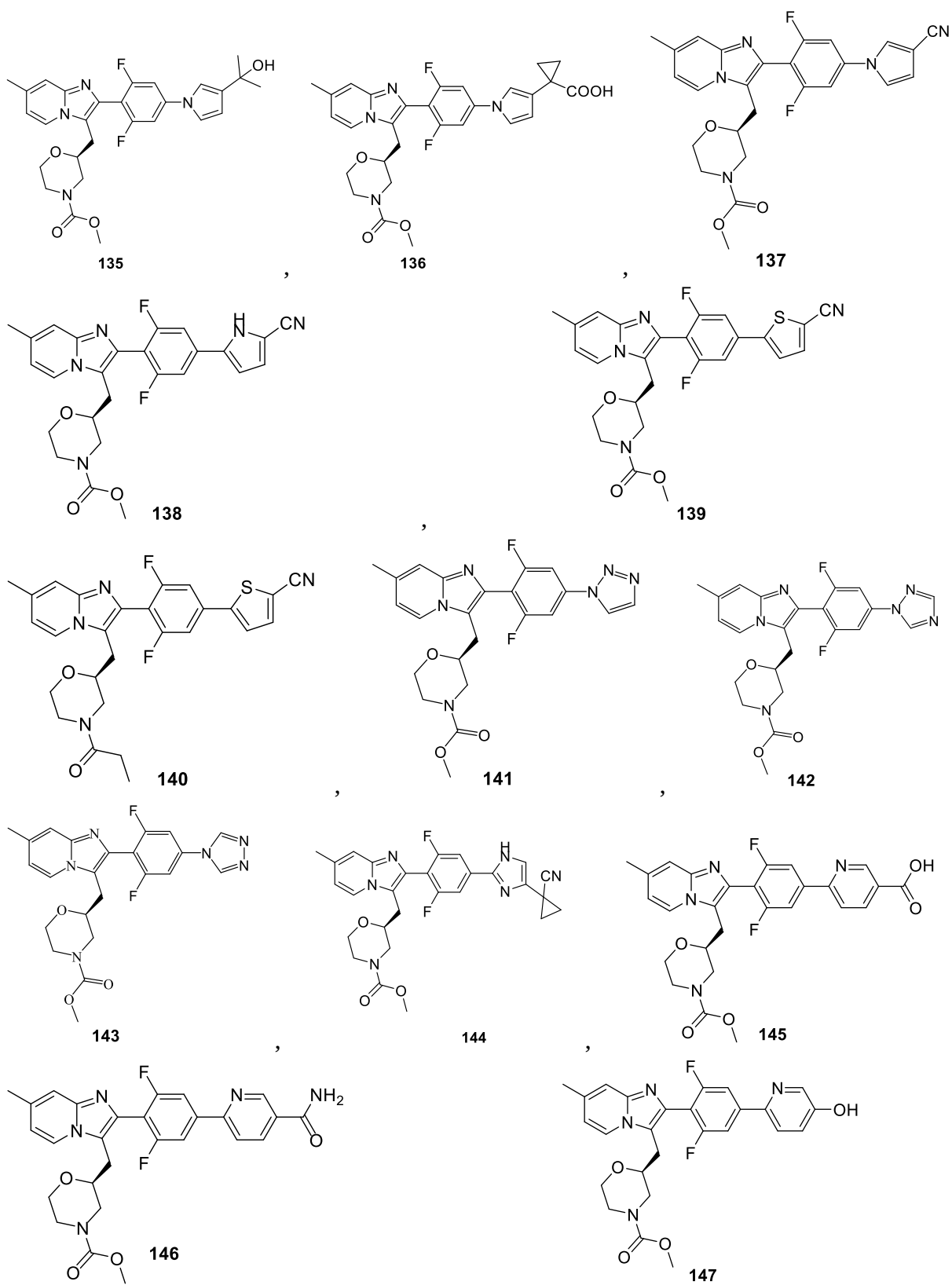


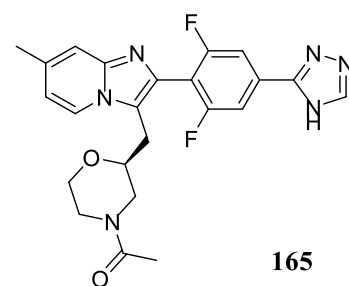
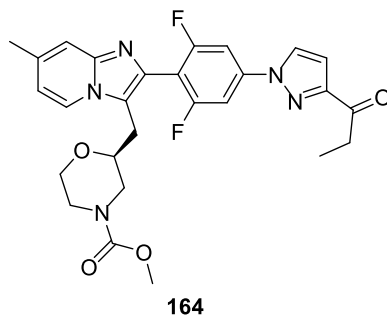
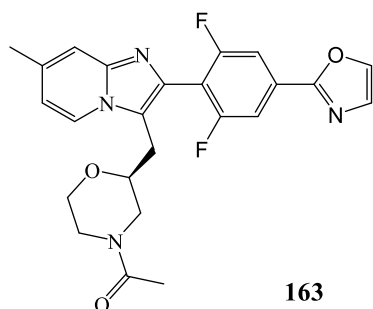
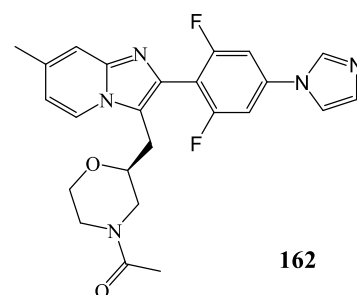
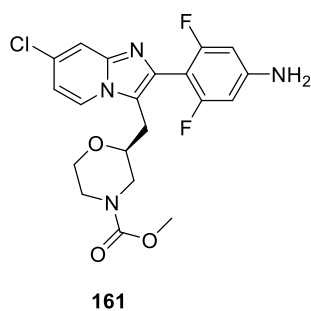
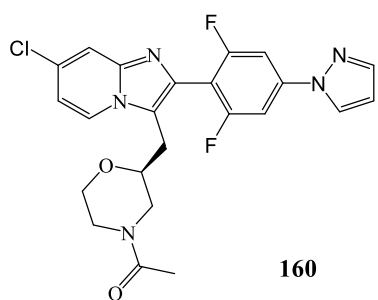
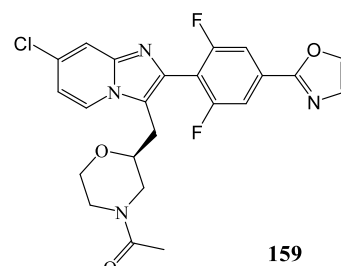
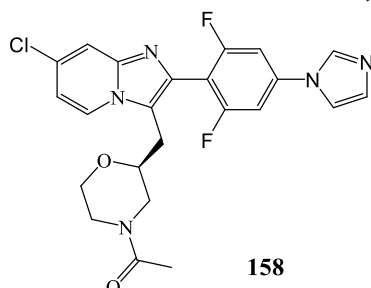
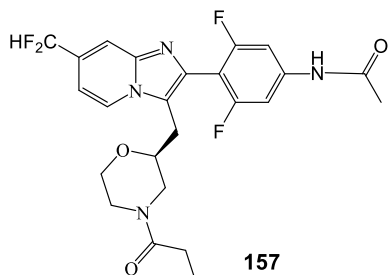
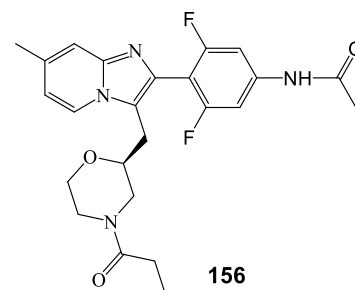
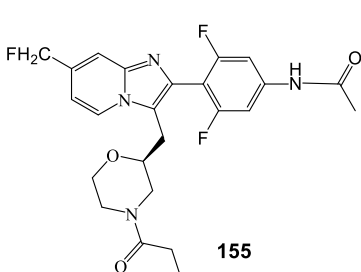
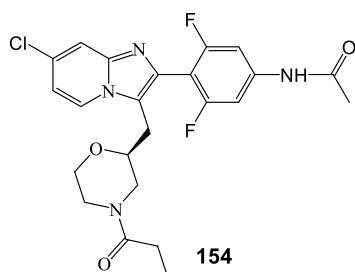
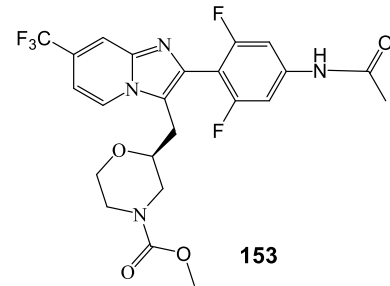
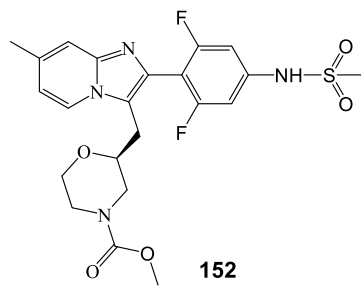
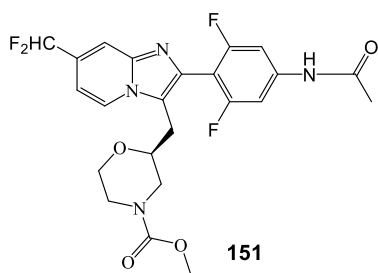
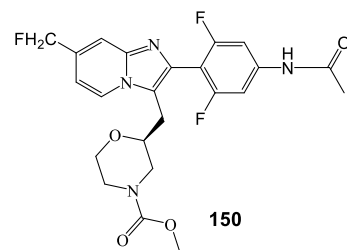
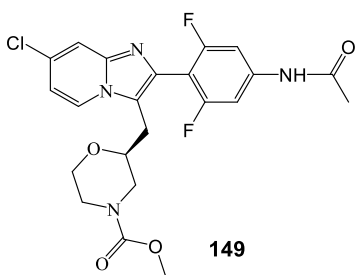
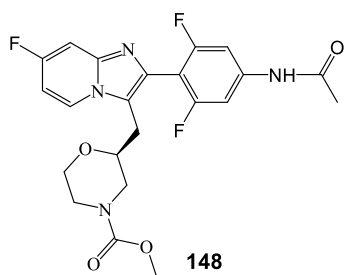




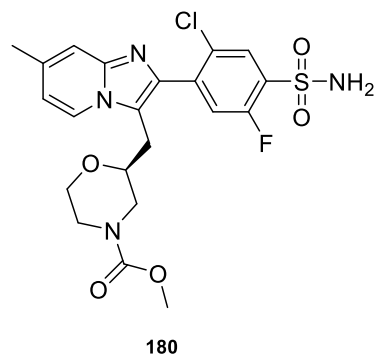
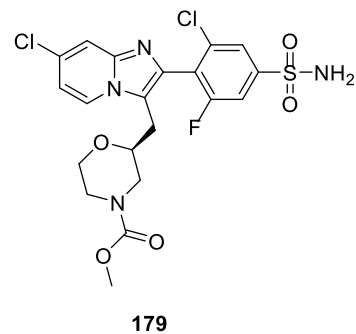
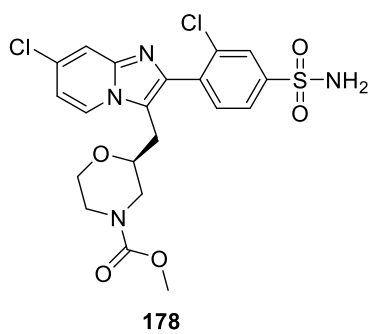
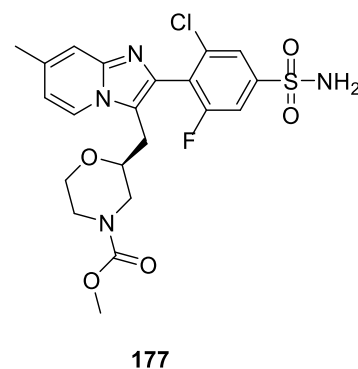
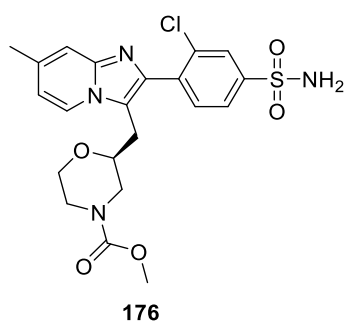
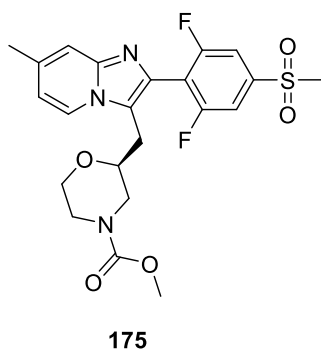
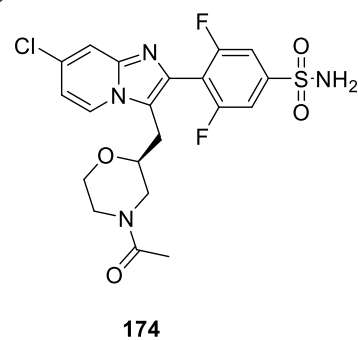
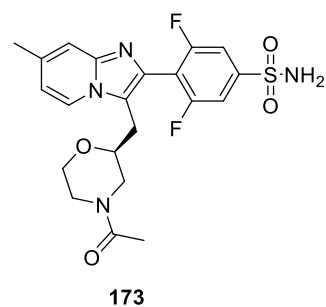
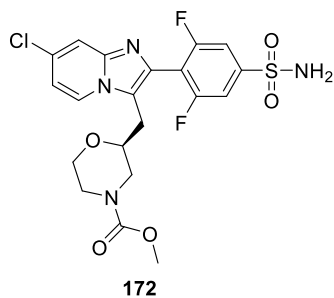
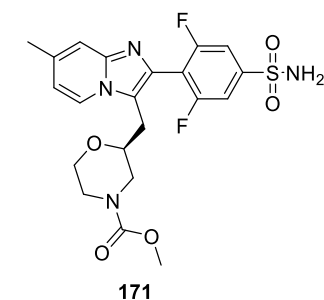
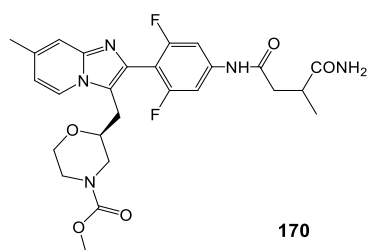
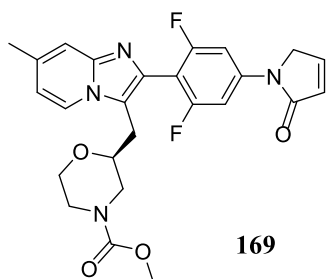
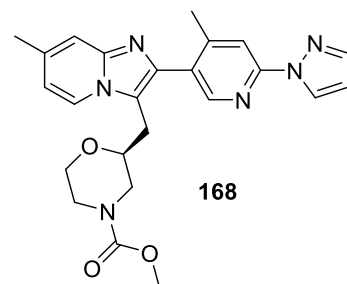
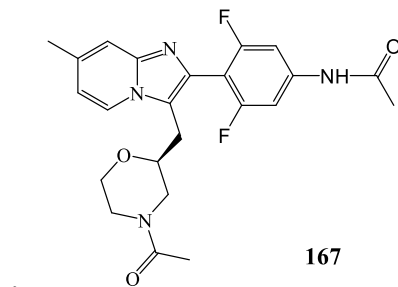
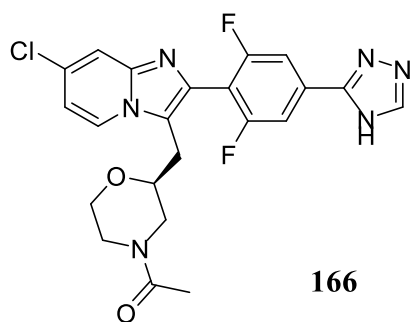


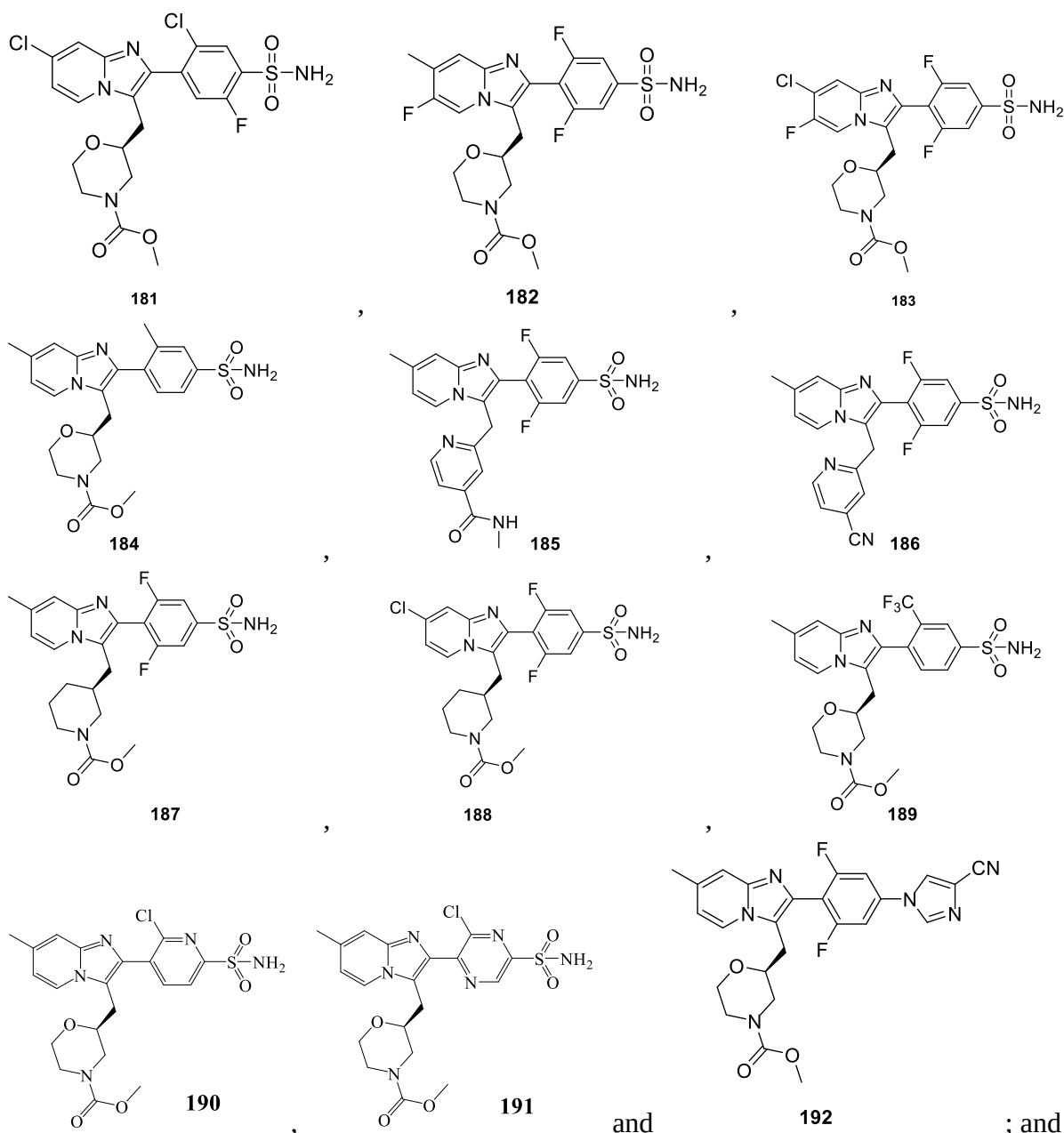




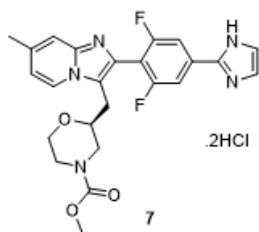


5



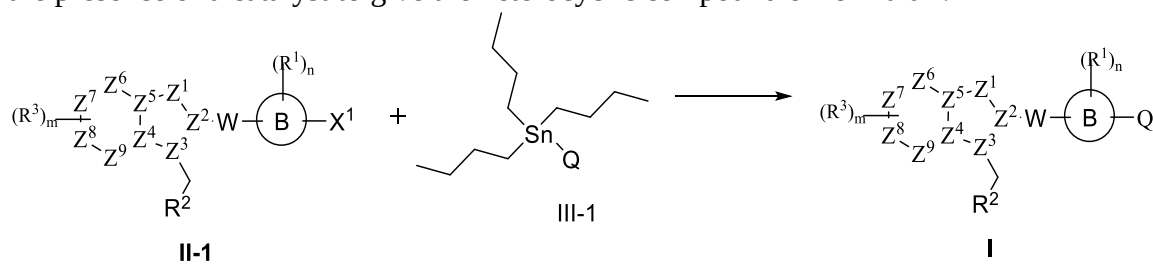


5 the pharmaceutically acceptable salt of the heterocyclic compound of the formula I is the following compound:



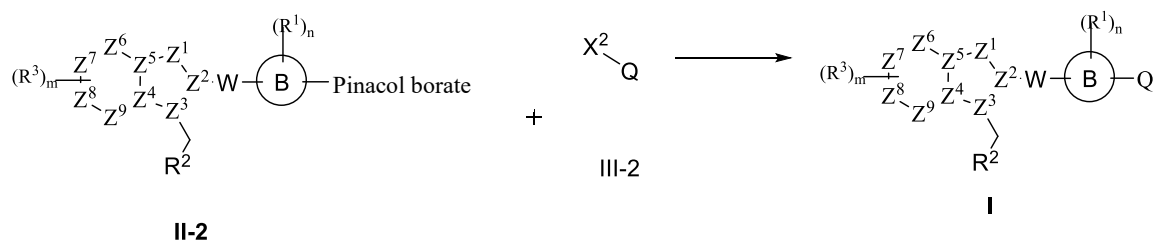
17. A preparation method of the heterocyclic compound of formula I according to any one of claims 1 to 16, wherein the preparation method is Scheme I, Scheme II, Scheme III, Scheme IV, Scheme V, Scheme VI, Scheme VII or Scheme VIII, wherein

Scheme I comprises the following step: performing Stille coupling reaction as shown below on a compound of formula II-1 and an organic tin reagent of formula III-1 in an organic solvent in the presence of a catalyst to give the heterocyclic compound of formula I:



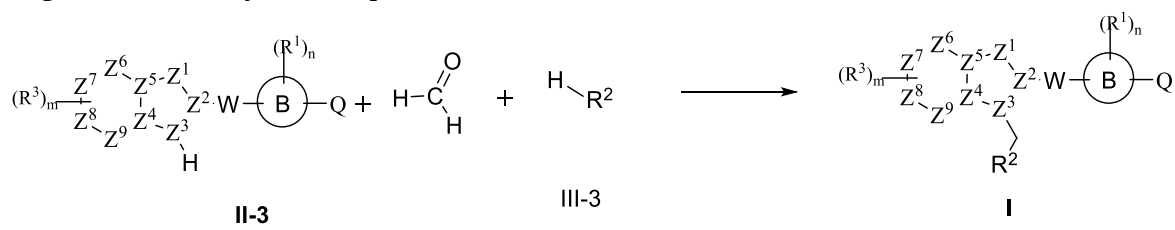
5 wherein X¹ is Br or Cl;

Scheme II comprises the following step: performing Suzuki coupling reaction as shown below on a compound of formula II-2 and a compound of formula III-2 in a solvent in the presence of a catalyst and a basic reagent to give the heterocyclic compound of formula I:

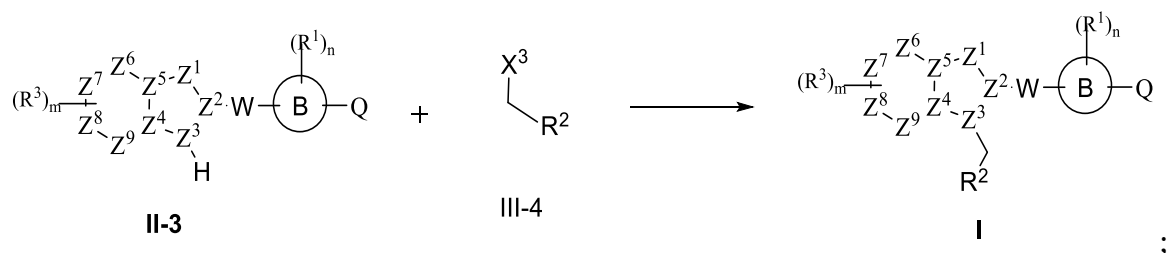


10 wherein X² is Br or Cl;

Scheme III comprises the following step: performing the reaction as shown below on a compound of formula II-3, a compound of formula III-3 and formaldehyde in an organic solvent to give the heterocyclic compound of formula I:

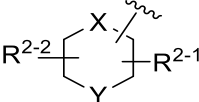


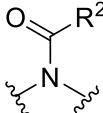
15 Scheme IV comprises the following step: performing the reaction as shown below on a compound of formula II-3 and a compound of formula III-4 in an organic solvent to give the heterocyclic compound of formula I:

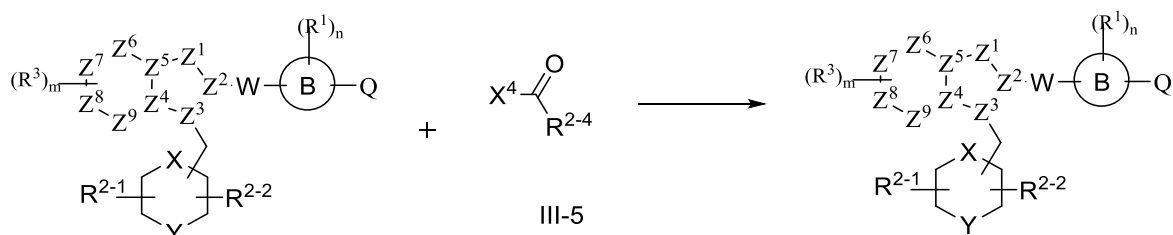


wherein X³ is a leaving group;

Scheme V comprises the following step: performing amidation reaction as shown below on

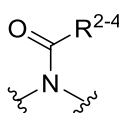
a compound of formula I, wherein R^2 is  and Y is -NH-, and a compound of formula III-5 in an organic solvent in the presence of a basic reagent to give the heterocyclic

compound of formula I, wherein Y is  :



Y is -NH-

I

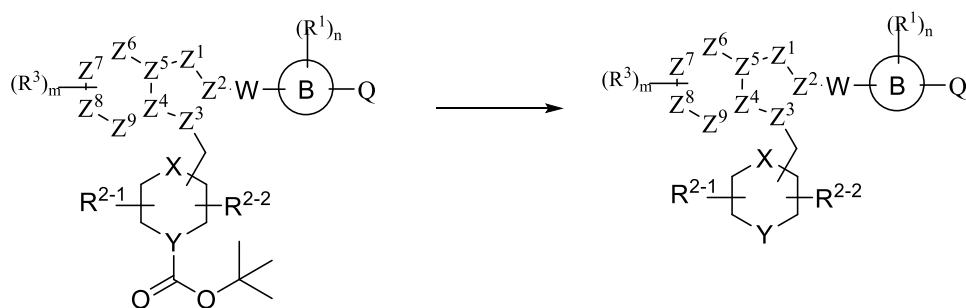
Y is 

I

;

wherein X^4 is halogen;

Scheme VI comprises the following step: performing deprotection reaction as shown below on a compound of formula II-4 and a compound of formula III-5 in an organic solvent in the presence of an acid to give the heterocyclic compound of formula I:



Y is -NH-

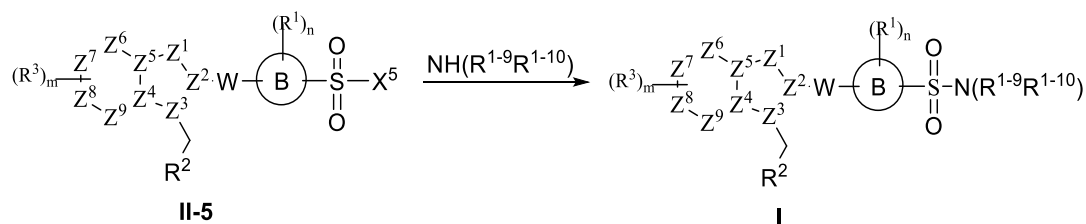
II-4

Y is -NH-

I

wherein Y is -NH-;

Scheme VII comprises the following step: performing substitution reaction as shown below on a compound of formula II-5 and $NH(R^{1-9}R^{1-10})$ in an organic solvent to give the heterocyclic compound of formula I, wherein Q is $-S(=O)_2-N(R^{1-9}R^{1-10})$:



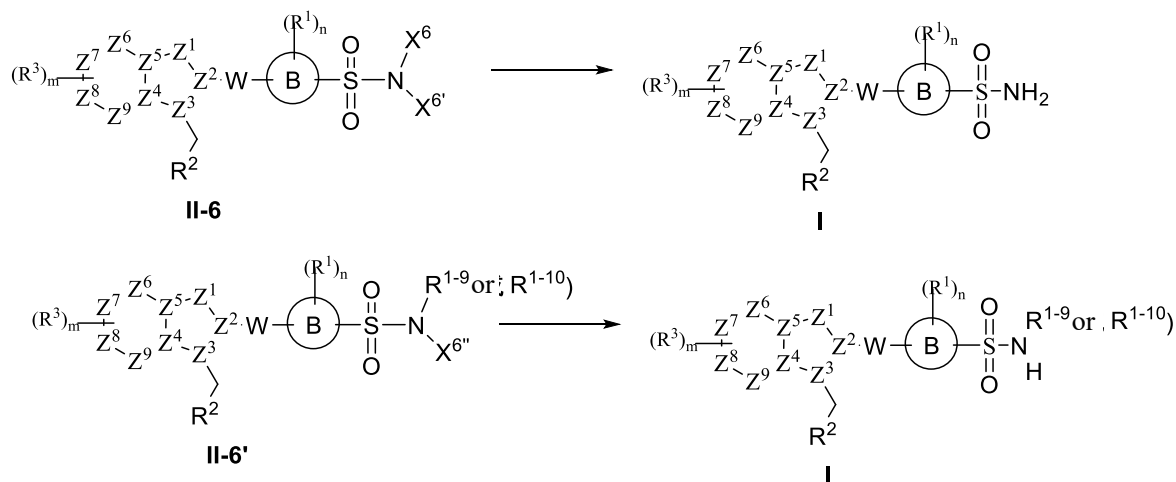
II-5

I

;

wherein X^5 is Br or Cl;

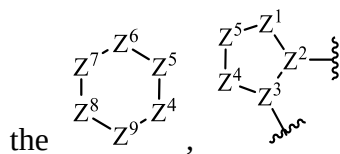
Scheme VIII comprises the following step: performing deprotection reaction as shown below on a compound of formula II-6 or formula II-6' in an organic solvent in the presence of an acid to give the heterocyclic compound of formula I:



5

;

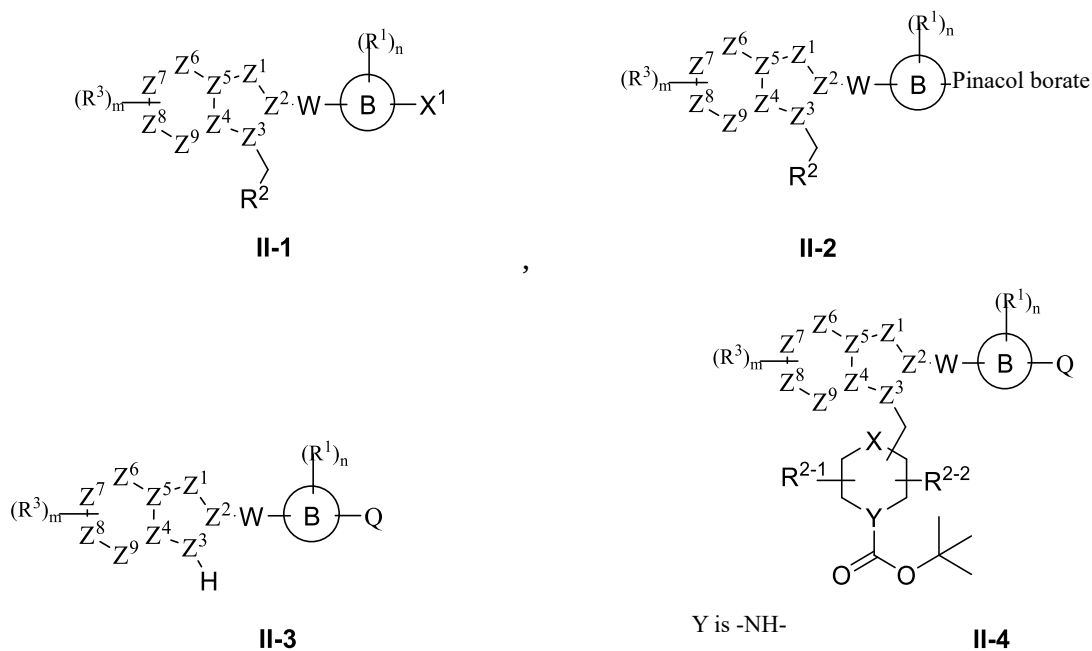
wherein X^6 , $X^{6'}$ and $X^{6''}$ are N protecting groups; and

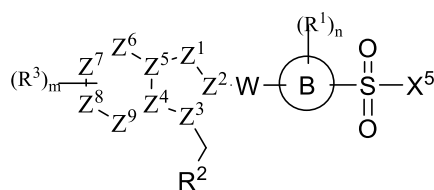


, R^3 , R^2 , R^1 , W, X, Y, R^{2-1} , R^{2-2} , n and m have the definitions as defined in any one of claims 1 to 16.

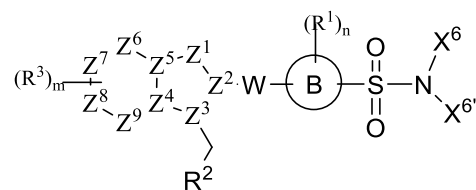
10

18. A compound of formula II-1, II-2, II-3, II-4, II-5, II-6 or II-6':



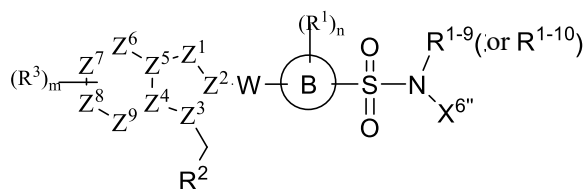


II-5



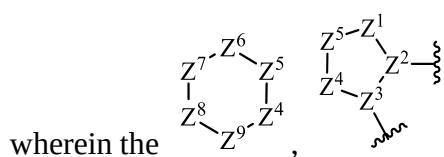
II-6

or



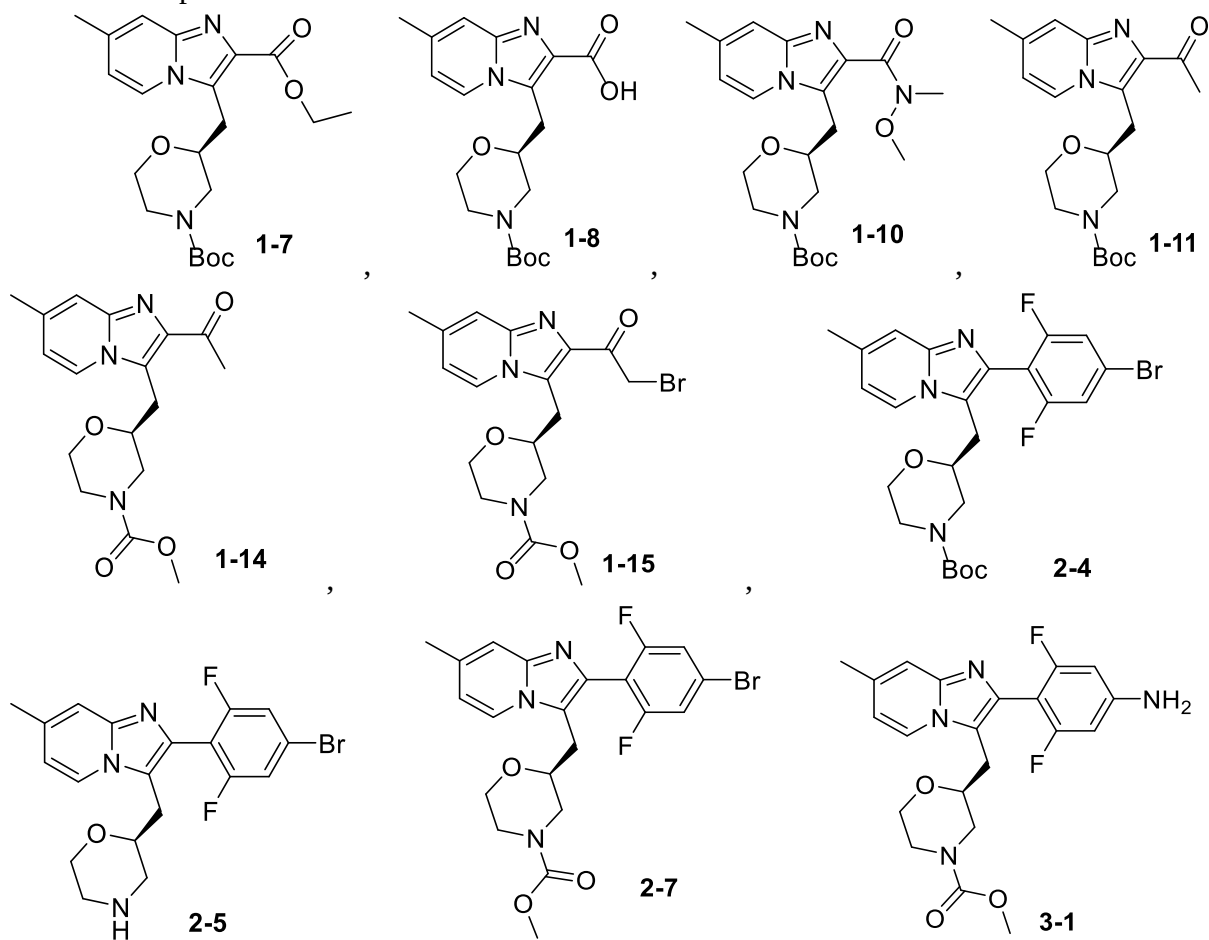
II-6'

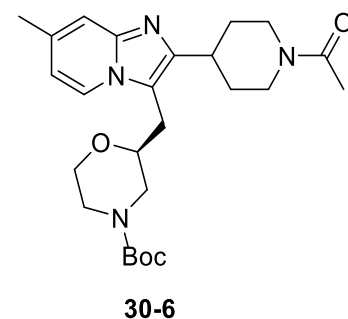
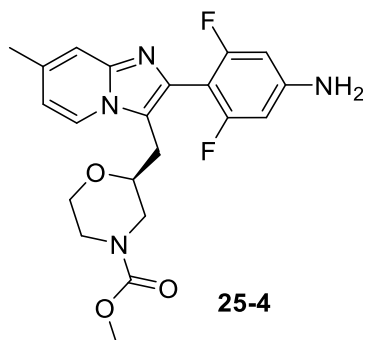
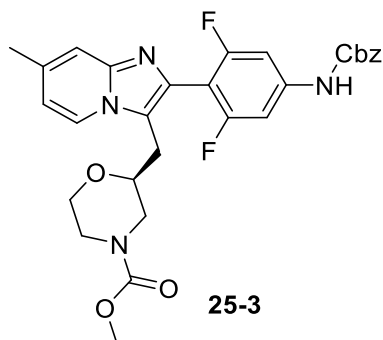
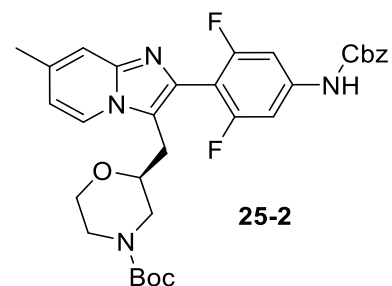
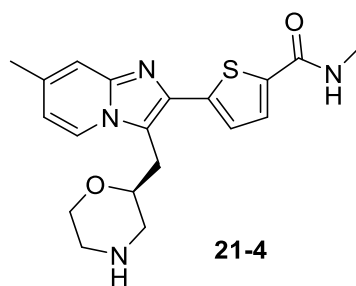
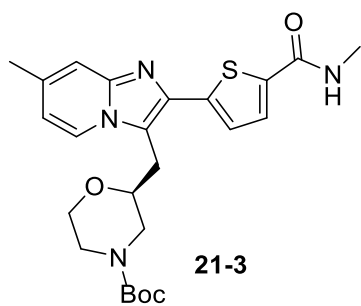
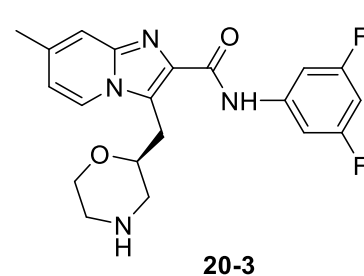
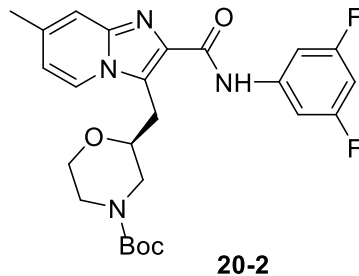
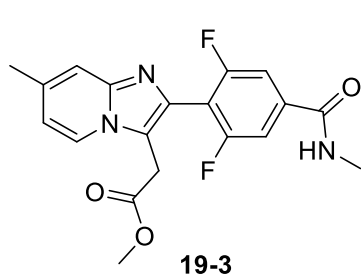
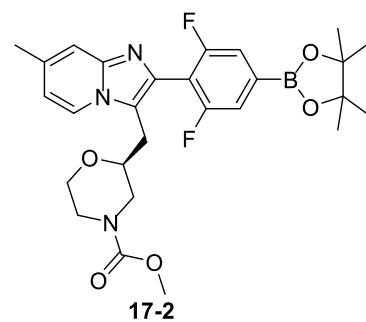
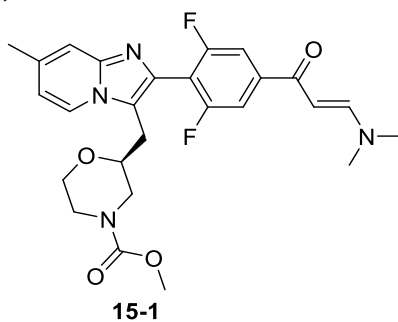
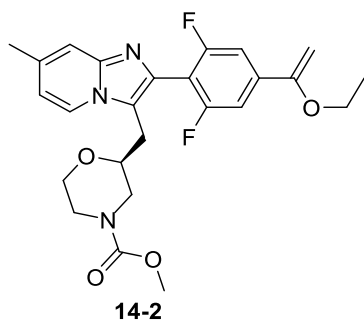
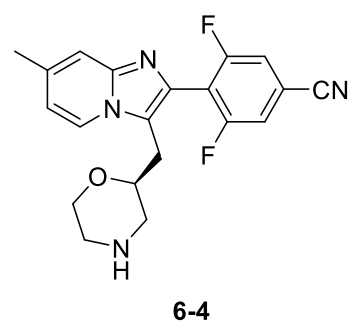
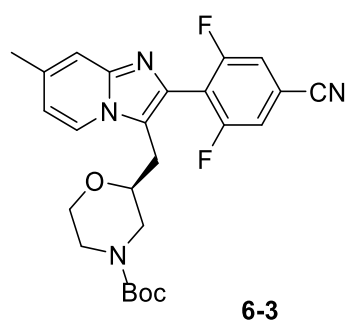
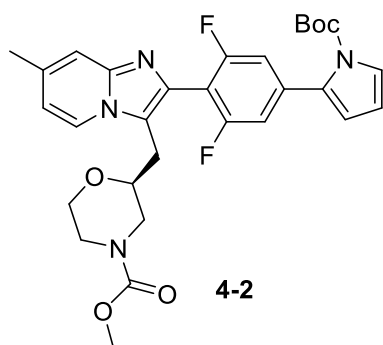
;

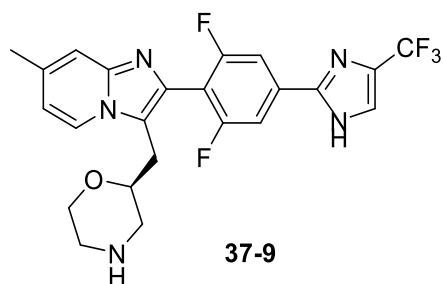
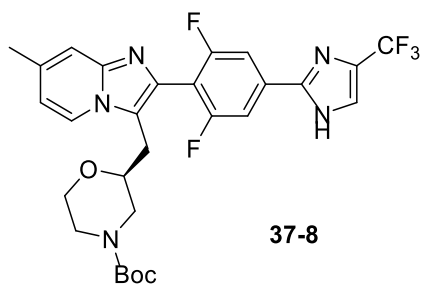
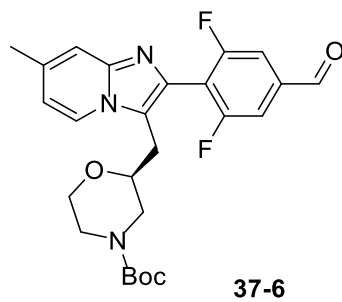
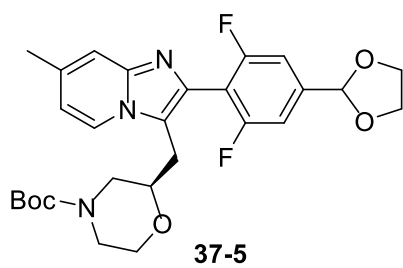
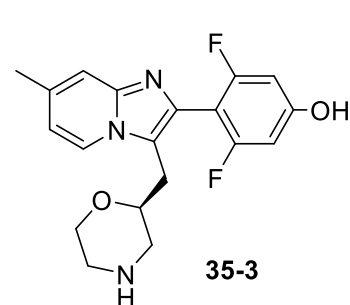
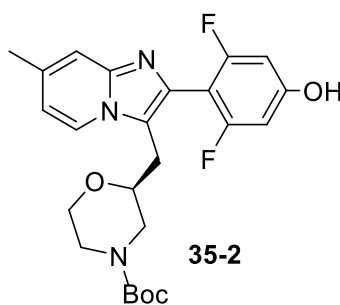
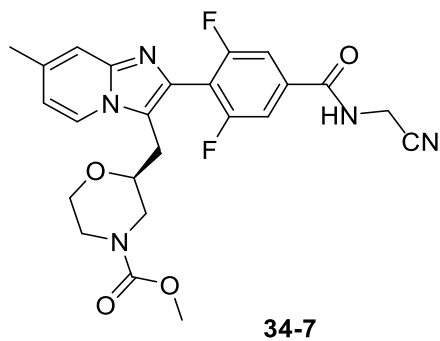
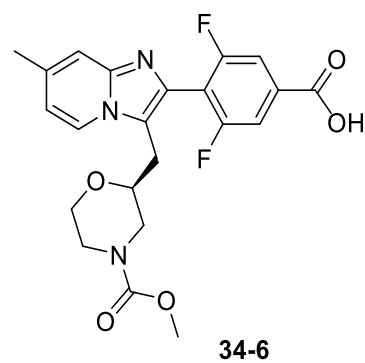
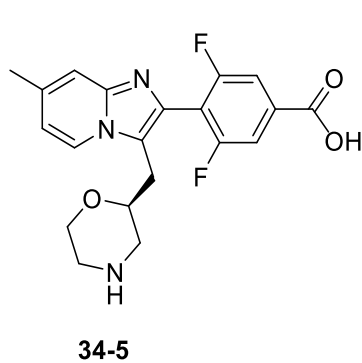
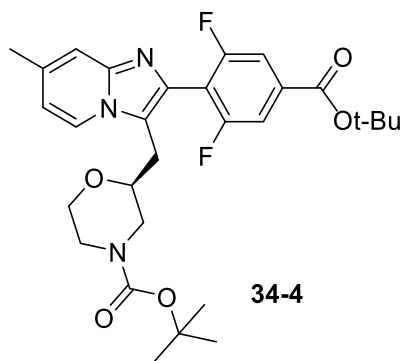
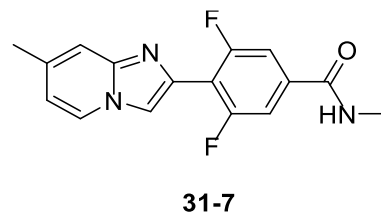
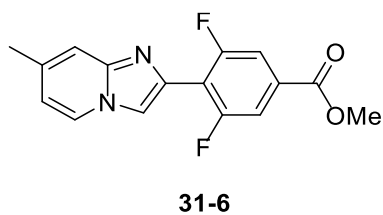
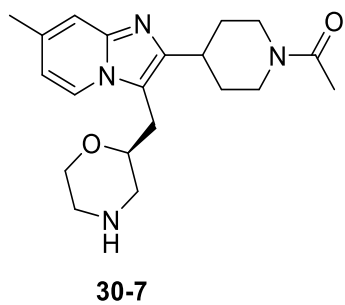


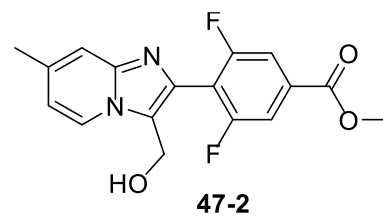
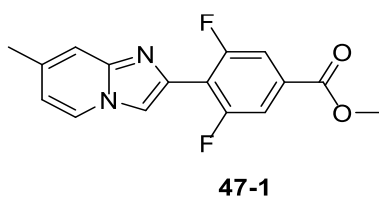
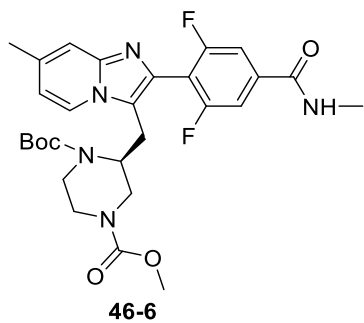
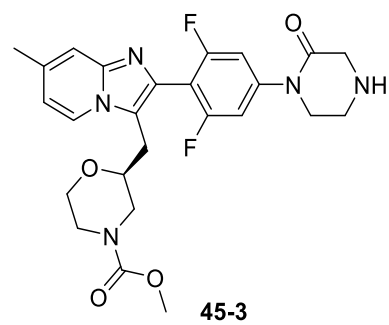
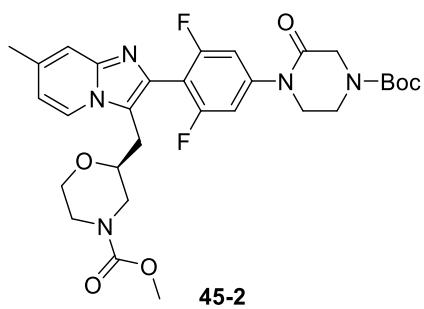
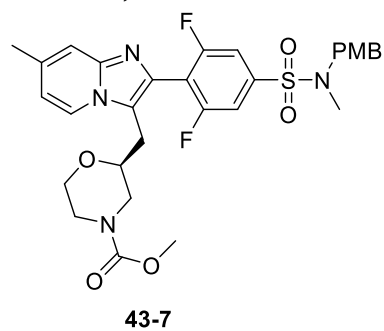
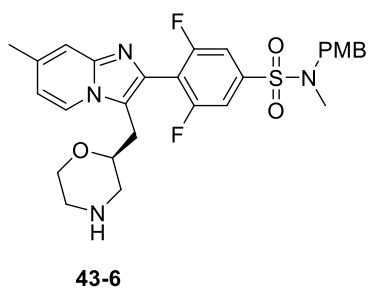
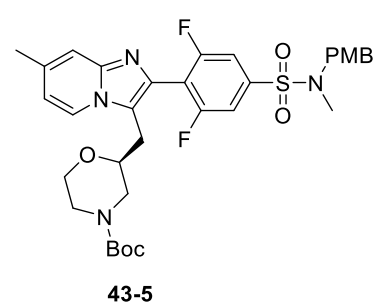
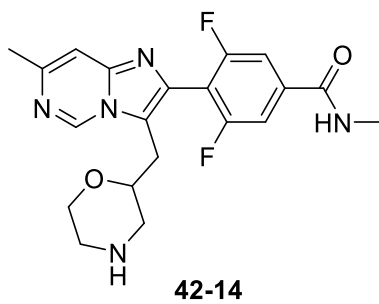
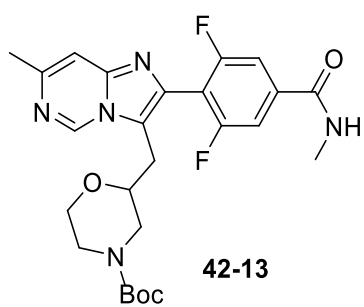
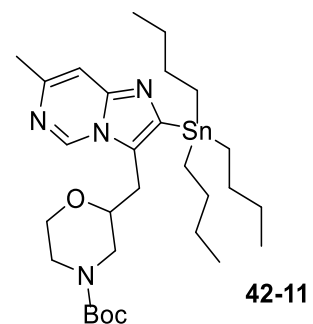
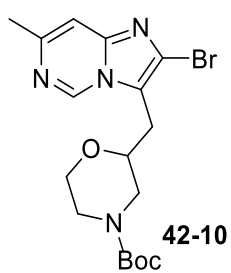
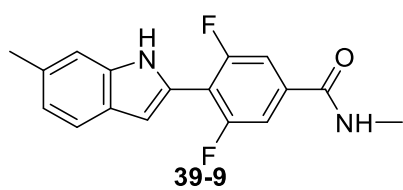
wherein the $Z^1, Z^2, Z^3, Z^4, Z^5, Z^6, Z^7, Z^8, Z^9$, $\textcircled{B}$, $R^3, R^2, R^1, W, X, Y, R^{2-1}, R^{2-2}, n$ and m have the definitions as defined in any one of claims 1 to 16, and X^1, X^5, X^6, X^6' and $X^{6''}$ have the definitions as defined in claim 15.

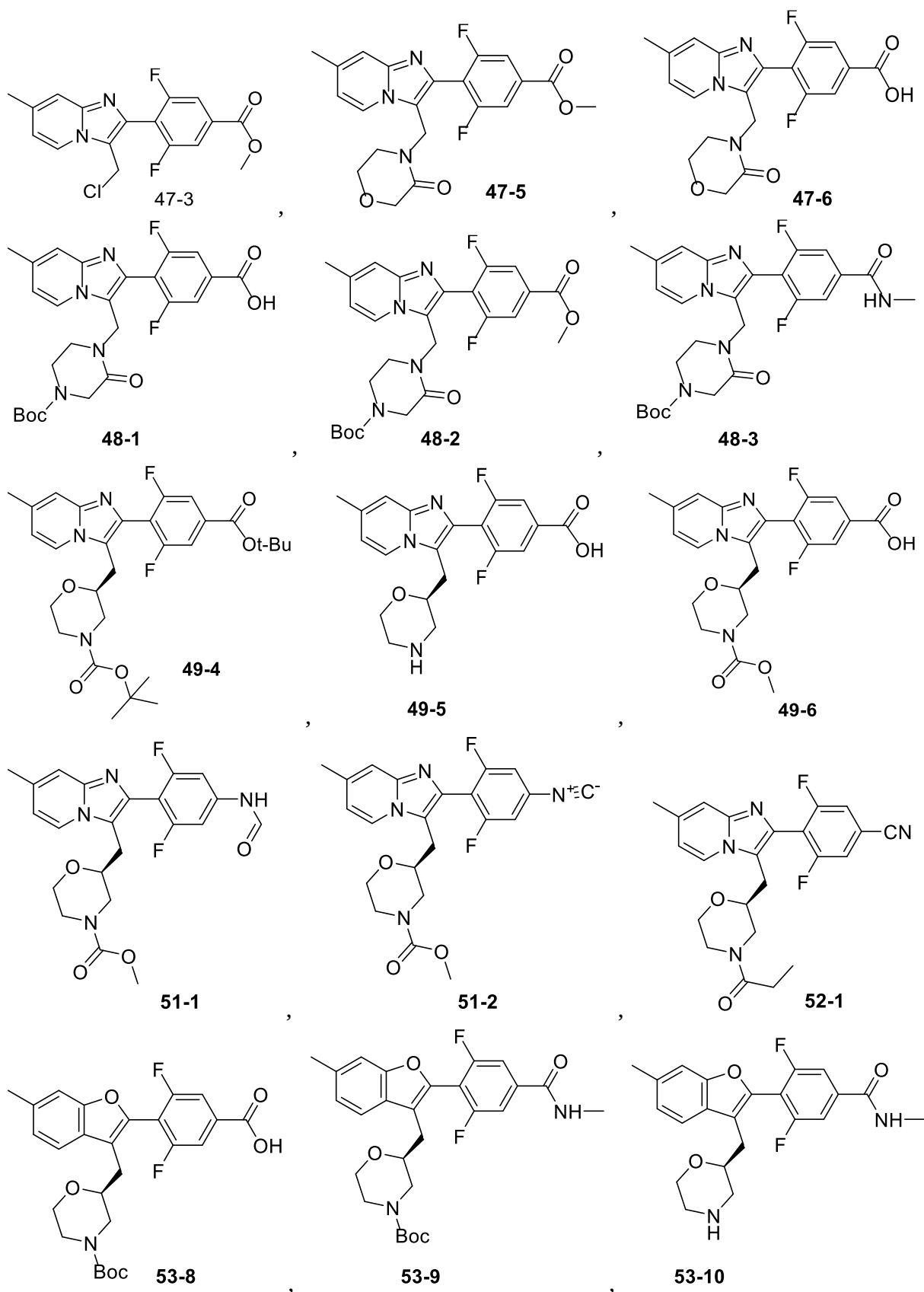
19. Compounds as shown below:

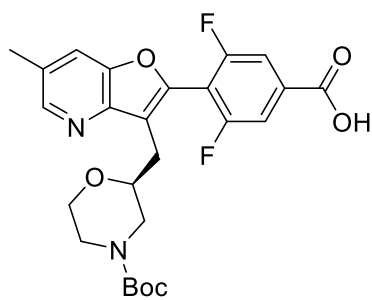




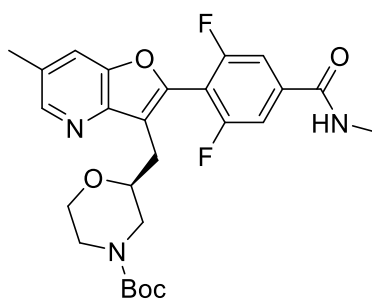




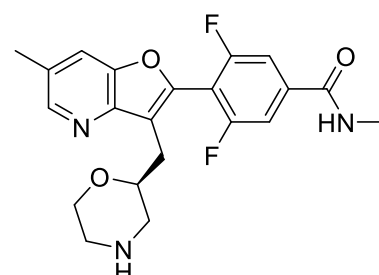




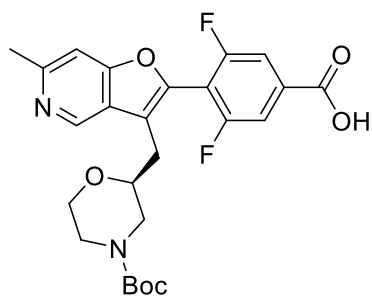
54-7



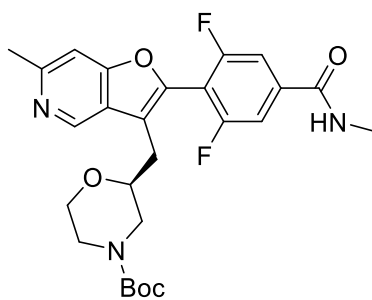
54-8



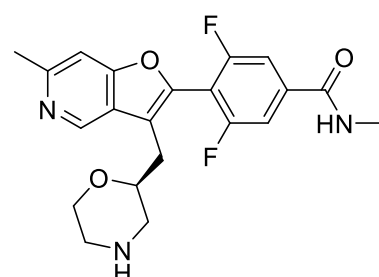
54-9



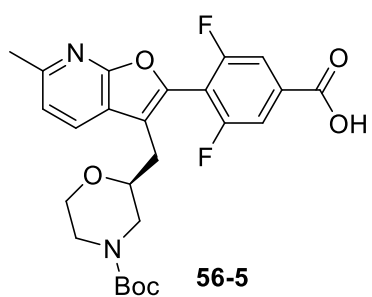
55-6



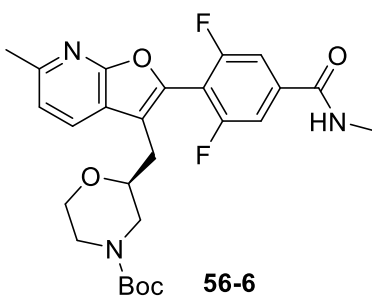
55-7



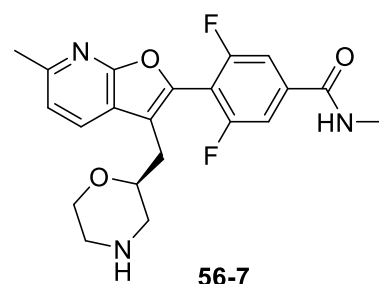
55-8



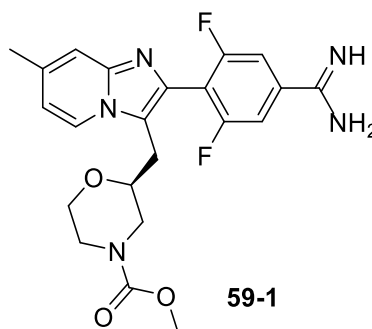
56-5



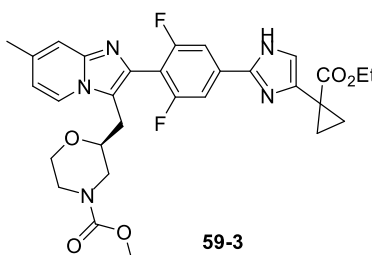
56-6



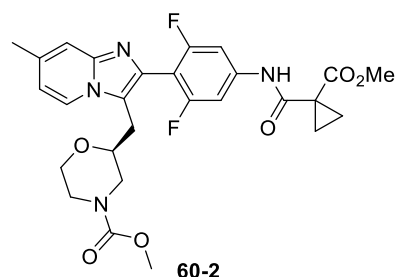
56-7



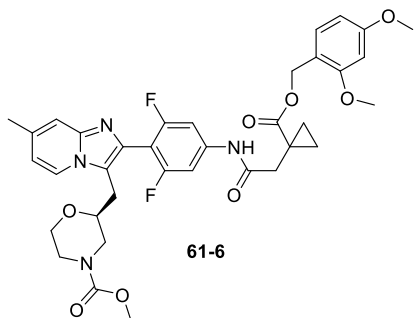
59-1



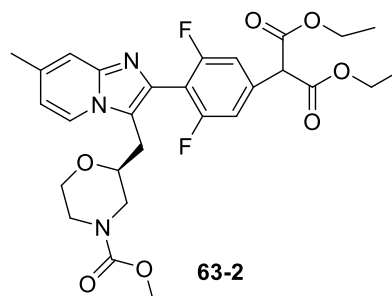
59-3



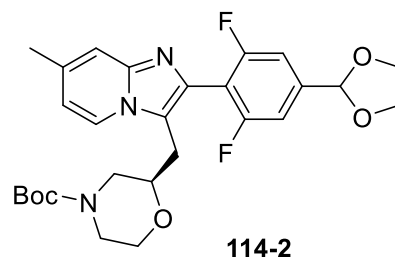
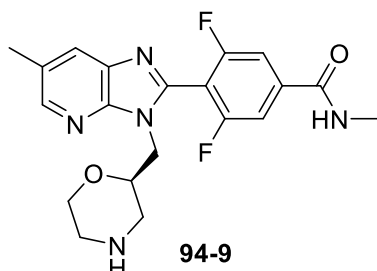
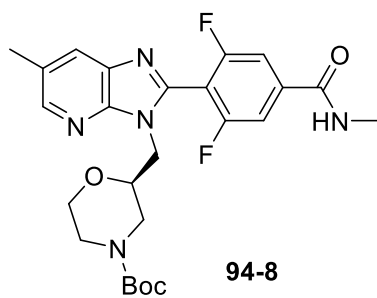
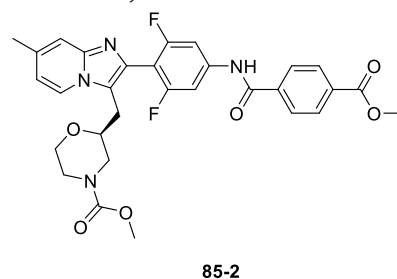
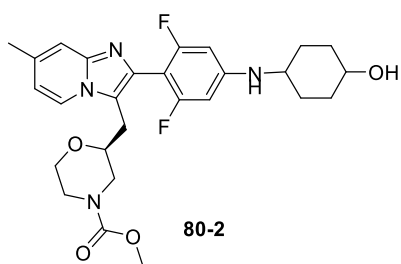
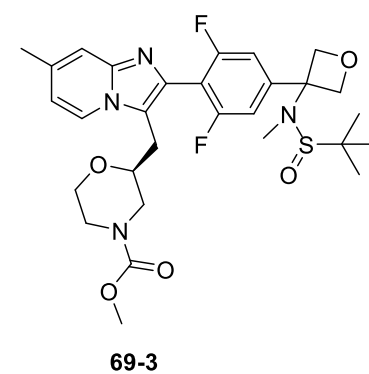
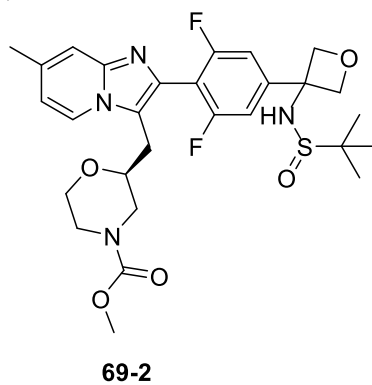
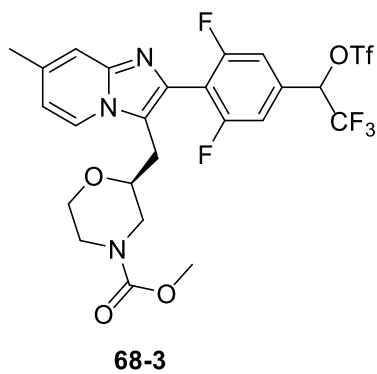
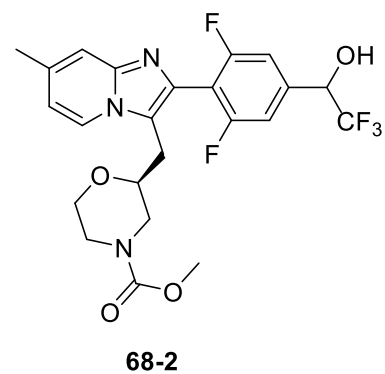
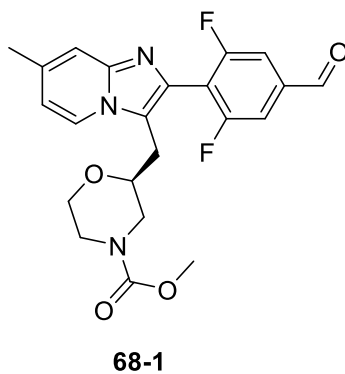
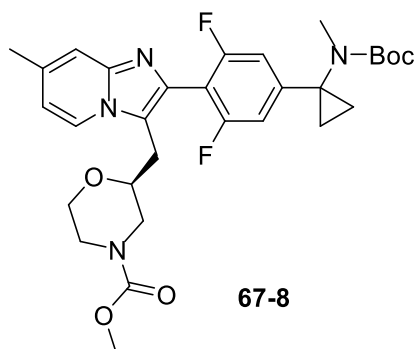
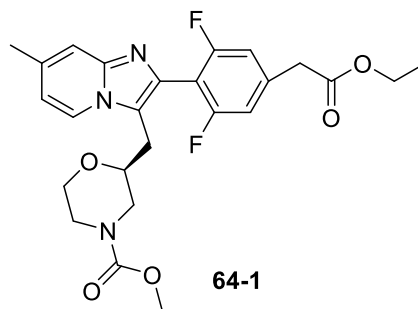
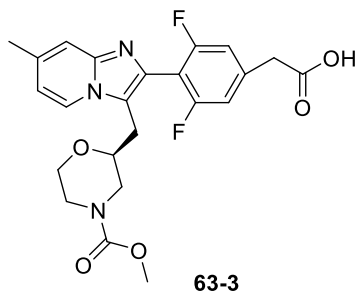
60-2

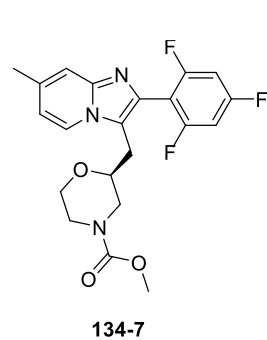
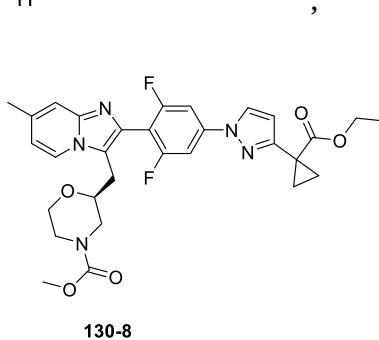
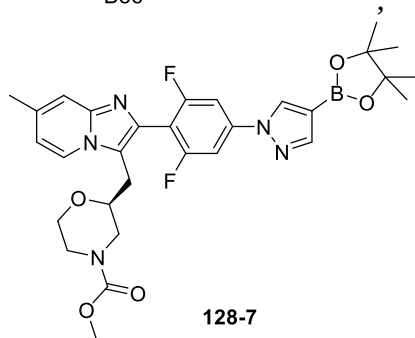
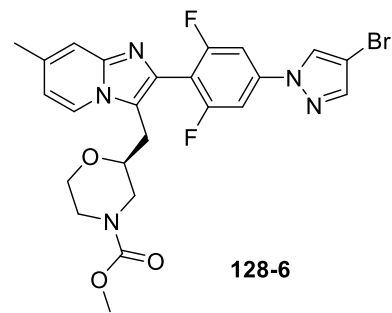
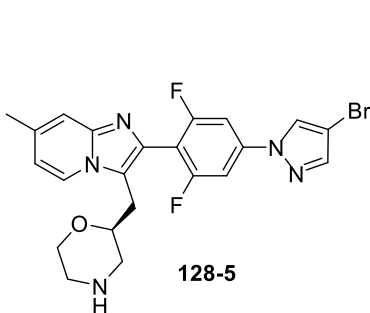
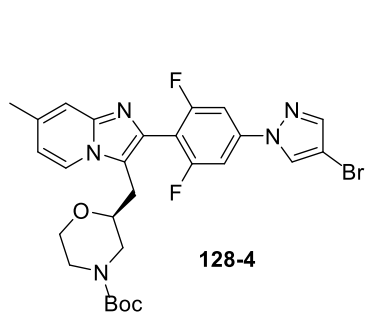
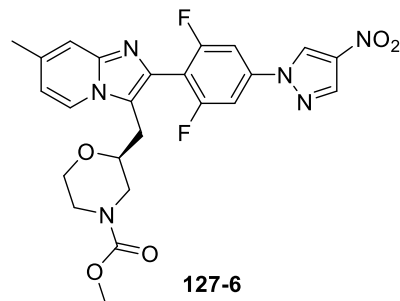
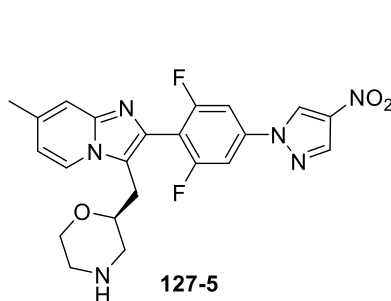
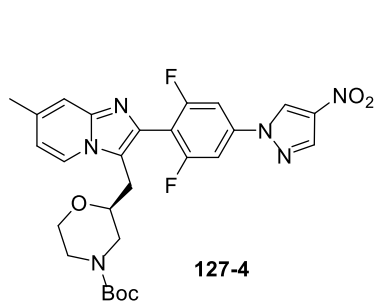
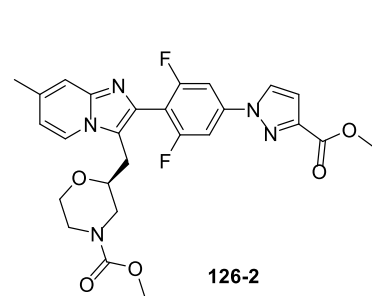
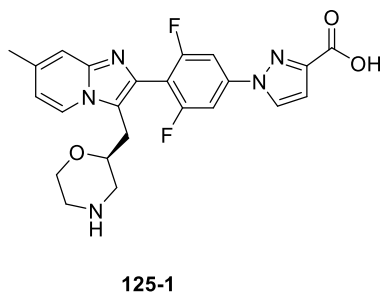
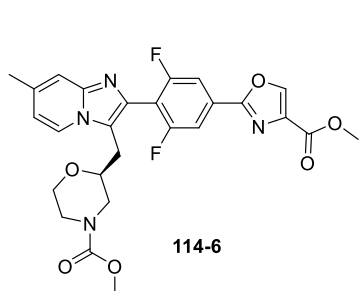
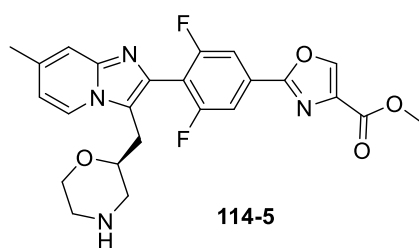
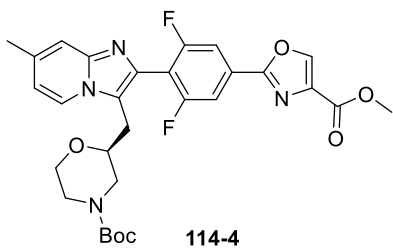
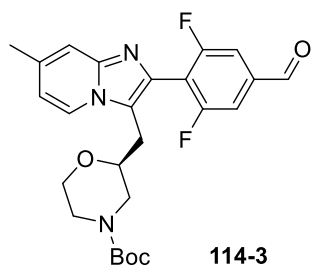


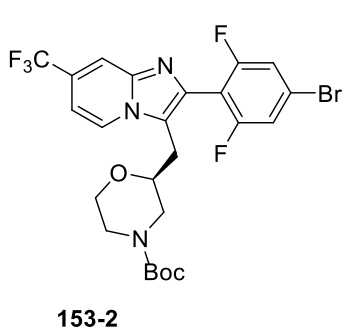
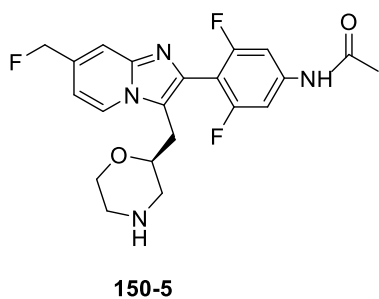
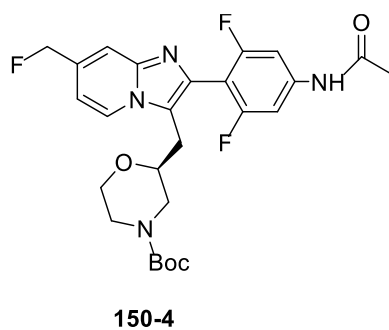
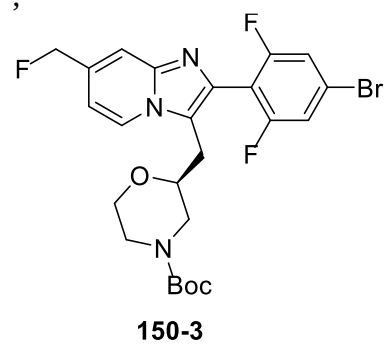
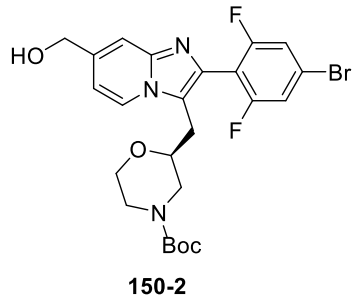
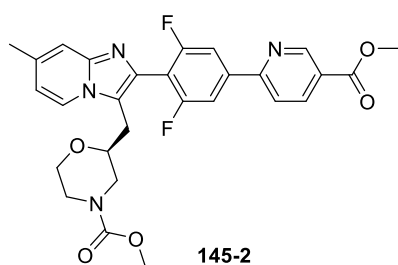
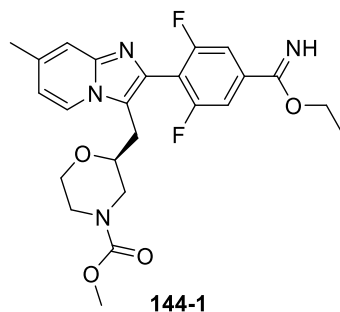
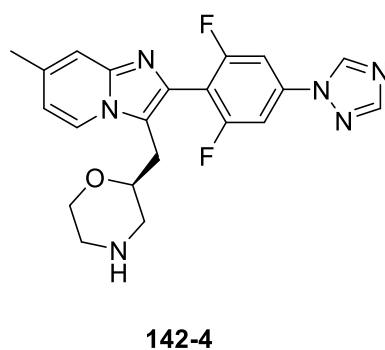
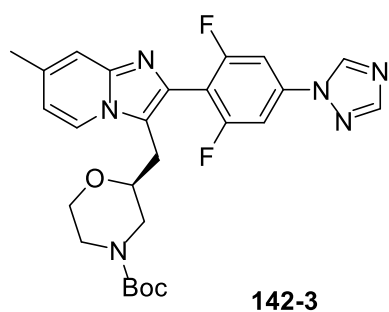
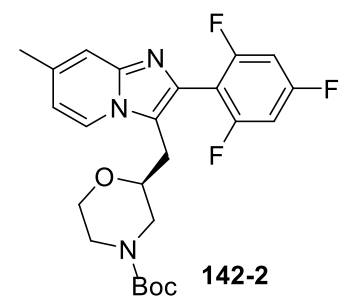
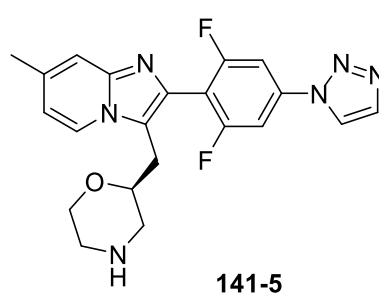
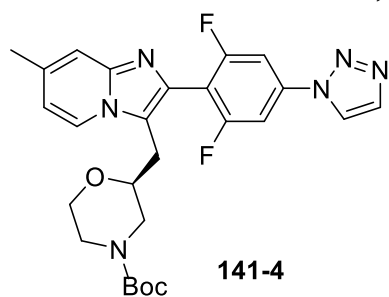
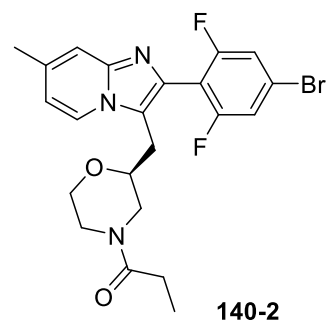
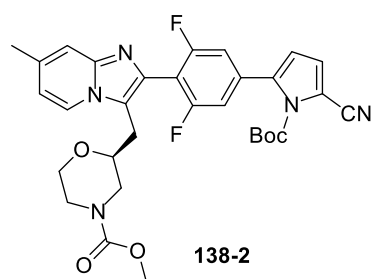
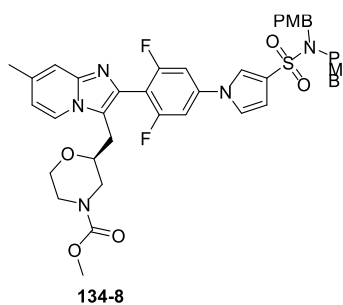
61-6

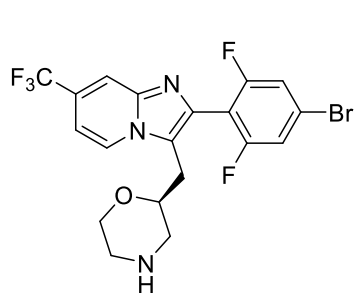


63-2

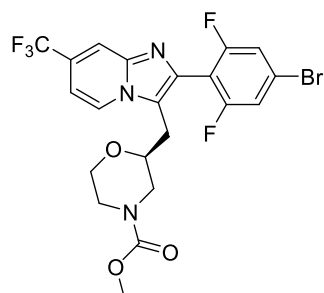




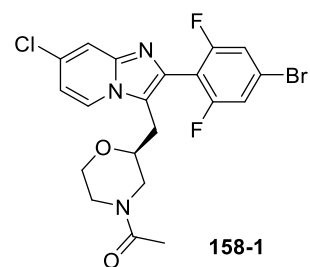




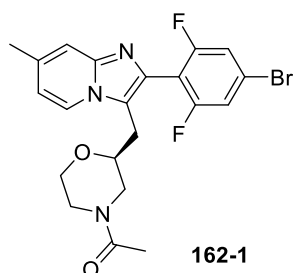
153-3



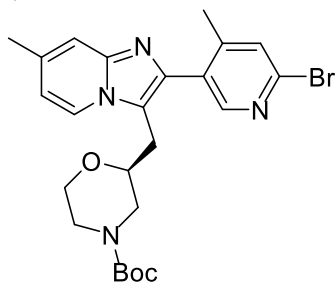
153-4



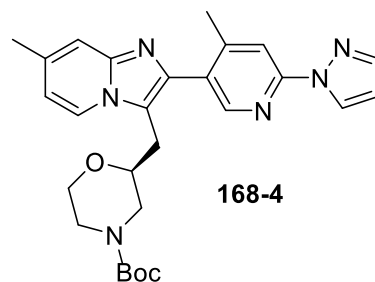
158-1



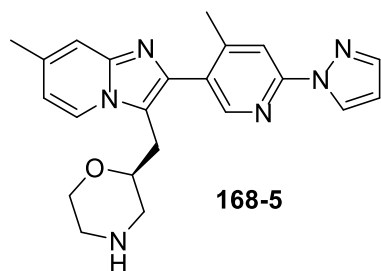
162-1



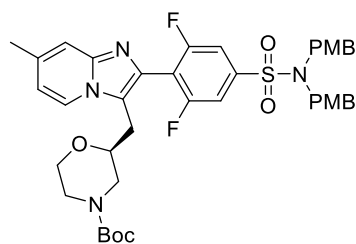
168-3



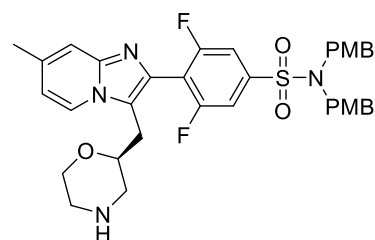
168-4



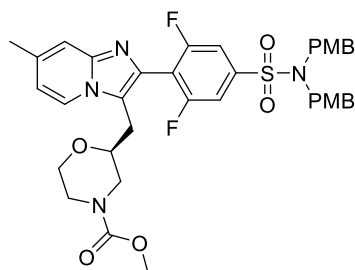
168-5



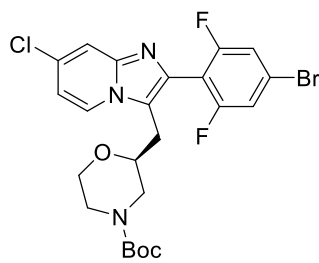
171-5



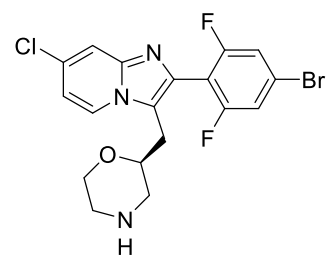
171-6



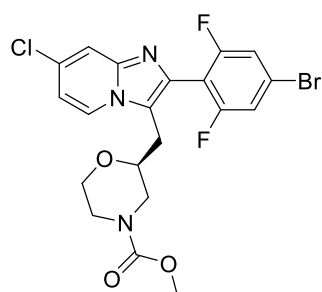
171-7



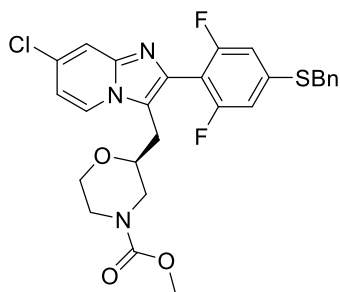
172-3



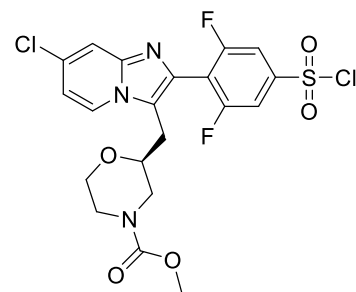
172-4



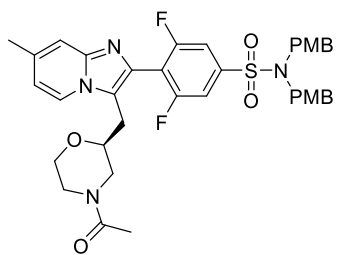
172-5



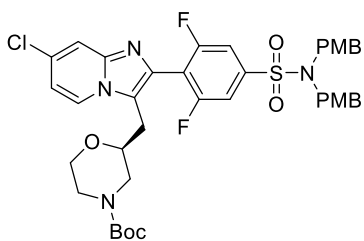
172-6



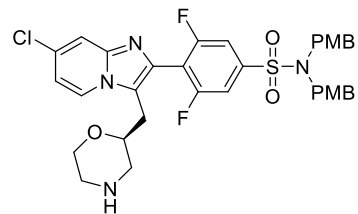
172-7



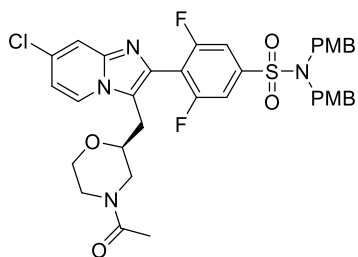
173-1



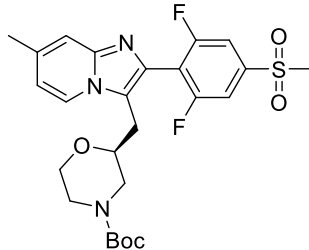
174-1



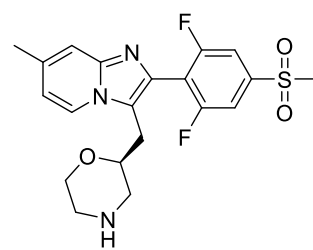
174-2



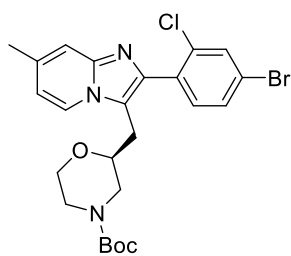
174-3



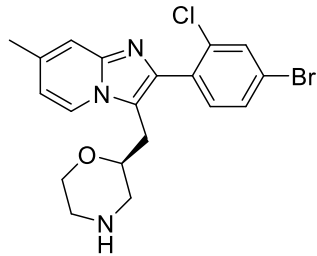
175-1



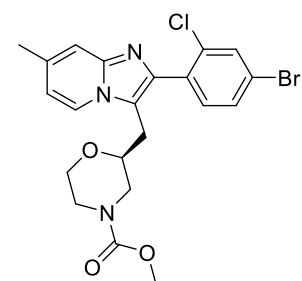
175-2



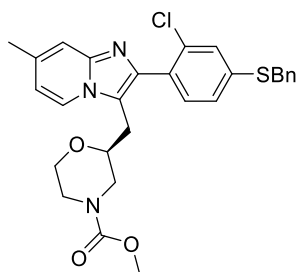
176-2



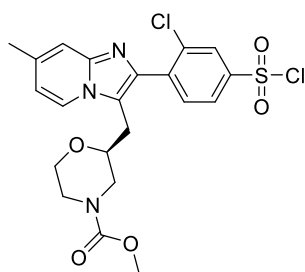
176-3



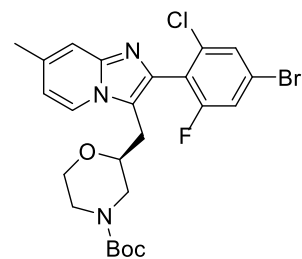
176-4



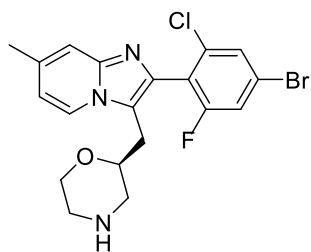
176-5



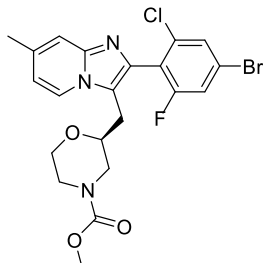
176-6



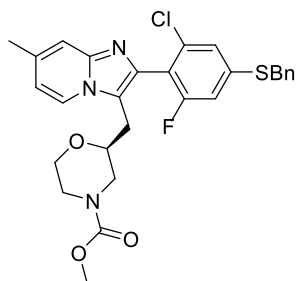
177-3



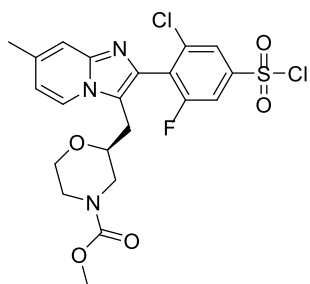
177-4



177-5

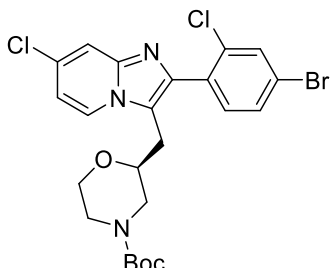


177-7



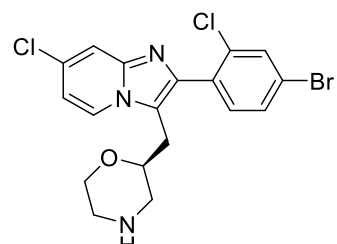
177-8

,



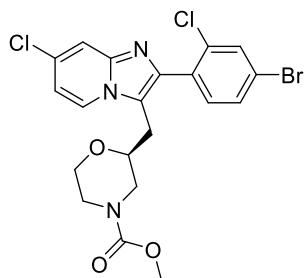
178-2

,



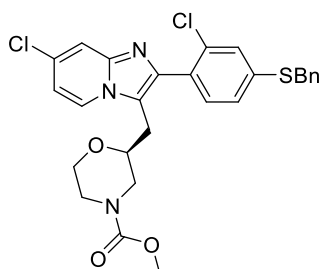
178-3

,



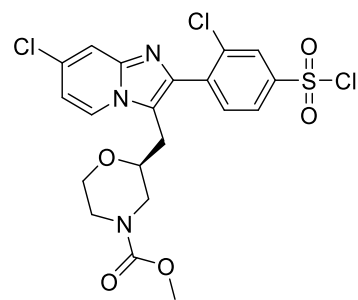
178-4

,



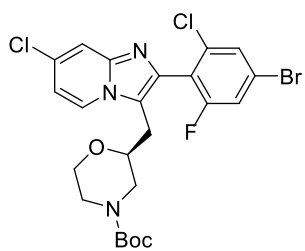
178-5

,



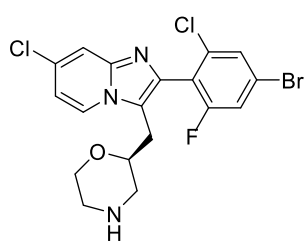
178-6

,



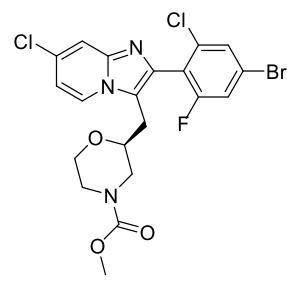
179-2

,



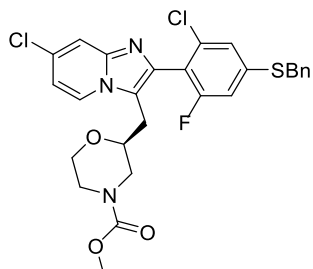
179-3

,



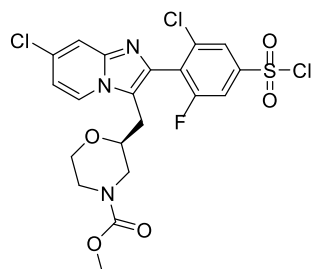
179-4

,



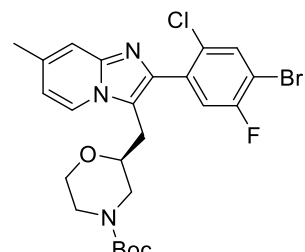
179-5

,



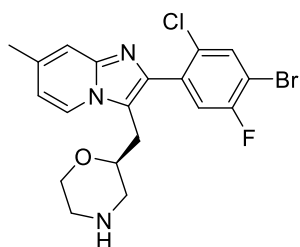
179-6

,



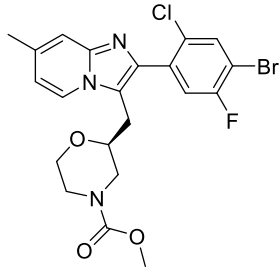
180-2

,



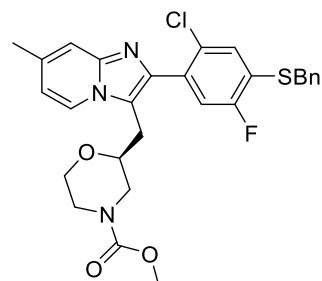
180-3

,



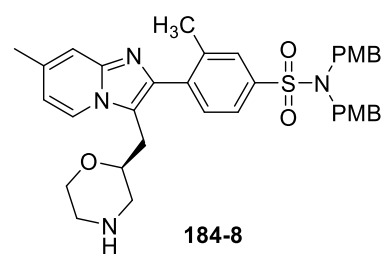
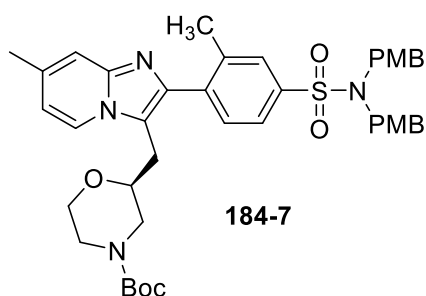
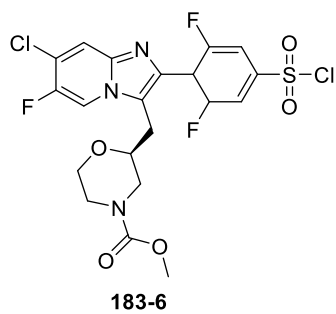
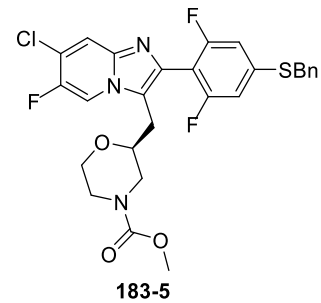
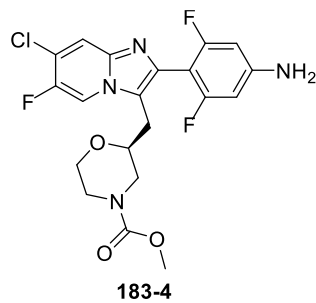
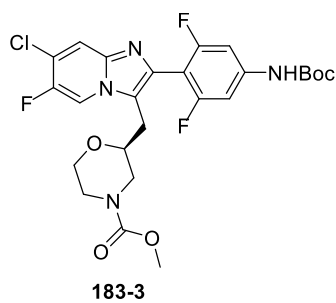
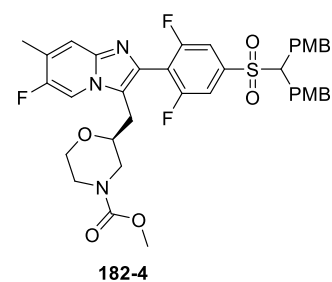
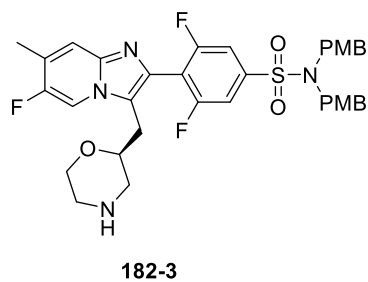
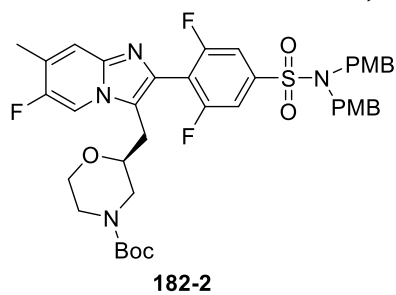
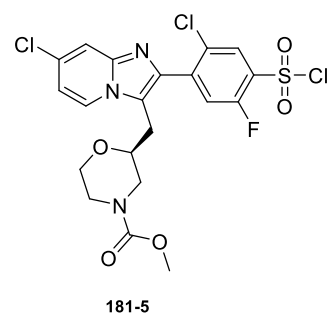
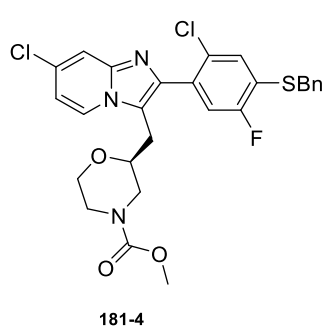
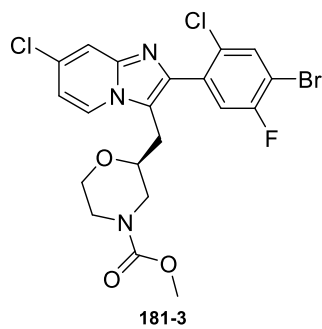
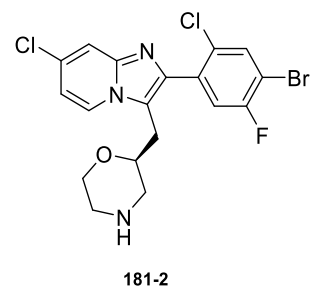
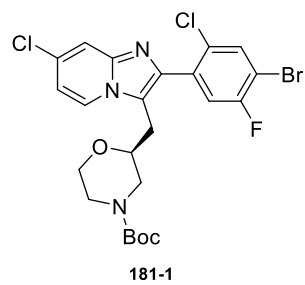
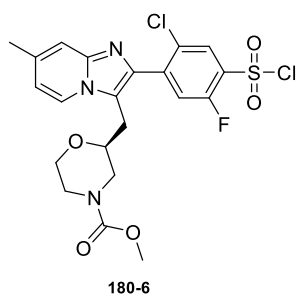
180-4

,



180-5

,



surgical pain, visceral pain, dental pain, premenstrual pain, central pain, pain resulting from burns, migraine or cluster headache; the urinary tract diseases comprise: urinary incontinence, overactive bladder, dysuria or cystitis; and the respiratory diseases preferably comprise: respiratory disorders including idiopathic pulmonary fibrosis, chronic obstructive pulmonary disease, asthma, bronchospasm or chronic cough.

5