

SYNTHESIS OF NOVEL EP4 ANTAGONIST AND USE IN CANCERS AND INFLAMMATIONS

CROSS-REFERENCE TO RELATED APPLICATIONS

[0001] This application is based on and claims the priority of Chinese Patent Application No. 202010144983.1, filed on March 4, 2020, which is incorporated herein by reference in its entirety.

FIELD

[0002] The present disclosure relates to the chemistry and medicine field, and in particular, to a pyrazole derivative and use thereof.

BACKGROUND

[0003] Prostaglandin E₂ (PGE₂) is an endogenous bioactive lipid. PGE₂ activates prostaglandin receptors to cause a broad upstream and downstream dependent biological response (Legler, D. F. et al, hit. J Biochem. Cell Biol. 2010, 42, p. 198-201), involved in the regulation of numerous physiological and pathological processes including inflammation, pain, renal function, cardiovascular system, pulmonary function, and cancer. It is reported that PGE₂ is highly expressed in cancerous tissues of various cancers, and it has been confirmed that PGE₂ is associated with the occurrence, growth and development of cancer and disease conditions in patients. It is generally believed that PGE₂ is associated with the activation of cell proliferation and cell death (apoptosis) and plays an important role in the processes of cancer cell proliferation, disease progression and cancer metastasis.

[0004] The receptors of PGE₂ are divided into 4 subtypes, i.e., EP1, EP2, EP3 and EP4, which are widely distributed in various tissues. Among these subtypes, PGE₂ effects on the EP4 receptor to interfere with inflammatory responses (including immunoinflammatory responses),

smooth muscle relaxation, pains, lymphocyte differentiation, hypertrophy or proliferation of vascular mesangial cells, secretion of gastrointestinal mucus, and the like. Thus, it can be considered that EP4 receptor antagonists are promising as anti-inflammatory and/or analgesic agents for the treatment of diseases associated with the PGE₂-EP4 pathway, such as inflammatory diseases, diseases accompanied by various pains, and the like.

[0005] EP4 is the primary receptor involved in arthritic pain in rodent models of rheumatoid arthritis and osteoarthritis (for example, see *J. Pharmacol. Exp. Ther.*, 325, 425 (2008)), which upon activation leads to an accumulation of the intracellular signaling molecule cAMP. There have been studies detecting EP4 receptor expression on peripheral nerve endings of pain receptors, macrophages, and neutrophils, and it has been confirmed that these cell types are extremely important for endometriosis. Studies have reported that oral EP4 antagonists can reduce proteinuria in type II diabetic mice, inhibiting the progression of diabetic nephropathy. Additional studies have reported that activation of EP4 and increased production of PGE₂ in the bladder mucosa may be the important causes of overactive bladder by prostatitis, and intravesical injection of EP4 antagonists may be effective in ameliorating overactive bladder following prostatitis. Thus, selective EP4 antagonists may be useful in the treatment of arthritis, including arthritis pain as well as endometriosis, diabetic nephropathy, overactive bladder. The existing treatments for arthritis are mainly conventional non-steroidal anti-inflammatory drugs (NSAIDs) or selective COX-2 inhibitors, which can produce cardiovascular and/or gastrointestinal side effects. However, the selective EP4 antagonists are less likely to produce cardiovascular side effects.

[0006] PGE₂ continuously activates EP receptors (abundantly produced by tumor cells) in the tumor microenvironment (Ochs et al, *J Neurochem.* 2016, 136, p. 1142-1154; Zelenay, S. et al, *Cell* 2015, 162, p. 1257-1270), which promotes the accumulation of a variety of immunosuppressive cells and enhances the activity thereof. The immunosuppressive cells include type II tumor-associated macrophages (TAMs), Treg cells, and myeloid-derived suppressor cells (MDSC). One of the main features of the immunosuppressive tumor microenvironment is the presence of a large number of MDSCs and TAMs, which in turn are closely associated with low overall survival in patients with gastric, ovarian, breast, bladder, hepatocellular carcinoma (HCC), head and neck cancer, and other types of cancer. In addition,

it was reported that PGE₂ can induce immune tolerance by suppressing the accumulation of antigen-presenting dendritic cells (DCs) in tumors as well as suppressing the activation of tumor-infiltrating DCs (Wang et al, Trends in Molecular Medicine 2016, 22, p. 1-3). All of these PGE₂-mediated effects may together help tumor cells evade immune surveillance. PGE₂ plays an important role in promoting the development of tumorigenesis. Increased expression levels of PGE₂ and its related receptors EP2 and EP4 have been found in various types of malignancies, including colon cancer, lung cancer, breast cancer, and head and neck cancer, and they are often closely associated with poor prognosis (Bhooshan, N. et al. Lung Cancer 101, 88-91). Thus, selective blockade of the EP2 and EP4 signaling pathways can suppress tumorigenesis by altering the tumor microenvironment and modulating tumor immune cells.

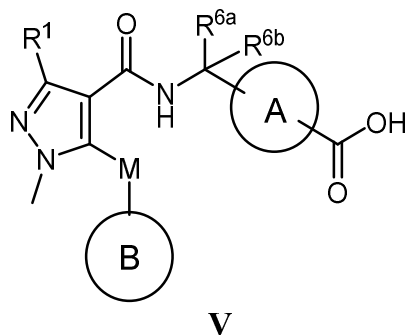
[0007] Existing preclinical research data indicate that EP2- and EP4-specific antagonists can prevent or inhibit tumor growth to varying degrees in animal models such as colon, esophageal, lung, and breast cancer. Among the PGE₂ receptor drugs entered into the clinic, Grapiprant, an EP4 antagonist developed by Pfizer, has been approved by the FDA for the treatment of arthritis in dogs, and meanwhile, it entered antitumor phase II clinical trials in 2015 for the treatment of multiple types of solid tumors such as prostate cancer, non-small cell lung cancer and breast cancer (De Vito, V. et al. J Pharm Biomed Anal 118, 251-258). E7046, an EP4 antagonist developed by Eisai, also launched a clinical phase I studies in 2015, and a phase Ib clinical trials in combination with radiotherapy or chemotherapy for rectal cancer was launched in 2017. ONO-4578, developed by Ono Pharmaceutical, entered a phase I clinical trials for advanced or metastatic solid tumors in 2017, and a phase I/II clinical trials for the treatment of advanced solid tumors either alone or in combination with nivolumab in 2018.

[0008] At present, EP4 antagonists have made some progress in the treatment of inflammatory diseases, pain, cancer, etc. However, it is still urgent to develop new drugs as improvements or replacements for current drugs.

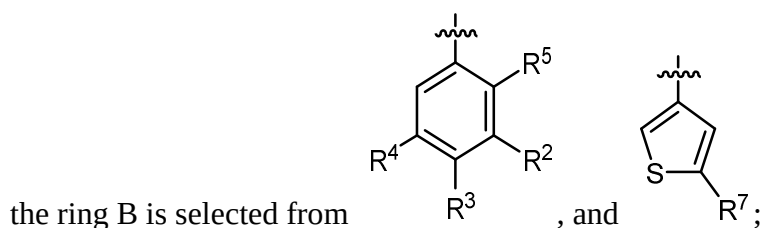
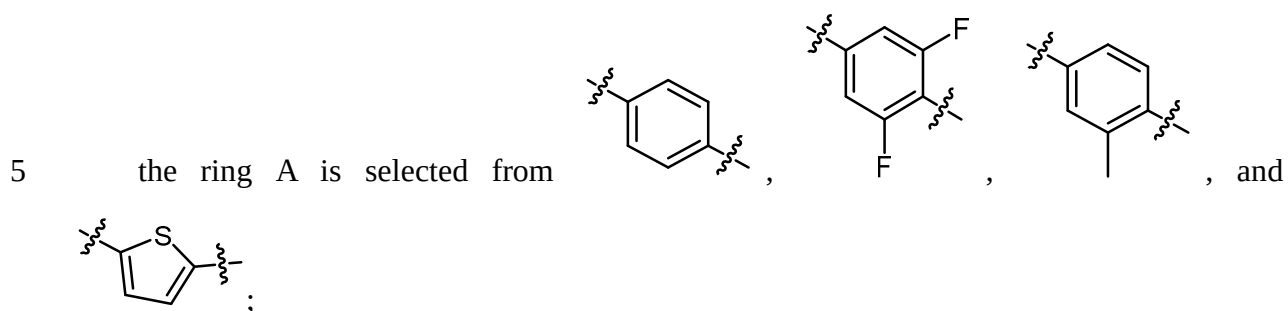
SUMMARY

[0009] The present disclosure provides a compound capable of effectively antagonizing EP4, which can be used as an improvement or replacement of the current drugs or EP4 antagonists.

[0010] To this end, in a first aspect, the present disclosure provides a compound, which is a compound represented by Formula V, or a tautomer, stereoisomer, hydrate, solvate, salt or prodrug of the compound represented by Formula V:



, in which



R^1 is selected from $-CH_3$, $-CHF_2$, and $-CF_3$;

R^2 is selected from C_2 - C_6 alkyl, C_3 - C_6 cycloalkyl, phenyl, trifluoromethyl, C_2 - C_6 halogen-substituted alkyl, C_3 - C_6 halogen-substituted cycloalkyl, C_2 - C_6 hydroxy-substituted alkyl, C_2 - C_6 cyano-substituted alkyl, $-SF_5$, and $-X-R^{2a}$, where X is selected from oxygen, sulfur, $-CO-$, $-SO_2-$, and $SO-$, and R^{2a} is selected from C_1 - C_6 alkyl and C_1 - C_6 halogen-substituted alkyl;

R^3 is selected from hydrogen, halogen, C_1 - C_2 alkyl, C_1 - C_2 fluorine-substituted alkyl, and phenyl;

15 R^4 is selected from hydrogen, halogen, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_1 - C_6 halogen-substituted alkyl, and C_1 - C_6 halogen-substituted alkoxy;

R^5 is selected from hydrogen and halogen;

one of R^{6a} and R^{6b} is hydrogen, and the other one of R^{6a} and R^{6b} is methyl; or R^{6a} and R^{6b}

together form cyclobutyl;

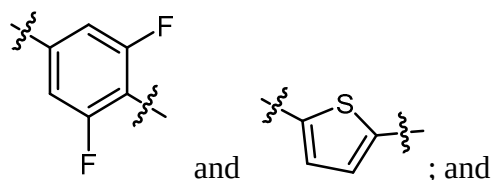
R^7 is selected from $-CH_3$, $-CHF_2$, and $-CF_3$; and

M is selected from oxygen, sulfur, and methylene;

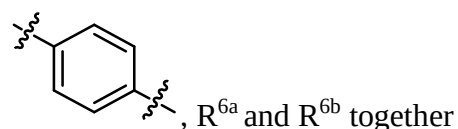
provided that:

- 5 when R^2 is trifluoromethyl, M is oxygen, one of R^{6a} and R^{6b} is hydrogen, and the other

one of R^{6a} and R^{6b} is methyl, the ring A is selected from

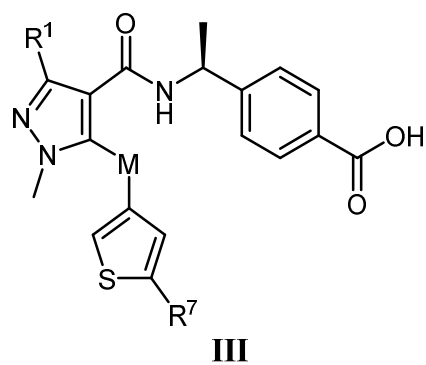


when R^2 is trifluoromethyl, M is oxygen, and the ring A is
form cyclobutyl.



[0011] According to the embodiments of the present disclosure, the above compound may
10 further include at least one of the following additional technical features.

[0012] According to the embodiments of the present disclosure, the compound is a compound
represented by Formula III, or the compound is a tautomer, stereoisomer, hydrate, solvate, salt
or prodrug of the compound represented by Formula III:

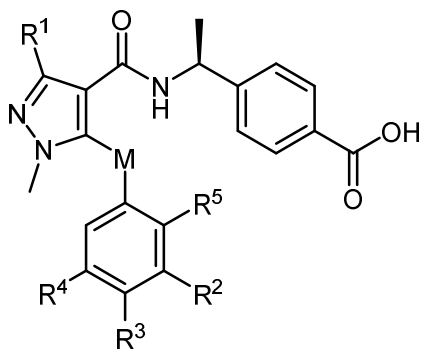


III

, in which R^1 is selected from $-CH_3$, $-CHF_2$, and $-CF_3$; R^7

- 15 is selected from $-CH_3$, $-CHF_2$, and $-CF_3$; and M is selected from oxygen, sulfur, and methylene.

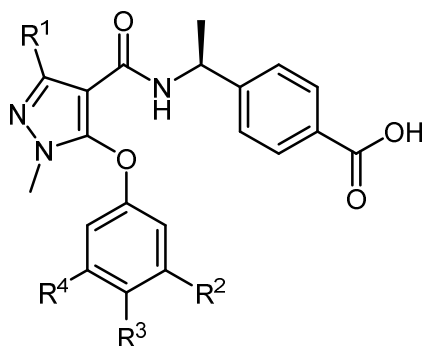
[0013] According to the embodiments of the present disclosure, the compound is a compound
represented by Formula II, or the compound is a tautomer, stereoisomer, hydrate, solvate, salt
or prodrug of the compound represented by Formula II:



II

, in which R¹ is selected from -CH₃, -CHF₂, and -CF₃, and preferably, R¹ is -CHF₂; R² is selected from ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl, fluoroethyl, fluoropropyl, fluoroisopropyl, fluorobutyl, fluoroisobutyl, hydroxyethyl, hydroxyisopropyl, cyanomethyl, cyanoethyl, phenyl, -SF₅, and -X-R^{2a}, where X is selected from oxygen, sulfur, and -CO-, and R^{2a} is selected from methyl, ethyl, fluoromethyl, and fluoroethyl; R³ is selected from hydrogen, fluorine, chlorine, methyl, ethyl, fluoromethyl, fluoroethyl, and phenyl; R⁴ is selected from hydrogen, fluorine, chlorine, methyl, ethyl, fluoromethyl, and fluoroethyl; R⁵ is selected from hydrogen, fluorine, and chlorine; and M is selected from oxygen, sulfur, and methylene.

[0014] According to the embodiments of the present disclosure, the compound is a compound represented by Formula I (also referred as to compound I), or the compound is a tautomer, stereoisomer, hydrate, solvate, salt or prodrug of the compound represented by Formula I:



I

, in which R¹ is selected from -CH₃, -CHF₂, and -CF₃; R² is selected from C₂-C₆ alkyl, C₃-C₆ cycloalkyl, C₂-C₆ halogen-substituted alkyl, and C₃-C₆ halogen-substituted cycloalkyl; R³ is selected from hydrogen, halogen, C₁-C₂ alkyl, and C₁-C₂ fluorine-substituted alkyl; R⁴ is selected from hydrogen, halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₆ halogen-substituted alkyl, and C₁-C₆ halogen-substituted alkoxy.

[0015] According to the embodiments of the present disclosure, the above compound may

further include at least one of the following additional technical features.

[0016] According to the embodiments of the present disclosure, R^2 is selected from C_2 - C_3 alkyl, C_3 - C_6 cycloalkyl, C_2 - C_3 fluorine-substituted alkyl, and C_3 - C_6 fluorine-substituted cycloalkyl.

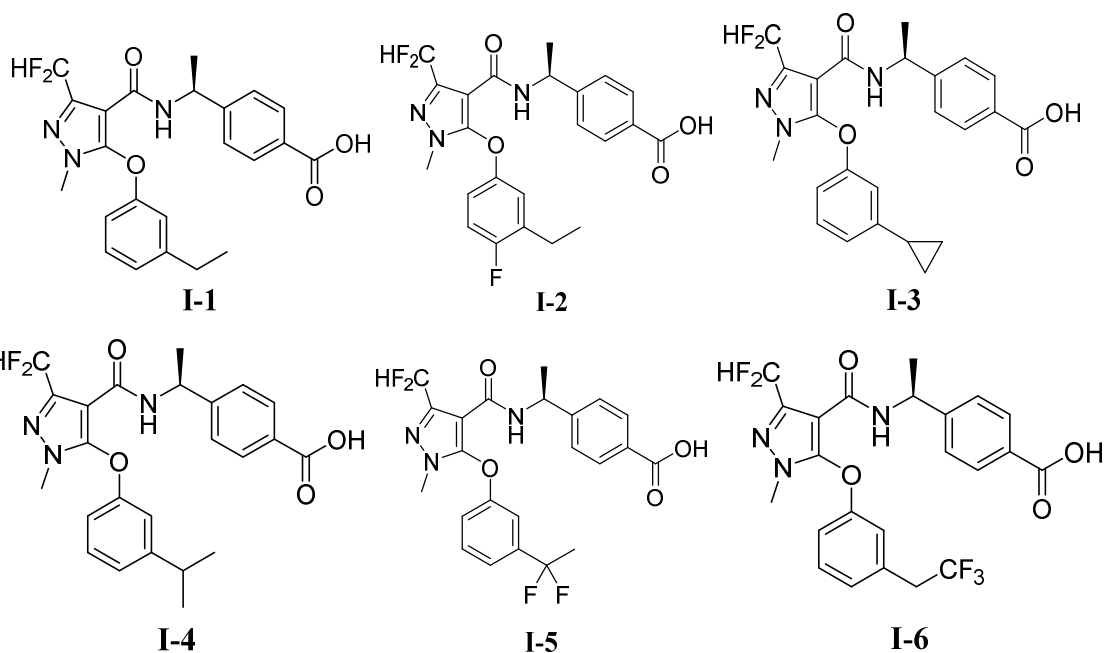
5 **[0017]** According to the embodiments of the present disclosure, R^2 is preferably selected from $-CH_2CH_3$, $-CH(CH_3)_2$, cyclopropyl, $-CF_2CH_3$, and $-CH_2CF_3$.

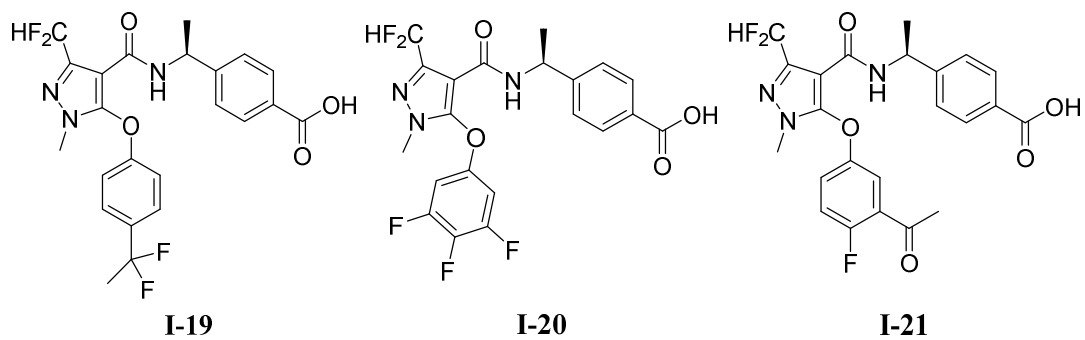
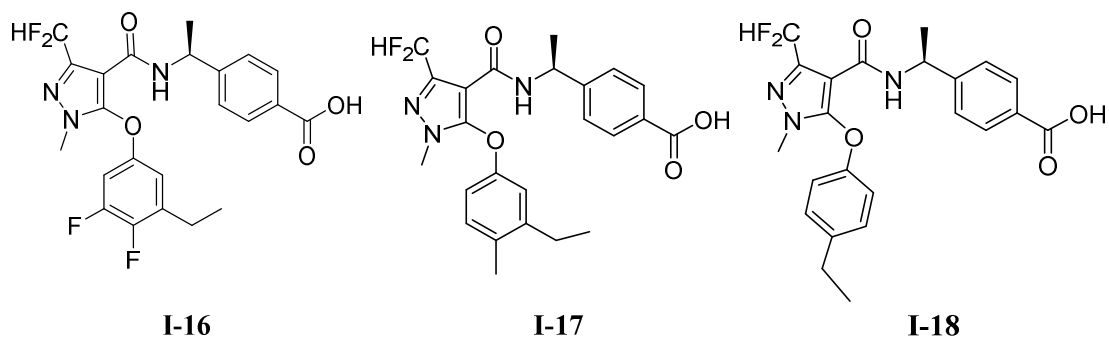
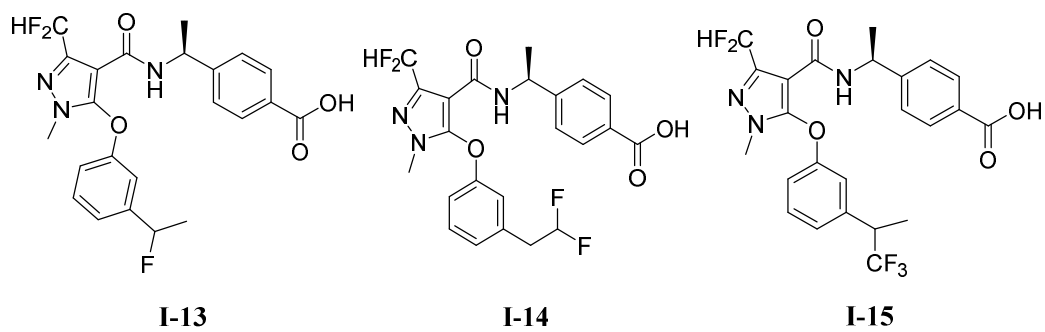
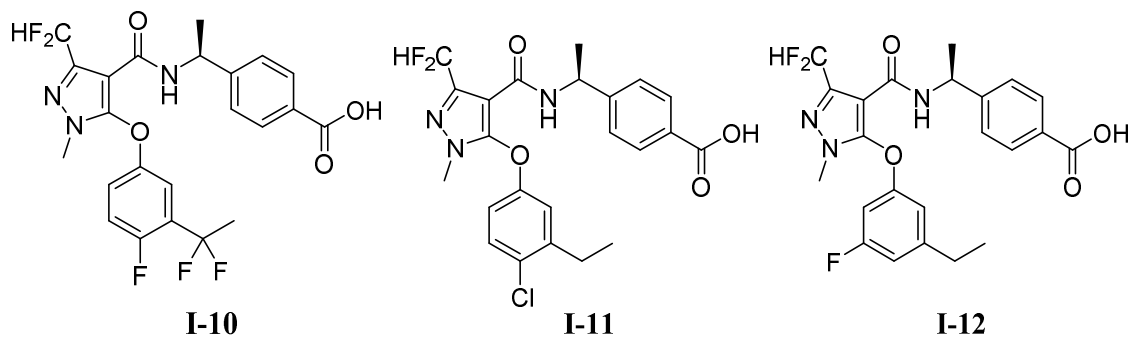
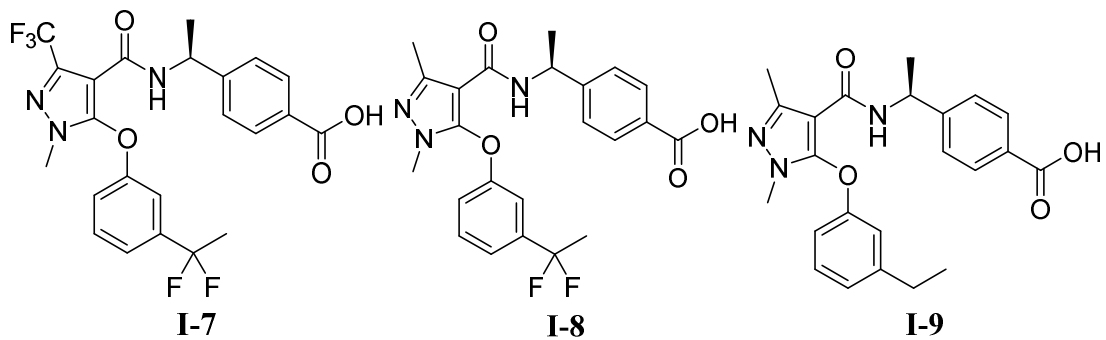
[0018] According to the embodiments of the present disclosure, R^3 is selected from hydrogen, fluorine, and chlorine.

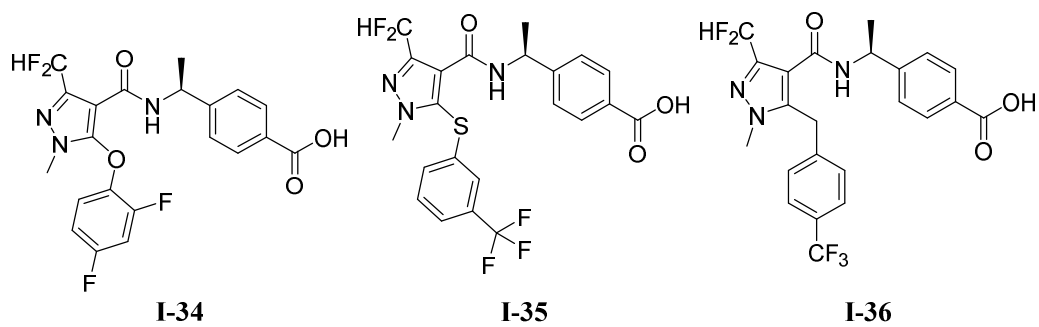
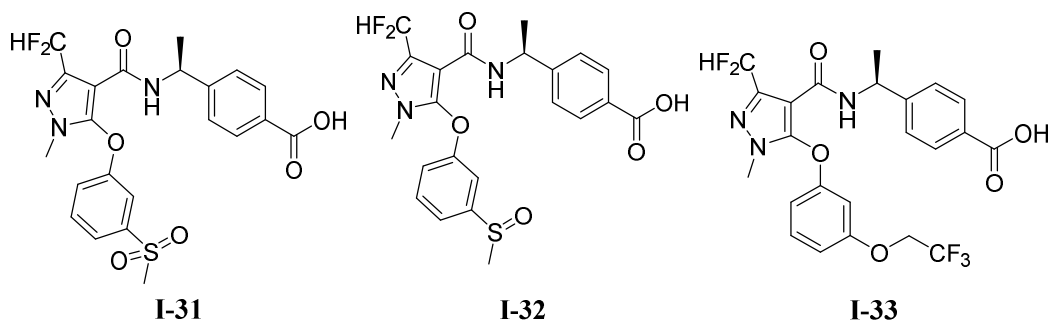
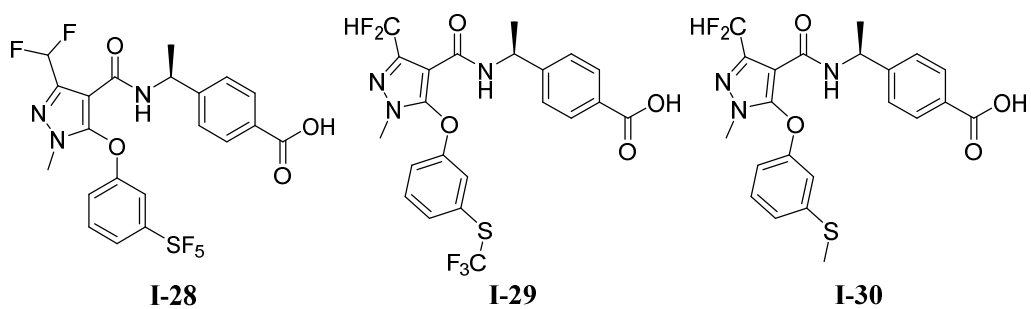
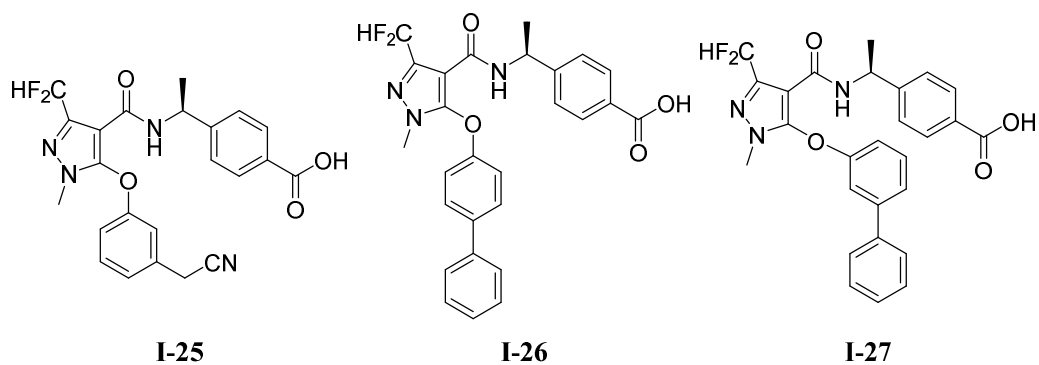
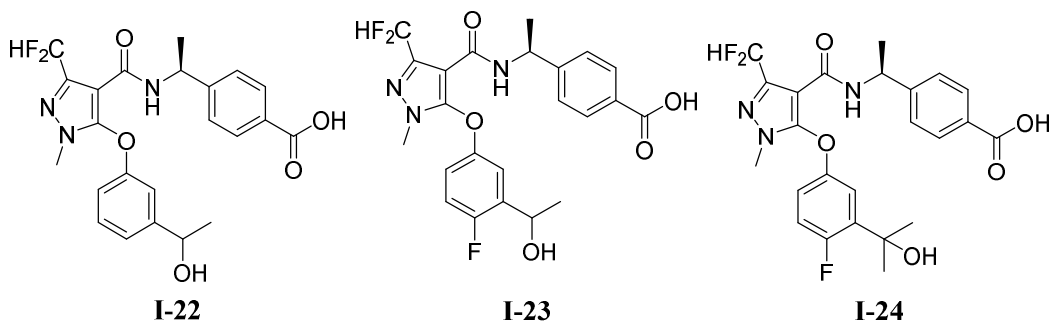
10 **[0019]** According to the embodiments of the present disclosure, R^4 is selected from hydrogen, halogen, C_1 - C_6 alkyl, C_1 - C_6 alkoxy, C_1 - C_6 halogen-substituted alkyl, and C_1 - C_6 halogen-substituted alkoxy.

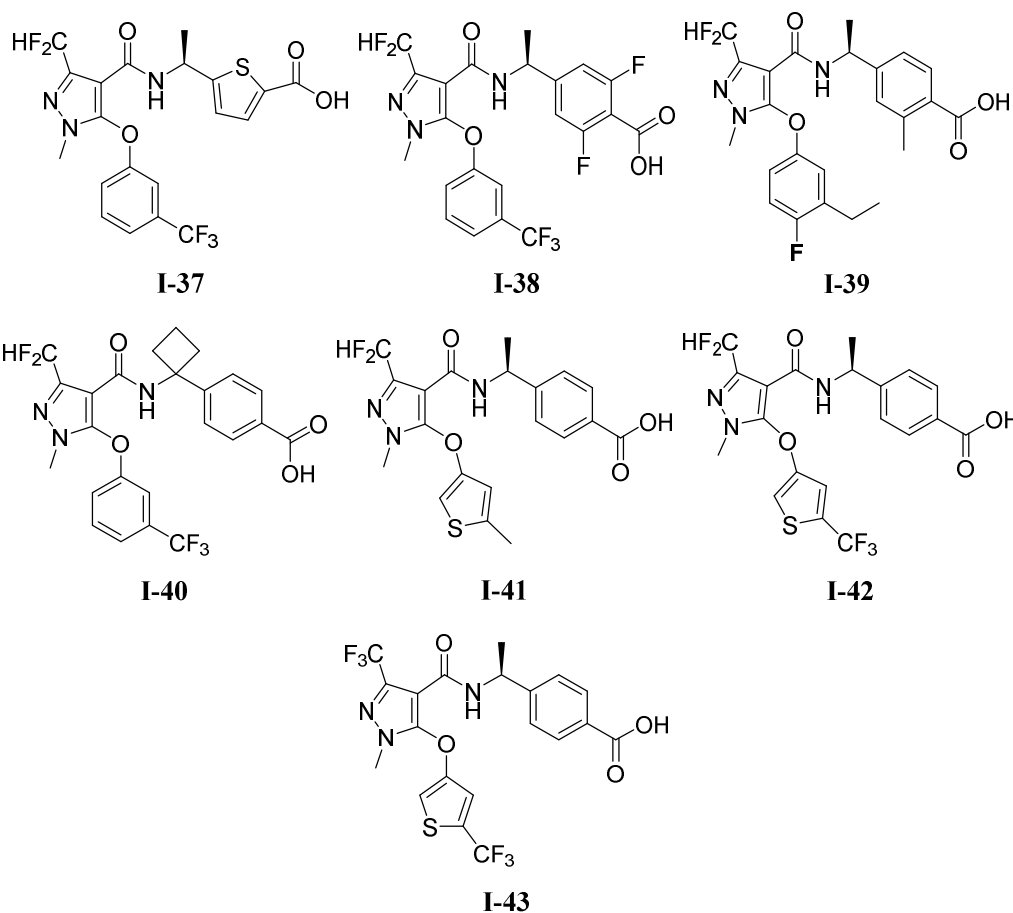
[0020] According to some embodiments of the present disclosure, R^4 is selected from hydrogen, fluorine, chlorine, C_1 - C_4 alkyl, C_1 - C_4 alkoxy, C_1 - C_4 fluorine- or chlorine-substituted alkyl, C_1 - C_4 fluorine- or chlorine-substituted alkoxy; and preferably, R^4 is selected from
15 hydrogen, fluorine, and chlorine.

[0021] According to the embodiments of the present disclosure, the compound is any one of the following compounds, or the compound is a tautomer, stereoisomer, hydrate, solvate, pharmaceutically acceptable salt or prodrug of any one of the following compounds:









[0022] According to the embodiments of the present disclosure, the salt includes a pharmaceutically acceptable salt and is at least one selected from sulfuric acid, phosphoric acid, nitric acid, hydrobromic acid, hydrochloric acid, formic acid, acetic acid, propionic acid, benzenesulfonic acid, benzoic acid, phenylacetic acid, salicylic acid, alginic acid, anthranilic acid, camphoric acid, citric acid, vinyl sulfonic acid, formic acid, fumaric acid, furoic acid, gluconic acid, glucuronic acid, glutamic acid, glycolic acid, isethionic acid, lactic acid, maleic acid, malic acid, mandelic acid, mucic acid, pamoic acid, pantothenic acid, stearic acid, succinic acid, sulfanilic acid, tartaric acid, *p*-toluenesulfonic acid, malonic acid, 2-hydroxypropionic acid, oxalic acid, glycolic acid, glucuronic acid, galacturonic acid, citric acid, lysine, arginine, aspartic acid, cinnamic acid, *p*-toluenesulfonic acid, methanesulfonic acid, ethanesulfonic acid or trifluoromethanesulfonic acid. Those skilled in the art can understand that, in addition to pharmaceutically acceptable salts, other salts can also be used in the present disclosure, acting as intermediates in the purification of compounds or in the preparation of other pharmaceutically acceptable salts, or for identifying, characterizing or purifying the compounds

of the present disclosure.

[0023] In a second aspect of the present disclosure, the present disclosure provides a pharmaceutical composition. According to the embodiments of the present disclosure, the pharmaceutical composition includes a pharmaceutically acceptable excipient, and the compound as described above.

[0024] In a third aspect of the present disclosure, the present disclosure provides uses of the compound as described above or the pharmaceutical composition as described above in the preparation of a medicament for the treatment or prevention of an EP4-related disease.

[0025] According to the embodiments of the present disclosure, the use may further include at least one of the following additional technical features.

[0026] According to an embodiment of the present disclosure, the EP4-related disease includes at least one selected from the group consisting of an inflammatory disease, a pain, a cancer, a metabolic disease and a urinary system disease.

[0027] According to an embodiment of the present disclosure, the inflammatory disease includes at least one selected from the group consisting of arthritis and rheumatoid arthritis.

[0028] According to an embodiment of the present disclosure, the pain includes osteoarthritis pain and endometriosis-induced pain.

[0029] According to an embodiment of the present disclosure, the compound or the pharmaceutical composition as described above may be administered in combination with a radiation therapy and/or an antibody therapy. The antibody therapy is selected from the group consisting of a CTLA4 antibody therapy, a PDL1 antibody therapy, a PD1 antibody therapy, and combinations thereof.

[0030] According to an embodiment of the present disclosure, the cancer includes a solid cancer.

[0031] According to embodiments of the present disclosure, the cancer includes breast cancer, cervical cancer, colorectal cancer, endometrial cancer, glioblastoma, head and neck cancer, kidney cancer, liver cancer, lung cancer, medulloblastoma, ovarian cancer, pancreatic cancer, prostate cancer, skin cancer, and urethral cancer.

[0032] According to an embodiment of the present disclosure, the metabolic disease includes diabetes, and the urinary disease includes overactive bladder.

[0033] According to an embodiment of the present disclosure, with the compound or pharmaceutical composition of the present disclosure, the patient in need thereof can be provided with a more optimal, more effective clinical treatment medication or regimen. According to the embodiments of the present disclosure, the present disclosure provides a series of EP4 antagonists having novel structures, better pharmacokinetic properties, better drug effect and good medicinal properties, which can effectively treat EP4-related diseases or disorders.

[0034] The present disclosure also relates to a method of treating a disease associated with EP4, the method including administering to a patient a therapeutically effective amount of a pharmaceutical formulation comprising a compound described herein, or a pharmaceutically acceptable salt thereof.

[0035] The present disclosure further provides a method of treating inflammatory diseases, pain, cancer, metabolic diseases, urinary system diseases. The method includes: administering, to a patient, a therapeutically effective amount of a pharmaceutical formulation containing the compound as described above or the pharmaceutically acceptable salt thereof. The present disclosure further provides a method for treating a disease by administering the compound or pharmaceutical composition in combination with a radiation therapy and/or an antibody therapy, in which the antibody therapy is selected from the group consisting of a CTLA4 antibody therapy, a PDL1 antibody therapy, a PD1 antibody therapy, and combinations thereof.

[0036] Term Definitions and Explanations

[0037] Unless otherwise stated, the definitions of groups and terms described in the specification and claims include actual definitions, exemplary definitions, preferred definitions, definitions recorded in tables, and definitions of specific compounds in the examples, etc., which can be arbitrarily combined and integrated with each other. The group definitions and compound structures that are combined and integrated should fall within the scope of the present disclosure.

[0038] Unless defined otherwise, all technical and scientific terms herein have the same meaning as commonly understood by one of ordinary skill in the art to which the claimed subject matter belongs. The patents, patent applications, publications cited herein are hereby incorporated by reference in their entireties, unless stated otherwise. When a term has multiple definitions, that defined in this chapter will prevail.

[0039] Unless otherwise indicated, conventional methods in the related art are employed, such as mass spectroscopy, NMR, IR and UV/Vis spectroscopy, and pharmacological methods. Unless specific definitions are set forth, the terms in the related description of analytical chemistry, organic synthetic chemistry, and medicinal and medicinal chemistry are those known in the art. Standard techniques may be used in chemical synthesis, chemical analysis, pharmaceutical preparation, formulation, and delivery, and treatment of patients. For example, reactions and purifications can be performed using the manufacturer's instructions of the kit, or in a manner well known in the related art or as described herein. The techniques and procedures described above may generally be performed with conventional methods well known in the art according to the description in a number of general and more specific documents cited and discussed throughout the present description. Throughout the description, groups and substituents thereof can be chosen by one skilled in the field to provide stable moieties and compounds. Where substituent groups are depicted by conventional chemical formulae, written from left to right, the substituent groups likewise encompass the chemically equivalent substituents that would result from writing the structural formula from right to left. For example, CH_2O is equivalent to OCH_2 .

[0040] As used herein, the description and claims recite numerical ranges, which, read as "integers", are to be understood as reciting both endpoints of the range and each integer within the range. For example, "an integer from 1 to 6" is to be understood as reciting each and every integer from 0, 1, 2, 3, 4, 5, and 6. When the numerical range is understood to be "a number", it is understood to recite both endpoints of the range and each integer within the range and each decimal number within the range. As an example, "a number from 1 to 10" is to be understood as reciting not only each of the integers 1, 2, 3, 4, 5, 6, 7, 8, 9, and 10, but also at least the sum of each of the integers with 0.1, 0.2, 0.3, 0.4, 0.5, 0.6, 0.7, 0.8, 0.9, respectively.

[0041] The term "pharmaceutically acceptable" means the compounds, materials, compositions and/or dosage forms that are suitable for use in contact with human and animal tissues without excess toxicity, irritation, allergic reactions or other problems or complications within the scope of reliable medical judgment, and are commensurate with a reasonable benefit/risk ratio.

[0042] The term "pharmaceutically acceptable salt" or "pharmaceutically acceptable salt

thereof" refers to salts of pharmaceutically acceptable non-toxic acids or bases, including salts of inorganic acids and bases, organic acids and bases.

[0043] In addition to the pharmaceutically acceptable salts, other salts may be adopted in the present disclosure, and they can serve as intermediates in the purification of compounds or in the preparation of other pharmaceutically acceptable salts or can be used for identifying, characterizing, or purifying the compounds of the present disclosure.

[0044] The term "stereoisomer" refers to an isomer produced by a different spatial arrangement of atoms in the molecule. The definitions and rules of stereochemistry used in the present disclosure generally follow "McGraw-Hill Dictionary of Chemical Terms (1984)", S. P. Parker, Ed., McGraw-Hill Book Company, New York; and "Stereochemistry of Organic Compounds", Eliel, E. and Wilen, S., John Wiley & Sons, Inc., New York, 1994. The compound of the present disclosure may contain an asymmetric center or chiral center, and thus different stereoisomeric forms may exist. All stereoisomeric forms of the compound of the present disclosure, including, but not limited to, diastereoisomers, enantiomers, atropisomers, geometric (or conformational) isomers, and mixtures thereof such as racemic mixtures, shall be fall within the scope of the present disclosure.

[0045] Many organic compounds exist in optically active forms, i.e., they are capable of rotating a plane of plane-polarized light. When describing optically active compounds, the prefixes *D* and *L*, or *R* and *S* are used to denote the absolute configurations of the molecule with respect to one or more chiral centers. The prefixes *D* and *L*, or (+) and (-) are symbols used to specify a rotation of plane-polarized light caused by a compound, where (-) or *L* indicates that the compound is levorotatory, and the prefix (+) or *D* indicates that the compound is dextrorotatory. For a given chemical structure, these stereoisomers are identical except that these stereoisomers are mirror images of each other. The specific stereoisomers can be referred as to enantiomers, and a mixture of such isomers is called an enantiomeric mixture. A mixture of enantiomers in 50:50 is called a racemic mixture or a racemate, which may occur when there is no stereoselectivity or stereospecificity in a chemical reaction or process.

[0046] In accordance with the selection of raw materials and methods, the compound of the present disclosure may exist in the form of one of the possible isomers or a mixture thereof, for example, as a pure optical isomer, or as a mixture of isomers such as racemic isomer and

diastereoisomeric mixture, depending on the number of asymmetric carbon atoms. The optically active (*R*)- or (*S*)-isomer can be prepared using chiral synthons or chiral preparations, or resolved using conventional techniques. If the compound contains a double bond, the substituents may be in *E*- or *Z*-configuration; if the compound contains a disubstituted cycloalkyl, the substituent of the cycloalkyl may have a *cis*- or *trans*-conformation.

[0047] When the bond with a chiral carbon in the formula of the present disclosure is depicted in a straight line, it should be understood that the two configurations (*R*) and (*S*) of the chiral carbon and both the resulting enantiomerically pure compound and mixture are included in the scope defined by the general formula. The diagrammatic presentation of the racemate or pure enantiomeric compound herein is from Maehr, J. Chem. Ed. 1985, 62: 114-120. Unless otherwise specified, the wedge bond and the dashed bond are used to represent the absolute configuration of a stereocenter.

[0048] The compounds of the present disclosure containing asymmetrically substituted carbon atoms can be separated in an optically active form or in a racemic form. The resolution of a racemic mixture of a compound can be carried out with any of a variety of methods known in the art. For example, the methods include fractional recrystallization using chiral resolving acids, which are optically active salt-forming organic acids. For example, the suitable resolving agents for fractional recrystallization are optically active acids, such as tartaric acid, diacetyl tartaric acid, dibenzoyl tartaric acid, mandelic acid, malic acid, lactic acid or various optically active camphorsulfonic acids such as the *D* and *L* forms of β -camphorsulfonic acid. Other resolving agents suitable for fractional crystallization include α -methyl-benzylamine in a pure stereoisomeric form (for example, *S* and *R* forms or a pure diastereomeric form), 2-phenylglycinol, norephedrine, ephedrine, N-methylephedrine, cyclohexylethylamine, 1,2-diaminocyclohexane, etc. The resolution of the racemic mixture can also be carried out by eluting a column filled with an optically active resolving agent (for example, dinitrobenzoylphenylglycine). High performance liquid chromatography (HPLC) or supercritical fluid chromatography (SFC) can also be employed. The specific method, elution conditions, and the chromatographic columns can be selected by those skilled in the art according to the structures of the compounds and the experimental results. Further, pure optically active starting materials or reagents with known configuration can also be used to

obtain any enantiomers or diastereomers of the compounds described in the present disclosure through stereo-organic synthesis.

[0049] Many geometric isomers of olefins, C=N double bonds, or the like may also be present in the compounds described herein, and all these stable isomers are considered in the present disclosure. When the compound described herein contains an ethylenic double bond, such a double bond includes *E*- and *Z*-geometric isomers, unless otherwise specified.

[0050] The term “tautomer” refers to an isomer of a functional group resulting from a rapid movement of an atom between two positions in a molecule. The compound of the present disclosure may exhibit tautomerism. Tautomeric compounds can be present in two or more mutually convertible species. The protonotropic tautomer are resulted from a transfer of covalently bonded hydrogen atoms between two atoms. The tautomer generally exists in an equilibrium form. When trying to separate a single tautomer, a mixture is usually produced, the physical and chemical properties of which are consistent with the mixture of compounds. The position of equilibrium depends on the intramolecularly chemical properties. For example, for many aliphatic aldehydes and ketones, such as acetaldehyde, ketonic type is dominant; and for phenols, enol type is dominant. All tautomeric forms of the compounds are included in the present disclosure.

[0051] The term “pharmaceutical composition” refers to a mixture of one or more of the compounds described herein or physiologically/pharmaceutically acceptable salts or prodrugs thereof and other chemical components. The other chemical components can be, for example, physiologically/pharmaceutically acceptable carriers and excipients. The pharmaceutical composition aims to facilitate the administration of the compound to an organism.

[0052] The term “solvate” refers to the compound of the present disclosure or a salt thereof including a stoichiometric or non-stoichiometric solvent bonded through an intermolecular non-covalent force. When the solvent is water, the solvate is a hydrate.

[0053] The term “prodrug” can be converted into the compound of the present disclosure having biological activity under physiological conditions or through solvolysis. The prodrug of the present disclosure is prepared by modifying the functional groups in the compound, and the modification moiety can be removed by conventional operations or *in vivo*, so as to obtain the parent compound. The prodrug includes a compound, which is formed by connecting a moiety

to a hydroxyl group or amino group in the compound of the present disclosure. When the prodrug of the compound of the present disclosure is administered to a mammal individual, the prodrug is dissociated to form a free hydroxyl or amino group.

[0054] The compound of the present disclosure may contain an unnatural ratio of atomic isotopes on one or more of the atoms constituting the compound. For example, the compound may be labeled with a radioisotope, such as tritium (^3H), iodine-125 (^{125}I) or C-14 (^{14}C). The transformation of all isotopic compositions of the compounds of the present disclosure, whether radioactive or not, are included within the scope of the present disclosure.

[0055] The term "excipient" refers to a pharmaceutically acceptable inert ingredient. Examples of the "excipient" include, but not limited to, binders, disintegrants, lubricants, glidants, stabilizers, fillers, diluents, and the like.

[0056] The term " $\text{C}_1\text{-C}_6$ alkyl" refers to a linear or branched, saturated, monovalent hydrocarbon group having 1, 2, 3, 4, 5 or 6 carbon atoms. Said alkyl is, for example, methyl, ethyl, propyl, butyl, pentyl, hexyl, isopropyl, isobutyl, sec-butyl, tert-butyl, isoamyl, 2-methylbutyl, 1-methylbutyl, 1-ethylpropyl, 1,2-dimethylpropyl, neopentyl, 1,1-dimethylpropyl, 4-methylpentyl, 3-methylpentyl, 2-methylpentyl, 1-methylpentyl, 2-ethylbutyl, 1-ethylbutyl, 3,3-dimethylbutyl, 2,2-dimethylbutyl, 1,1-dimethylbutyl, 2,3-dimethylbutyl, 1,3-dimethylbutyl, 1,2-dimethylbutyl, etc., or an isomer thereof. Specifically, said group has 1, 2 or 3 carbon atoms (" $\text{C}_1\text{-C}_3$ alkyl"), e.g., methyl, ethyl, n-propyl, or isopropyl.

[0057] The term " $\text{C}_3\text{-C}_6$ cycloalkyl" refers to a saturated, monovalent, mono-or bicyclic hydrocarbon ring having 3 to 6 carbon atoms, including fused or bridged polycyclic ring systems, e.g., cyclopropyl, cyclobutyl, cyclopentyl, and cyclohexyl.

[0058] The term " $\text{C}_1\text{-C}_6$ alkoxy" is to be understood as $-\text{O}(\text{C}_1\text{-C}_6 \text{ alkyl})$, in which the " $\text{C}_1\text{-C}_6$ alkyl" has the above definition.

[0059] The term "halogeno-group" or "halogen" is fluorine, chlorine, bromine, or iodine.

[0060] "Halogen-substituted alkyl" refers to a branched and straight-chain saturated aliphatic hydrocarbon group having a specified number of carbon atoms and substituted with one or more halogen atoms (e.g., $-\text{C}_v\text{F}_w$, where $v = 1$ to 3, $w = 1$ to $(2v+1)$). Examples of halogen-substituted alkyl include, but are not limited to, trifluoromethyl, trichloromethyl, pentafluoroethyl, pentachloroethyl, 2,2,2-trifluoroethyl, heptafluoropropyl, and heptachloropropyl.

[0061] Beneficial Effects

[0062] According to the embodiments of the present disclosure, the compounds and/or compositions thereof described herein have the activity of effectively antagonizing the EP4 receptor, and they have the advantages of superior hepatic metabolic stability and cardiac safety and also have better pharmacokinetic properties, higher *in vivo* exposure, lower dosing and a better compliance. Therefore, they have good application perspective in the preparation of medicaments for the treatment of EP4-related diseases.

[0063] Additional aspects and advantages of the present disclosure will be set forth in part in the following description, and in part will be obvious from the description, or may be learned by practice of the present disclosure.

BRIEF DESCRIPTION OF DRAWINGS

[0064] FIG. 1 is a result of tumor inhibition of the compounds according to an embodiment of the present disclosure.

DESCRIPTION OF EMBODIMENTS

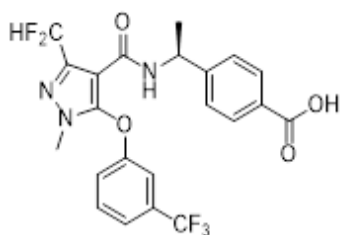
[0065] Solutions of the present disclosure will be explained below in connection with the examples. It will be appreciated by those skilled in the art that the following examples are merely illustrative of the present disclosure and should not be taken as limiting the scope of the present disclosure. Techniques or conditions that are not specified in the examples shall be performed in accordance with the techniques or conditions described in the literatures in the field or in accordance with the product instruction. Reagents or instruments without indicating the manufacturer and are those commercially available.

[0066] Compounds of the present application are identified by nuclear magnetic resonance (NMR) and/or mass spectrometry (MS), unless otherwise specified. The unit of NMR shift is 10^{-6} (ppm). Solvents for NMR were deuterated dimethyl sulfoxide, deuterated chloroform, deuterated methanol, etc., and internal standard was tetramethylsilane (TMS).

[0067] Abbreviations in the present disclosure are defined as follows:

[0068] BAST: bis(2-methoxyethyl)aminosulfur trifluoride

- [0069]** m-CPBA: m-chloroperoxybenzoic acid
- [0070]** L-selectride: lithium tri-sec-butyl borohydride
- [0071]** Pd (dppf) Cl₂: 1,1-bis(diphenylphosphino)ferrocene palladium chloride
- [0072]** DCM: dichloromethane
- 5 **[0073]** HATU: O-(7-Azabenzotriazol-1-yl)-N, N, N', N'-tetramethyluronium hexafluorophosphate
- [0074]** DIPEA: diisopropylethylamine i.e., N, N-diisopropylethylamine
- [0075]** DMF: N, N-dimethylformamide
- [0076]** N: normality, e.g., 1N hydrochloric acid means 1 mol/L hydrochloric acid solution
- 10 **[0077]** THF: tetrahydrofuran
- [0078]** DMA: N, N-dimethylacetamide
- [0079]** DMSO: dimethyl sulfoxide
- [0080]** EA: ethyl acetate
- [0081]** IC₅₀: half inhibitory concentration, indicating a concentration at which half of the
- 15 maximal inhibitory effect is achieved.
- [0082]** CHO: Chinese hamster ovary cells
- [0083]** HBSS: Hank's Balanced Salt Solution
- [0084]** BSA: albumin from bovine serum
- [0085]** HEPES: hydroxyethylpiperazine ethanethiosulfonic acid
- 20 **[0086]** IBMX: 3-isobutyl-1-methyl-7H-xanthine
- [0087]** FLIPR: fluorescence imaging plate reader
- [0088]** EC₈₀: a concentration at which 80% of maximal effect is achieved
- [0089]** Unless otherwise indicated, the compounds exemplified herein are named and numbered using ChemBioDraw Ultra 13.0.
- 25 **[0090]** Control Example 1: Preparation of Control Compound



Control compound

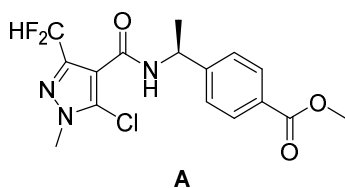
[0091]

[0092] The control compound was synthesized with reference to patent application WO2012039972A1.

[0093] The control compounds in the following test examples are all referred to as the compound as described in Control Example 1.

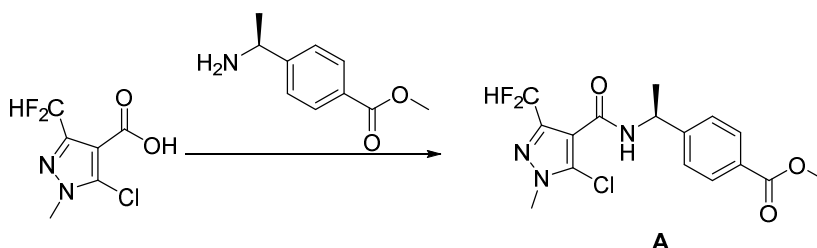
[0094] Preparation Example 1: Preparation of Intermediate A

[0095] Methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (Intermediate A)



[0096]

[0097] The synthesis scheme for **Intermediate A** is shown below:



[0098]

[0099] A starting material 5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxylic acid (5g, 23.8mmol), which was synthesized with reference to patent application WO2011151369A1, was added to DCM (200 mL). Methyl (S)-4-(1-aminoethyl) benzoate (5.1 g, 28.6 mmol), HATU (10.9 g, 28.6 mmol) and DIPEA (4.6 g, 35.7 mmol) were added. The mixture was stirred at room temperature for 16 hours; water (200 mL) was added, and the mixture was extracted with DCM (50 mL ×3) and separated, and the organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to

obtain a white solid of methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate.

[00100] LCMS (ESI) *m/z*: 372.5 [M+H]⁺

[00101] Preparative Example 2: Acidic Preparation Method A

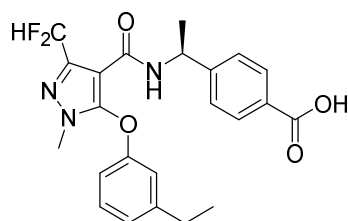
5 **[00102]** This example is an example for product purification, in which the purification is performed using high performance liquid chromatography with the following purification conditions: Welch, Ultimate C18 column, 10μm, 21.2 mm × 250 mm.

[00103] The mobile phase A was 1% trifluoroacetic acid in pure water, the mobile phase B was acetonitrile. Gradient Conditions: within 0 to 3 min, the mobile phase A was kept at 90%; after gradient elution from 3 min to 18 min, the mobile phase A was changed from 90% to 5%, and
10 the mobile phase A was kept at 5% from 18 min to 22 min).

[00104] The "Acidic Preparation Method A" described in the following Examples all refer to the Acidic Preparation Method A of the Preparation Example 2.

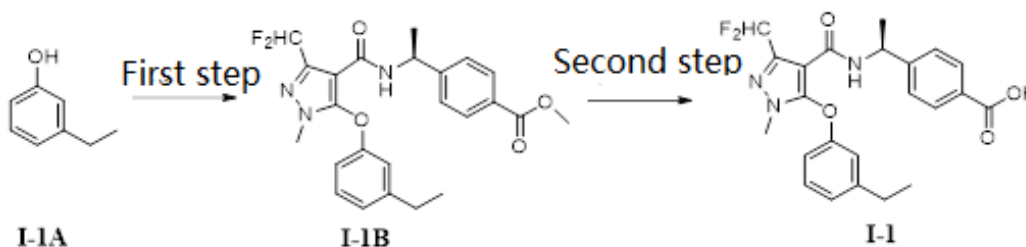
[00105] Example 1: Preparation of Compound I-1

15 **[00106]** (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-1**)



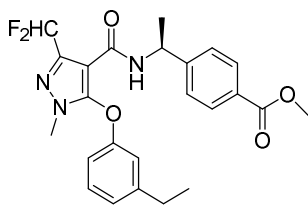
[00107]

[00108] The synthesis scheme for **Compound I-1** was shown below:



[00109]

20 **[00110]** First step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-1B**)

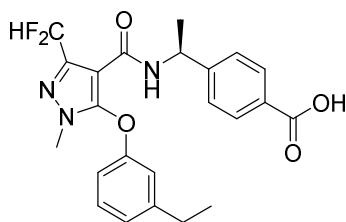


[00111] I-1B

[00112] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (370 mg, 1.0 mmol) was added to DMF (10 mL) at room temperature, and 3-ethylphenol (**I-1A**) (183 mg, 1.5 mmol) and KOH (168 mg, 3.0 mmol) were added; the mixture was heated to 120°C and stirred for 6h, and then was cooled to room temperature and diluted with water (40 mL); the pH was adjusted to 7 with 1N hydrochloric acid; the solution was extracted with ethyl acetate (20 mL ×3) and separated the organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-1B**) (220 mg, yield 48.1%).

[00113] LCMS (ESI) m/z: 458.1 [M+H]⁺

[00114] Second step: (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-1**)



[00115] I-1

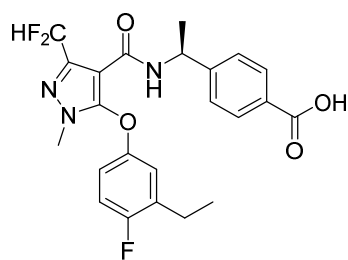
[00116] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-1B**) (220 mg, 0.48 mmol) was added to THF (4 mL) at room temperature; water (2 mL) and lithium hydroxide monohydrate (42 mg, 1.0 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(3-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-1**) (80 mg, yield 37.5%).

[00117] ^1H NMR (400m Hz, DMSO- d_6) δ 12.8 (s, 1H), 7.90 (d, 1H), 7.71 (d, 2H), 7.32 (t, 1H), 7.25 (t, 1H), 7.12 (d, 2H), 7.07 (d, 1H), 6.90 (s, 1H), 6.76 (dd, 1H), 4.90 (t, 1H), 3.72 (s, 3H), 2.61 (q, 2H), 1.22 (d, 3H), 1.14 (t, 3H).

[00118] LCMS (ESI) m/z : 444.1 $[\text{M}+\text{H}]^+$

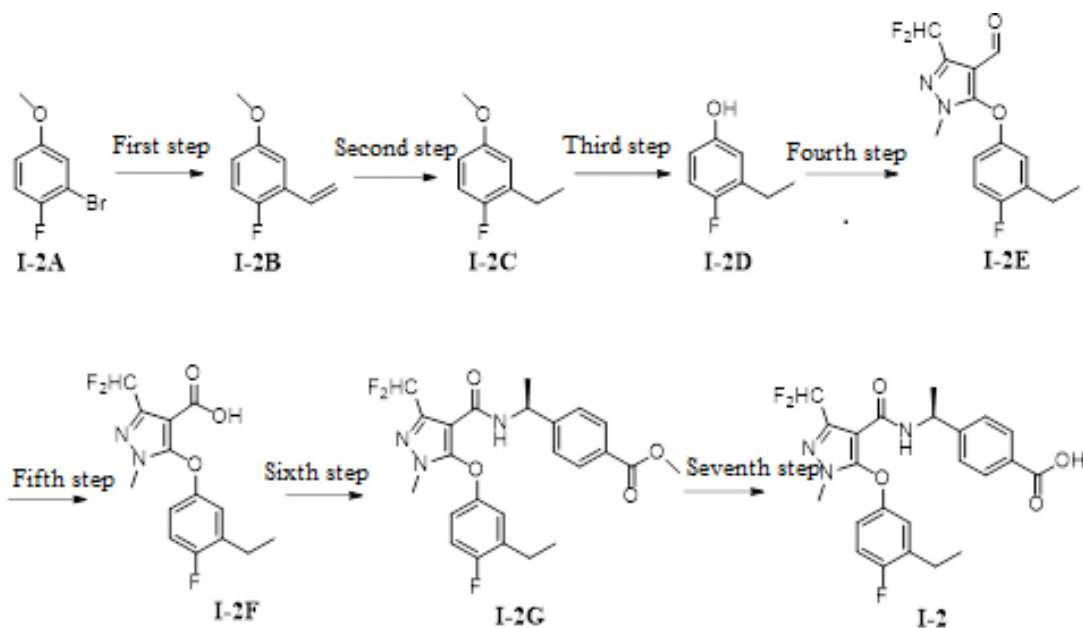
5 **[00119]** Example 2: Preparation of Compound I-2

(*S*)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-2**)



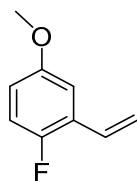
[00120]

[00121] The synthesis scheme for **Compound I-2** is shown below:



10 **[00122]**

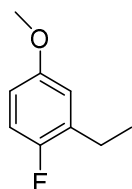
[00123] First step: 1-fluoro-4-methoxy-2-vinylbenzene (**Compound I-2B**)



[00124] **I-2B**

[00125] 2-bromo-1-fluoro-4-methoxybenzene (**Compound I-2A**) (1.02 g, 5.0 mmol) was added to 1, 4-dioxane (20 mL) at room temperature; potassium vinylfluoroborate (740 mg, 5.52 mmol) and [1, 1-bis(diphenylphosphino) ferrocene]dichloropalladium (430 mg, 0.50 mmol) and potassium carbonate (1.52 g, 11.0 mmol) were added. The mixture was heated to 100°C under nitrogen protection and stirred for 14 hours, and then cooled to room temperature, diluted with water (200 mL), extracted with DCM (80 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (pure petroleum ether) to obtain a colorless liquid crude product of 1-fluoro-4-methoxy-2-vinylbenzene (**Compound I-2B**) (680 mg, yield 89.9%).

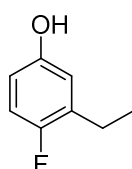
[00126] Second step: 2-ethyl-1-fluoro-4-methoxybenzene (**Compound I-2C**)



[00127] **I-2C**

[00128] 1-fluoro-4-methoxy-2-vinylbenzene (3.60 g, 23.7 mmol) was added to methanol (50 mL) at room temperature; 10% palladium on carbon (200 mg) was added; H₂ was purged, and then the solution was stirred for 16 hours at room temperature. The solution was filtrated and the filtrate washed with methanol (30 mL×3), and the organic phases were combined and concentrated to obtain a colorless liquid crude product of 2-ethyl-1-fluoro-4-methoxybenzene (**Compound I-2C**) (2.90 g, yield 79.5%).

[00129] Third step: 3-ethyl-4-fluorophenol (**Compound I-2D**)

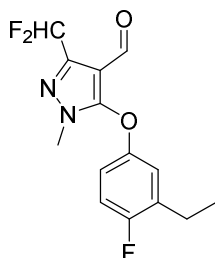


[00130] **I-2D**

[00131] 2-ethyl-1-fluoro-4-methoxybenzene (100 mg, 0.65 mmol) was added to DCM (3 mL) at room temperature, and then the mixture was cooled to -60°C. 1 mol/L BBr₃ DCM solution (2 mL) was added, the mixture was warmed up naturally to room temperature, and stirred at room temperature for 4 H. The residue was purified by silica gel column (petroleum ether: ethyl

acetate (V/V) = 4: 1) to obtain a colorless liquid of 3-ethyl-4-fluorophenol (**Compound I-2D**) (60 mg, yield 89.9%).

[00132] Fourth step: 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (**Compound I-2E**)

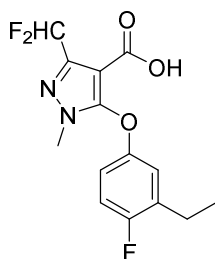


5 **[00133]** **I-2E**

[00134] The compound 5-chloro-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carbaldehyde (350 mg, 1.18 mmol) was added to DMF (5 mL) at room temperature; 3-ethyl-4-fluorophenol (379 mg, 2.70 mmol) and potassium carbonate (546 mg, 3.95 mmol) were added; and the mixture was heated to 100°C and stirred for 1.5 h. Then the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (15 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a colorless liquid crude product of 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (**Compound I-2E**) (600 mg, yield 100%).

[00135] LCMS (ESI) *m/z*: 299.1 [M+H]⁺

[00136] Fifth step: 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic acid (**Compound I-2F**)



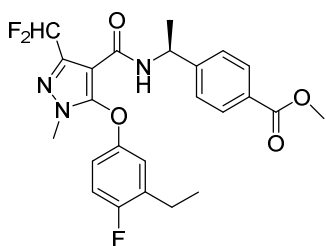
[00137] **I-2F**

20 **[00138]** The compound 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (350 mg, 1.80 mmol) was added to tert-butanol (10 mL) and water (2

mL) at room temperature; 2-methyl-2-butene (246 mg, 3.52 mmol), sodium chlorite (316 mg, 3.52 mmol) and sodium dihydrogen phosphate (281 mg, 2.34 mmol) were added. The mixture was stirred for 4 hours at room temperature. The solution was diluted with water (5 mL), extracted with ethyl acetate (10 mL× 3) and separated, and the organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic acid (**Compound I-2F**) (350 mg, yield 94.9%).

[00139] LCMS (ESI) *m/z*: 315.1 [M+H]⁺

[00140] Sixth step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-2G**)

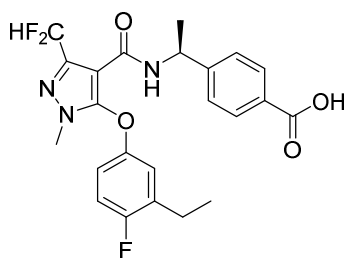


[00141] **I-2G**

[00142] The compound 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic acid (350 mg, 1.11 mmol) was added to DMF (5 mL) at room temperature, and methyl (S)-4-(1-aminoethyl) benzoate (220 mg, 1.23 mmol), HATU (467 mg, 1.23 mmol) and DIPEA (301 mg, 2.33 mmol) were added. The mixture was stirred at room temperature for 16 hours, diluted with water (20 mL), extracted with ethyl acetate (10 mL ×3) and separated, and the organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated, and purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a colorless liquid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-2G**) (400 mg, yield 75.5%).

[00143] LCMS (ESI) *m/z*: 476.2 [M+H]⁺

[00144] Seventh step: (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-2**)



I-2

[00145]

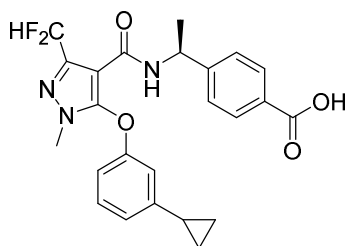
[00146] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (400 mg, 0.84 mmol) was added to THF (5 mL), water (5 mL) and methanol (5 mL) at room temperature, and lithium hydroxide monohydrate (141 mg, 3.36 mmol) was added. The mixture was stirred at room temperature for 16 hours, and the reaction mixture was concentrated to obtain a white solid of ((S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (300 mg, 77.2% yield).

[00147] ¹H NMR (400 m Hz, DMSO-d₆) 12.8 (s, 1H), 8.03 (d, 1H), 7.74 (d, 2H), 7.17 (t, 1H), 7.14-7.10 (m, 3H), 7.01-6.97 (m, 1H), 6.82-6.79 (m, 1H), 4.92 (t, 1H), 3.73 (s, 3H), 2.59 (q, 2H), 1.25 (d, 3H), 1.11 (t, 3H).

[00148] LCMS (ESI) m/z: 462.2 [M+H]⁺

[00149] Example 3: Preparation of Compound I-3

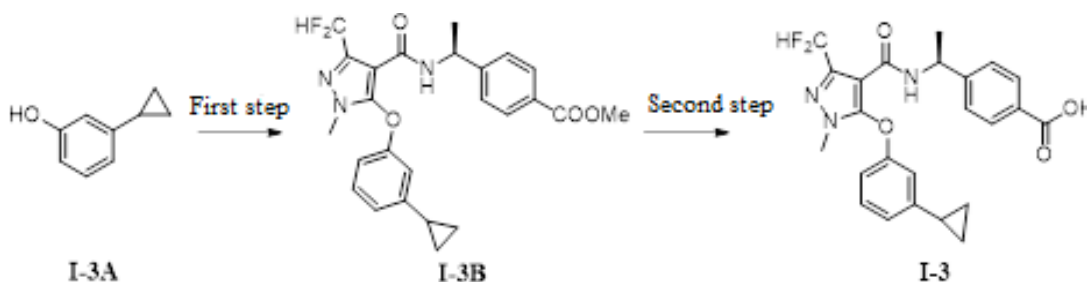
[00150] (S)-4-(1-(5-(3-cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-3**)



I-3

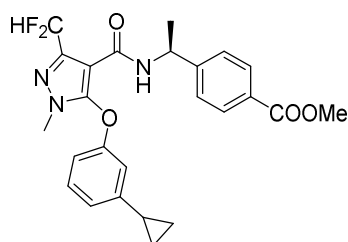
[00151]

[00152] The synthesis scheme for **Compound I-3** is shown below:



[00153]

[00154] First step: methyl (S)-4-(1-(5-(3-cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-3B**)

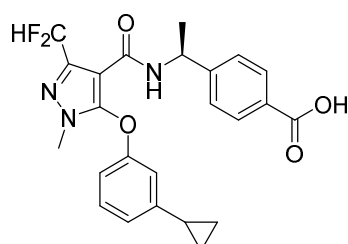


[00155] **I-3B**

5 **[00156]** The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (145 mg, 0.39 mmol) was added to DMA (2 mL) at room temperature, 3-cyclopropylphenol (80 mg, 0.59 mmol) and KOH (34 mg, 0.61 mmol) was added, and the mixture was heated to 120°C, stirred for 2 hours, and then cooled to room temperature, diluted with water (100 mL), extracted with ethyl acetate (10 mL ×3) and
10 separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of methyl (S)-4-(1-(5-(3-cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-3B**) (20 mg, yield 10.9%).

[00157] LCMS (ESI) m/z : 470.6 $[M+H]^+$

15 **[00158]** Second step: (S)-4-(1-(5-(3-cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-3**)



[00159] **I-3**

[00160] A starting material methyl (S)-4-(1-(5-(3-cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-3B**) (20 mg, 0.04 mmol) was added to THF (1 mL) at room temperature, and water (1 mL) and lithium hydroxide monohydrate (2 mg, 0.048 mmol) were added. The mixture was stirred at room temperature for
20 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(3-

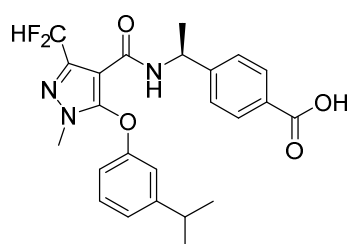
cyclopropylphenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-3**) (1.5 mg, 7.7% yield).

[00161] ¹H NMR (400m Hz, DMSO-*d*₆) δ 12.8 (s, 1H), 7.92 (d, 1H), 7.72 (d, 2H), 7.27 (t, 1H), 7.11 (t, 1H), 7.10 (d, 2H), 6.90 (d, 1H), 6.80 (t, 1H), 6.70 (dd, 1H), 4.90 (t, 1H), 3.72 (s, 3H), 1.92-1.98 (m, 1H), 1.23 (d, 3H), 0.96-0.92 (m, 2H), 0.66-0.27 (m, 2H).

[00162] LCMS (ESI) *m/z*: 456.6 [M+H]⁺

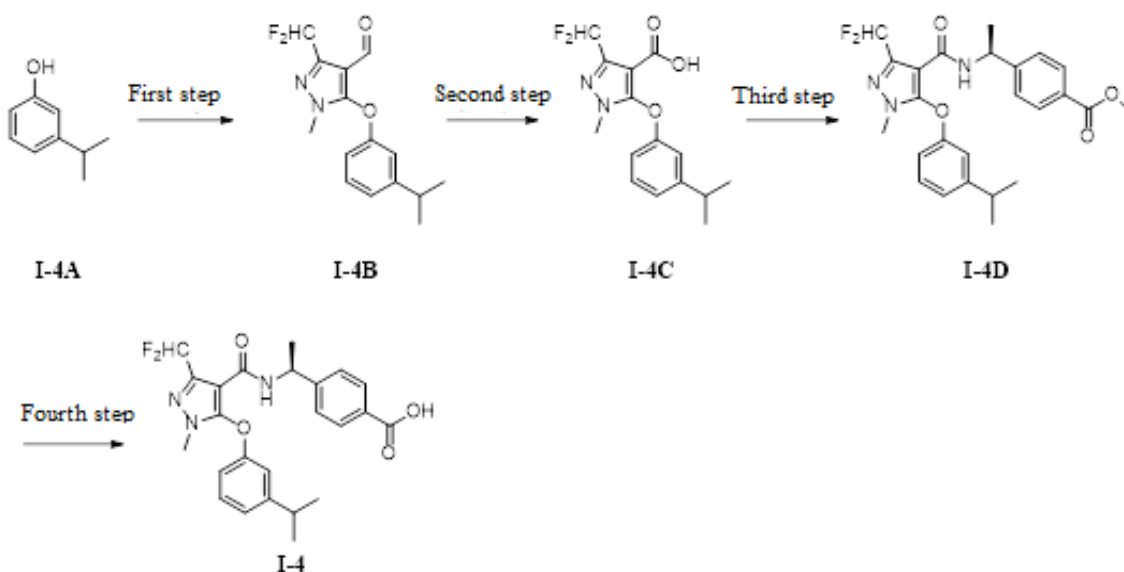
[00163] Example 4: Preparation of Compound I-4

[00164] (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-4**)



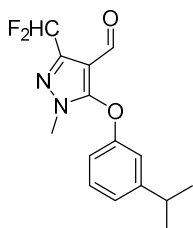
I-4

[00166] The synthesis scheme for **Compound I-4** is shown below:



[00167]

[00168] First step: 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (**Compound I-4B**)

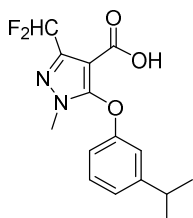


[00169] I-4B

[00170] The compound 5-chloro-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carbaldehyde (500 mg, 2.57 mmol) was added to DMF (5 mL) at room temperature; 3-isopropylphenol (386 mg, 2.80 mmol) and KOH (216 mg, 3.85 mmol) were added; and the mixture was heated to 150°C and stirred for 4 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (15 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow liquid crude product of 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (**Compound I-4B**) (750 mg, 98.9% yield).

[00171] LCMS (ESI) *m/z*: 295.1 [M+H]⁺

[00172] Second step: 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic acid (**Compound I-4C**)



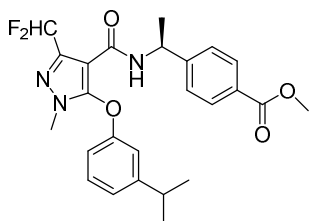
[00173] I-4C

[00174] The compound 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carbaldehyde (750 mg, 2.55 mmol) was added to tert-butanol (6 mL) and water (7 mL) at room temperature, 2-methyl-2-butene (355 mg, 5.07 mmol), sodium chlorite (456 mg, 5.07 mmol), and sodium dihydrogen phosphate (669 mg, 5.57 mmol) were added. The mixture was stirred for 14 hours at room temperature. The mixture was diluted with water (15 mL) and extracted with ethyl acetate (30 mL ×3) and separated, and the organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a light yellow solid crude product of 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic

acid (**Compound I-4C**) (800 mg, yield 100%).

[00175] LCMS (ESI) m/z : 311.1 $[M+H]^+$

[00176] Third step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-4D**)

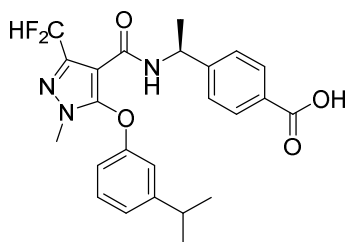


5 [00177] **I-4D**

[00178] The compound 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxylic acid (800 mg, 2.58 mmol) was added to DCM (20 mL); (S)-methyl 4-(1-aminoethyl)benzoate (459 mg, 2.56 mmol), HATU (1.40 g, 3.68 mmol) and DIPEA (991 mg, 7.68 mmol) were added; and the mixture was stirred at room temperature for 16 hours, diluted with DCM (40 mL), washed with water (20 mL $\times$ 3) and separated. The organic phases were dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow liquid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-4D**) (720 mg, yield 59.2%).

15 [00179] LCMS (ESI) m/z : 472.2 $[M+H]^+$

[00180] Fourth step: (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-4**)



[00181] **I-4**

[00182] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-4D**) (440 mg, 0.93 mmol) was added to methanol (10 mL) and water (1 mL) at room temperature, and sodium hydroxide (93 mg, 2.32 mmol) was added. The mixture was stirred at room temperature for 16 hours. The

20

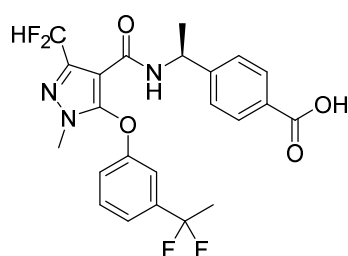
reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (93 mg, 21.7% yield).

[00183] ¹H NMR (400m Hz, DMSO-d₆) δ 12.7 (s, 1H), 7.89 (d, 1H), 7.70 (d, 2H), 7.32 (t, 1H), 7.12 (t, 1H), 7.10 (s, 1H), 7.06 (d, 2H), 6.99 (s, 1H), 6.72-6.70 (m, 1H), 4.91 (t, 1H), 3.73 (s, 3H), 2.91-2.85 (m, 1H), 1.21 (d, 3H), 1.17 (d, 6H).

[00184] LCMS (ESI) m/z: 458.3 [M+H]⁺

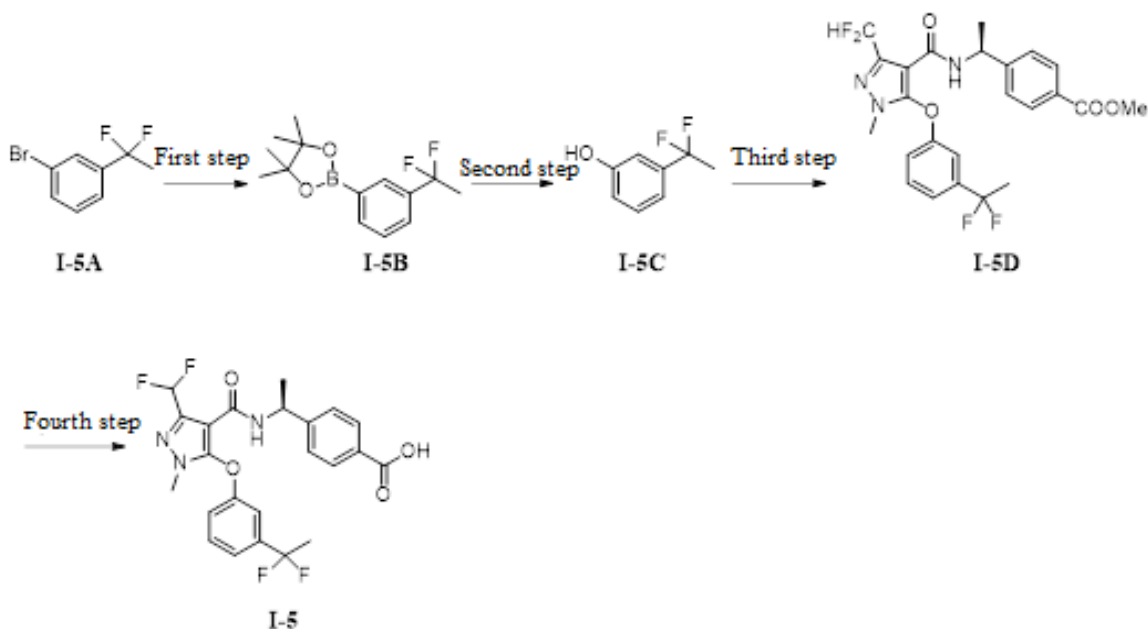
[00185] Example 5: Preparation of Compound I-5

[00186] (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-5**)



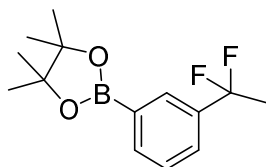
[00187] **I-5**

[00188] The synthesis scheme for **Compound I-5** is shown below:



[00189]

[00190] First step: 2-(3-(1,1-difluoroethyl)phenyl)-4,4,5,5-tetramethyl-1,3,2-dioxaborolane (**Compound I-5B**)

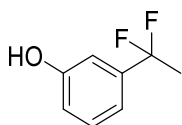


I-5B

[00191]

[00192] 1-bromo-3-(1, 1-difluoroethyl) benzene (800 mg, 3.62 mmol) was added to 1, 4-dioxane (30 mL) at room temperature, and bis(pinacolato)diboron (17.0 g, 156.3 mmol), copper iodide (2.5 g, 13.0 mmol), L-proline (2.76 g, 10.86 mmol), potassium acetate (710 mg, 7.24 mmol) and [1, 1-bis(diphenylphosphino) ferrocene] dichloropalladium (295 mg, 0.36 mmol) were added. The mixture was heated to 90°C under nitrogen protection and stirred for 16 hours, and then the mixture was cooled to room temperature, diluted with water (200 mL), extracted with dichloromethane (80 mL ×3) and separated. The organic phases were combined and dried with anhydrous sodium sulfate, filtered and concentrated, and the residue was purified the by silica gel column separation (pure petroleum ether) to obtain a colorless liquid crude product of 2-(3-(1,1-difluoroethyl)phenyl)-4,4,5,5-tetramethyl-1,3,2-dioxaborolane (**Compound I-5B**) (900 mg, yield 92.7%).

[00193] Second step: 3-(1,1-difluoroethyl)phenol (**Compound I-5C**)

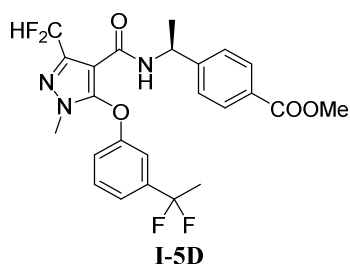


I-5C

[00194]

[00195] 2-(3-(1,1-difluoroethyl)phenyl)-4,4,5,5-tetramethyl-1,3,2-dioxaborolane (**Compound I-5B**) (900 mg, 3.36 mmol) was added to THF (15 mL) and water (15 mL) at room temperature, and sodium perborate monohydrate (1.01 g, 10.07 mmol) was added. The mixture was stirred for 16 hours at room temperature, and the mixture was diluted with water (200 mL), extracted with DCM (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 8: 1) to obtain a colorless liquid of 3-(1,1-difluoroethyl)phenol (**Compound I-5C**) (280 mg, yield 52.7%).

[00196] Third step: methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-5D**)

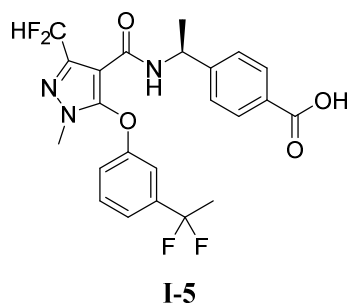


[00197]

[00198] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (650 mg, 1.75 mmol) was added to DMF (12 mL) at room temperature; 3-(1, 1-difluoroethyl) phenol (360 mg, 2.27 mmol) and potassium hydroxide (147 mg, 2.62 mmol) were added; and the mixture was heated to 120°C and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (200 mL), extracted with ethyl acetate (80 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, and concentrated to obtain a colorless liquid crude product of methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-5D**) (1.2 g, crude product).

[00199] LCMS (ESI) *m/z*: 494.6 [M+H]⁺.

[00200] Fourth step: (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-5**)



[00201]

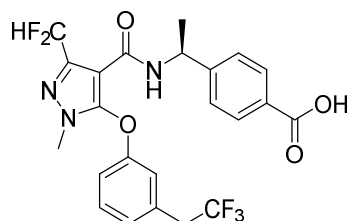
[00202] A starting material methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-5D**) (1.0 g, 2.03 mmol) was added to THF (5 mL) at room temperature, and water (4 mL) and lithium hydroxide monohydrate (340 mg, 8.11 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-5**) (88 mg, 7.9% yield).

[00203] LCMS (ESI) *m/z*: 480.5 [M+H]⁺

[00204] ^1H NMR (400m Hz, DMSO- d_6) δ 12.8 (s, 1H), 8.10 (d, 1H), 7.71 (d, 2H), 7.53 (t, 1H), 7.47 (d, 1H), 7.27 (d, 1H), 7.11 (t, 1H), 7.11 (d, 2H), 7.07 (dd, 1H), 4.88 (t, 1H), 3.74 (s, 3H), 1.96 (t, 3H), 1.96 (d, 3H).

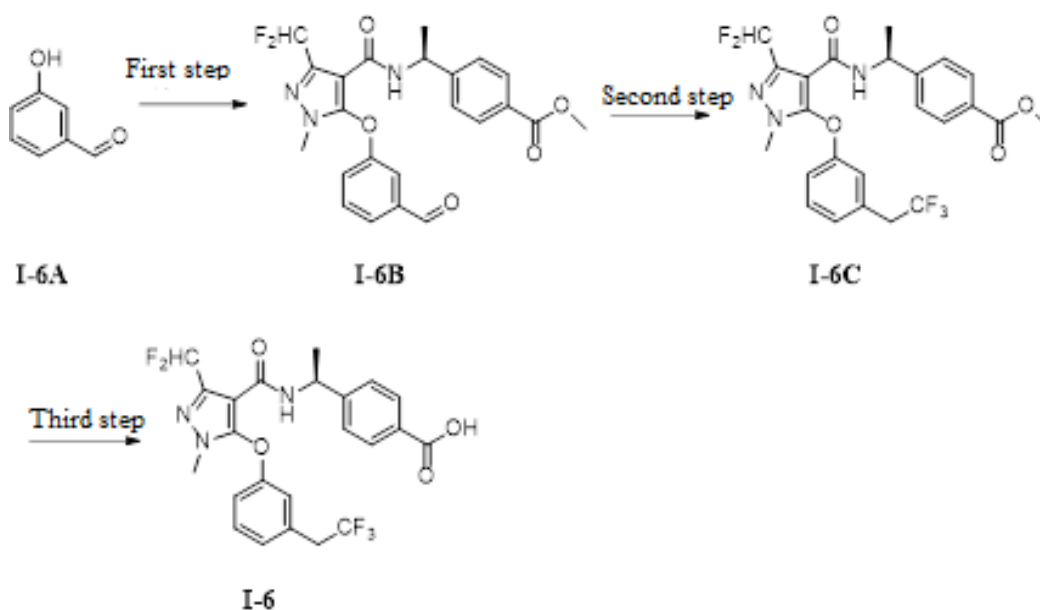
[00205] Example 6: Preparation of Compound I-6

- 5 **[00206]** (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-6**)



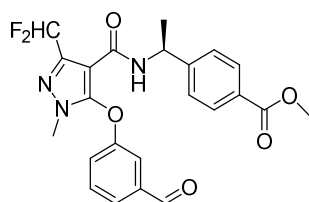
[00207] **I-6**

[00208] The synthesis scheme for **Compound I-6** is shown below:



[00209] **I-6**

- 10 **[00210]** First step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6B**)



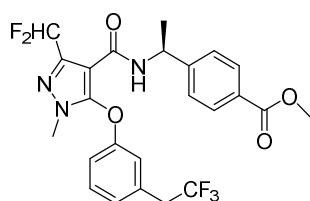
[00211] **I-6B**

[00212] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-

4-carboxamido)ethyl)benzoat (**Intermediate A**) (371 mg, 1.0 mmol) was added to DMSO (5 mL) at room temperature, and 3-hydroxybenzaldehyde (122 mg, 1.0 mmol), K₂CO₃ (270 mg, 2.0 mmol) and cuprous iodide (76 mg, 0.4 mmol), phenanthroline (72 mg, 0.4 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (20 ml), extracted with ethyl acetate (10 ml × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 8: 1) to obtain a colorless liquid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6B**) (180 mg, yield 39.3%).

[00213] LCMS (ESI) m/z: 458.1 [M+H]⁺

[00214] Second step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6C**)



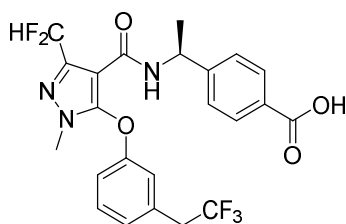
[00215] **I-6C**

[00216] Methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6B**) (41.4 mg, 0.09 mmol) was added to DMF (2 mL) and 2,2-difluoro-2-triphenylphosphoniumylacetate (64 mg, 0.18 mmol) was added at room temperature, and the mixture was heated to 60°C, and stirred for 2 hours. A solution of tetrabutylammonium fluoride in tetrahydrofuran (0.3 mL, 0.30 mmol) was added and stirring was continued for 4 hours. Then, the mixture was cooled to room temperature, the reaction mixture was concentrated and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 8: 1) to obtain a colorless solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6C**) (40 mg, 91.2% yield).

[00217] LCMS (ESI) m/z: 512.1 [M+H]⁺

[00218] Third step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-

trifluoroethyl)phenoxy)-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-6**)



[00219] **I-6**

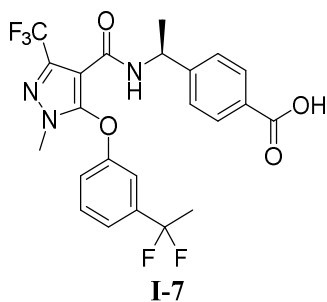
[00220] A starting material (*S*)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethyl)phenoxy)-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-6C**) (40 mg, 0.082 mmol) was added to methanol (2 mL) at room temperature, and water (2 mL) and lithium hydroxide monohydrate (12 mg, 0.3 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (*S*)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethyl)phenoxy)-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-6**) (4.8 mg, 12.3% yield).

[00221] ¹H NMR (400m Hz, DMSO-*d*₆) δ 12.7 (s, 1H), 7.95 (d, 1H), 7.73 (d, 2H), 7.43 (t, 1H), 7.22 (d, 1H), 7.13 (t, 1H), 7.12 (d, 2H), 7.10 (s, 1H), 6.97 (dd, 1H), 4.89 (t, 1H), 3.74 (s, 3H), 3.72-3.63 (m, 2H), 1.24 (d, 3H).

[00222] LCMS (ESI) *m/z*: 498.5 [M+H]⁺

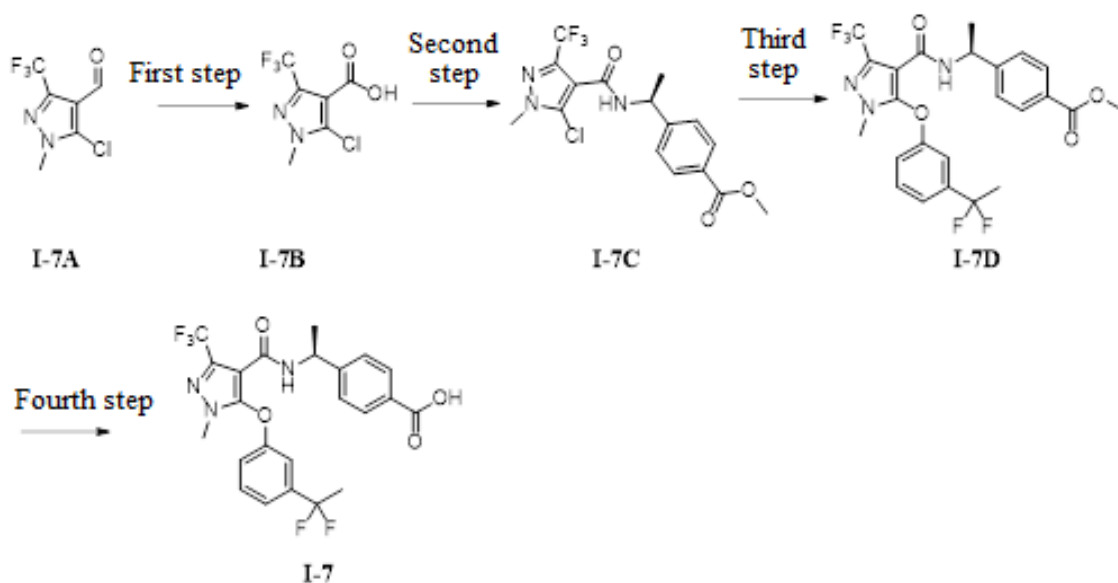
[00223] Example 7: Preparation of Compound I-7

[00224] (*S*)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-7**)



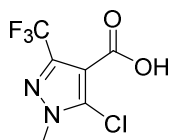
[00225]

[00226] The synthesis scheme for **Compound I-7** is shown below:



[00227]

[00228] First step: 5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxylic acid (**Compound I-7B**)



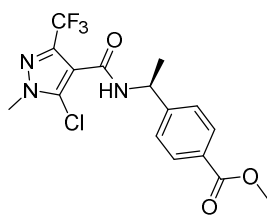
I-7B

[00229]

[00230] A compound 5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carbaldehyde (700 mg, 3.30 mmol) was added to tert-butanol (20 mL) and water (5 mL) at room temperature, and 2-methyl-2-butene (1.80 g, 25.7 mmol), sodium chlorite (1.48 g, 16.4 mmol) and sodium dihydrogen phosphate (3.10 g, 25.8 mmol) were added. The mixture was stirred for 14 hours at room temperature, and then the mixture was diluted with water (50 mL) and extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain the crude residue. The crude residue was then purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid of 5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxylic acid (**Compound I-7B**) (680 mg, yield 90.3%).

[00231] LCMS (ESI) m/z : 229.6 $[M+H]^+$

[00232] Second step: methyl (*S*)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7C**)

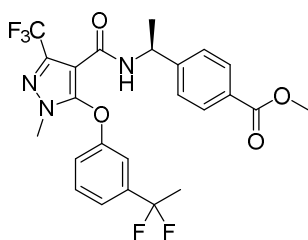


[00233] I-7C

[00234] The compound 5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxylic acid (**Compound I-7B**) (680 mg, 2.98 mmol) was added to DMF (20 mL), and methyl (S)-4-(1-aminoethyl) benzoate (537 mg, 3.00 mmol), HATU (1.70 g, 4.47 mmol) and DIPEA (1.90 g, 14.7 mmol) were added. The mixture was stirred at room temperature for 16 hours. Then, the mixture was diluted with water (200 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filter and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid of methyl (S)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7C**) (700 mg, 60.3% yield).

[00235] LCMS (ESI) *m/z*: 390.5 [M+H]⁺

[00236] Third step: methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7D**)



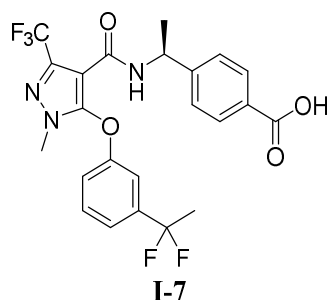
[00237] I-7D

[00238] The compound methyl (S)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7C**) (700 mg, 1.00 mmol) was added to DMF (10 mL) at room temperature, and 3-(1, 1-difluoroethyl) phenol (284 mg, 1.00 mmol) and KOH (264 mg, 4.63 mmol) were added. The mixture was heated to 120°C and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (20 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered, and concentrated to obtain a colorless liquid crude product

of methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7D**) (100 mg, yield 10.8%).

[00239] LCMS (ESI) m/z: 512.3 [M+H]⁺

[00240] Fourth step: (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-7**)



[00241]

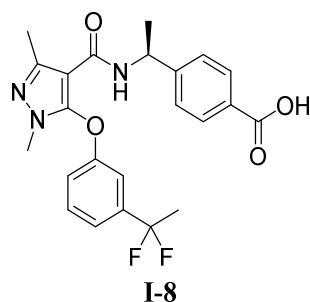
[00242] A starting material methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-7D**) (100 mg, 0.19 mmol) was added to THF (5 mL) at room temperature, and water (4 mL) and lithium hydroxide monohydrate (10 mg, 0.24 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-7**) (47.2 mg, 48.5% yield).

[00243] ¹H NMR (400 MHz, DMSO-d₆) δ 12.8 (s, 1H), 8.58 (d, 1H), 7.73 (d, 2H), 7.53 (t, 1H), 7.40 (d, 1H), 7.27 (s, 1H), 7.15 (d, 2H), 7.11 (d, 1H), 4.85 (t, 1H), 3.29 (s, 3H), 1.96 (t, 3H), 1.17 (d, 3H).

[00244] LCMS (ESI) m/z: 498.3 [M+H]⁺

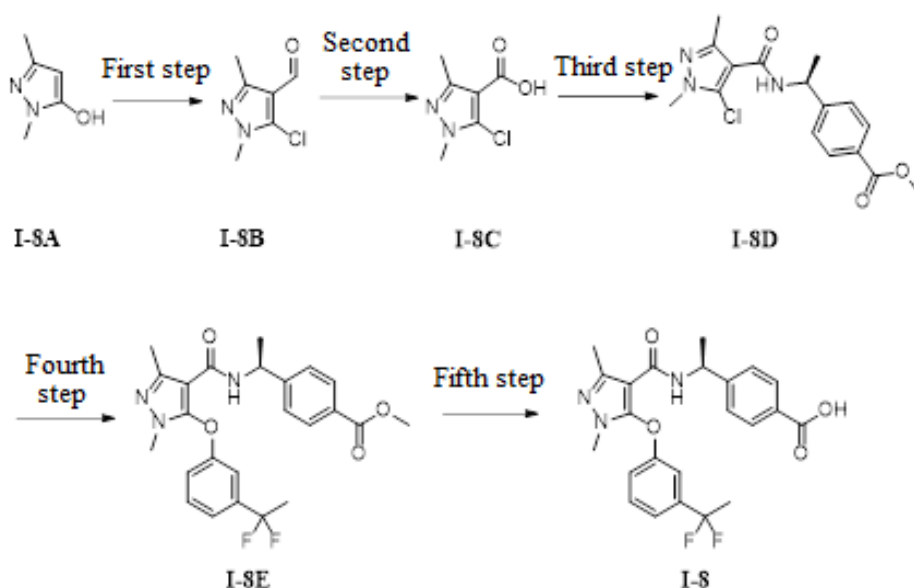
[00245] Example 8: Preparation of Compound I-8

[00246] (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-8**)



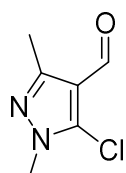
[00247]

[00248] The synthesis scheme for **Compound I-8** is shown below:



[00249]

[00250] First step: 5-chloro-1,3-dimethyl-1H-pyrazole-4-carbaldehyde (**Compound I-8B**)

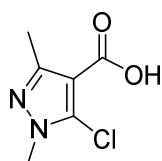


[00251] **I-8B**

- 5 **[00252]** 1, 3-dimethyl-5-hydroxypyrazole (5.5 g, 49.1 mmol) was added to DMF (10.9 g) at room temperature, and the mixture was cooled to 0°C, and POCl₃ (53.0 g, 346.4 mmol) was added and then naturally warmed to room temperature, heated to 120°C and stirred for 1 h. Then, the mixture was cooled to room temperature, diluted with water (200 mL), extracted with EA (200 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless liquid of 5-chloro-1,3-dimethyl-1H-pyrazole-4-carbaldehyde (**Compound I-8B**) (4.3 g, yield 55.4%).
- 10

[00253] LCMS (ESI) m/z: 159.6 [M+H]⁺

[00254] Second step: 5-chloro-1,3-dimethyl-1H-pyrazole-4-carboxylic acid (**Compound I-8C**)



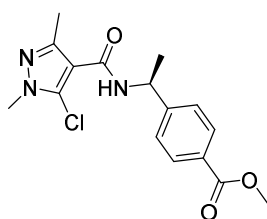
[00255] **I-8C**

15

[00256] The compound 5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carbaldehyde (**Compound I-8B**) (3.20 g, 20.2 mmol) was added to tert-butanol (50 mL) and water (15 mL) at room temperature; and 2-methyl-2-butene (11.3 g, 161.4 mmol), sodium chlorite (9.10 g, 101.1 mmol) and sodium dihydrogen phosphate (19.4 g, 161.6 mmol) were added. The mixture was stirred for 14 hours at room temperature; and then the mixture was diluted with water (50 mL), extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain the crude colorless liquid residue of the captioned compound. The residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid of 5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carboxylic acid (**Compound I-8C**) (3.10 g, yield 87.9%).

[00257] LCMS (ESI) *m/z*: 175.6 [M+H]⁺

[00258] Third step: methyl (S)-4-(1-(5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8D**)

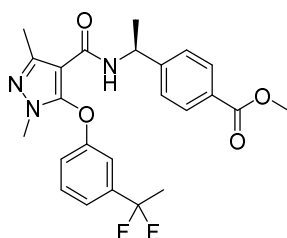


[00259] **I-8D**

[00260] The compound 5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carboxylic acid (**Compound I-8C**) (1.20 g, 6.89 mmol) was added to DMF (30 mL); and (S)-methyl 4-(1-aminoethyl) benzoate (1.23 g, 6.87 mmol), HATU (3.90 g, 10.1 mmol) and DIPEA (4.50 g, 34.8 mmol) were added. The mixture was stirred at room temperature for 16 hours, diluted with water (200 mL), extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid of methyl (S)-4-(1-(5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8D**) (1.90 g, yield 82.2%).

[00261] LCMS (ESI) *m/z*: 336.6 [M+H]⁺

[00262] Fourth step: methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8E**)

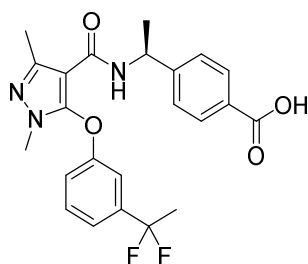


[00263] **I-8E**

[00264] The compound methyl (S)-4-(1-(5-chloro-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8D**) (335 mg, 1.00 mmol) was added to DMF (10 mL) at room temperature, and 3-(1, 1-difluoroethyl) phenol (158 mg, 1.00 mmol) and KOH (150 mg, 2.67 mmol) were added. The mixture was heated to 120°C and stirred for 16 hours. Then the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (20 mL ×3) and separated. The organic phases were combined and dried with anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of ethyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8E**) (170 mg, yield 37.1%).

[00265] LCMS (ESI) *m/z*: 458.5 [M+H]⁺

[00266] Fifth step: (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-8**)



[00267] **I-8**

[00268] A starting material ethyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8E**) (500 mg, 1.09 mmol) was added to THF (5 mL) at room temperature, and water (4 mL) and lithium hydroxide monohydrate (340 mg, 8.11 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(3-(1,1-difluoroethyl)phenoxy)-1,3-dimethyl-1*H*-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-8**) (53 mg, 10.9% yield).

[00269] ¹H NMR (400m Hz, DMSO-*d*₆) δ 12.8 (s, 1H), 7.73 (d, 2H), 7.71 (d, 1H), 7.52 (t, 1H),

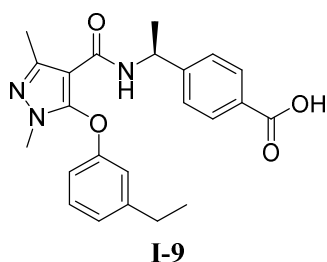
7.37 (d, 1H), 7.21 (s, 1H), 7.14 (d, 2H), 7.01 (dd, 1H), 4.91 (t, 1H), 3.59 (s, 3H), 2.27 (s, 3H), 1.96 (t, 3H), 1.22 (d, 3H).

[00270] LCMS (ESI) m/z : 444.5 $[M+H]^+$

[00271] Example 9: Preparation of Compound I-9

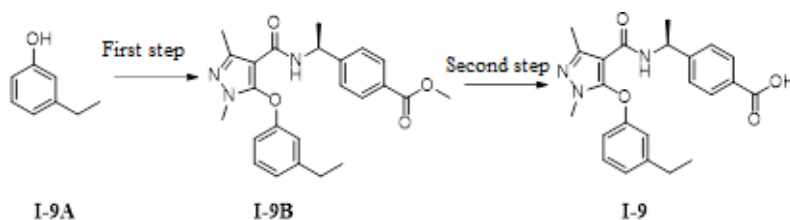
5 [00272] (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-9**)

[00273] The synthesis scheme for Compound I-9 is shown below:



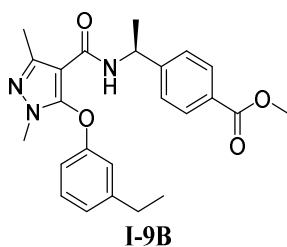
[00274]

[00275] The synthesis scheme for **Compound I-9** is shown below:



10 [00276]

[00277] First step: methyl (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-9B**)



[00278]

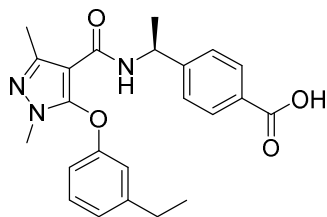
[00279] The compound methyl (S)-4-(1-(5-chloro-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-8D**) (100 mg, 0.30 mmol) was added to DMF (3

15 mL) at room temperature; 3-ethylphenol (36 mg, 0.30 mmol) and KOH (49 mg, 0.86 mmol) were added; and the mixture was heated to 120°C and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (20 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium

sulfate, filtered and concentrated to obtain a colorless liquid crude product of methyl (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-9B**) (60 mg, 47.7% yield).

[00280] LCMS (ESI) m/z: 422.6 [M+H]⁺

5 [00281] Second step: (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-9**)



[00282] I-9

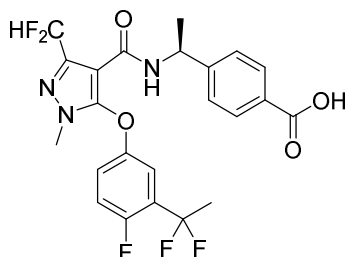
[00283] A starting material methyl (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-9B**) (60 mg, 0.14 mmol) was added to THF (2 mL) at room temperature, and water (2 mL) and lithium hydroxide monohydrate (6 mg, 0.14 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(3-ethylphenoxy)-1,3-dimethyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-9**) (10 mg, 17.2% yield).

[00284] ¹H NMR (400m Hz, DMSO-d₆) δ 12.6 (s, 1H), 7.72 (d, 2H), 7.50 (d, 1H), 7.31 (t, 1H), 7.12 (d, 2H), 7.04 (d, 1H), 6.85 (s, 1H), 6.71 (dd, 1H), 4.92 (t, 1H), 3.56 (s, 3H), 2.67 (q, 2H), 2.27 (s, 3H), 1.24 (d, 3H), 1.14 (t, 3H).

[00285] LCMS (ESI) m/z: 408.6 [M+H]⁺

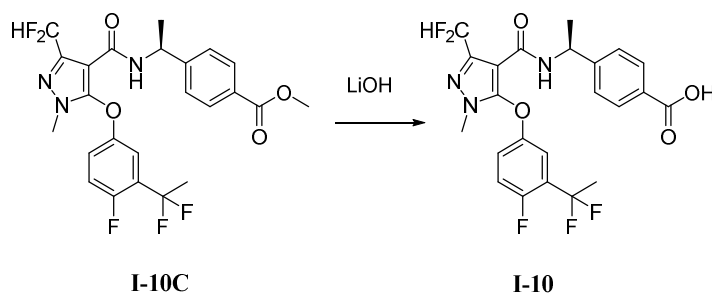
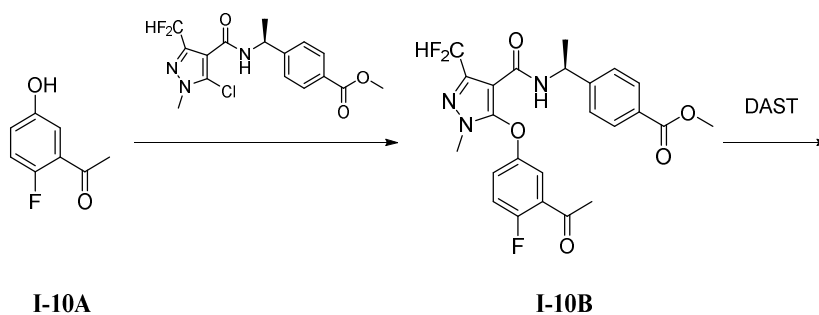
[00286] Example 10: Preparation of Compound I-10

[00287] (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-10**)



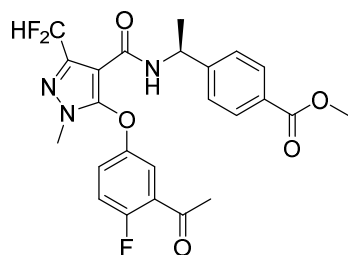
[00288] I-10

[00289] The synthesis scheme for **Compound I-10** is shown below:



[00290]

[00291] First step: methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-10B**)



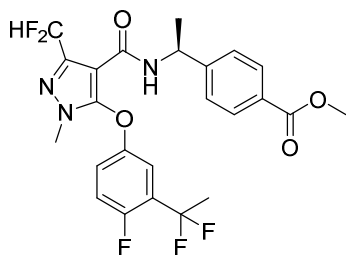
[00292]

I-10B

5 [00293] The compound 1-(2-fluoro-5-hydroxyphenyl) ethan-1-one (1.58 g, 10.8 mmol) was added to N, N-dimethylformamide (30 mL) at room temperature; methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (2.0 g, 5.4 mmol) and potassium hydroxide (450 mg, 8.1 mmol) were added; and the mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours. The mixture was cooled to room temperature and added with water (30 mL), extracted with ethyl acetate (30 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a white solid of methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl) benzoate (**Compound I-10B**) (346 mg, 13% yield).

[00294] LC-MS, M/Z (ESI): 490.2 [M+H]⁺

[00295] Second step: methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-10C**)

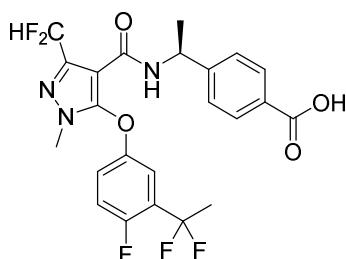


[00296] **I-10C**

[00297] The compound methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-10B**) (346 mg, 0.71 mmol) was added to (diethylamino) sulfur trifluoride (10 mL) at room temperature. The mixture was heated to 50°C and stirred for 16 hours. Then, the mixture was cooled to 0°C, diluted with water (50 mL), extracted with EA (100 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white crude product of methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-10C**) (390 mg, yield 100%).

[00298] LC-MS, M/Z (ESI): 512.2 [M+H]⁺

[00299] Third step 3: (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-10**)



[00300] **I-10**

[00301] The compound methyl (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-10C**) (390 mg, 0.76 mmol) was added to tetrahydrofuran (10 mL), methanol (10 mL) and water (3 mL) at room temperature, and lithium hydroxide (64 mg, 1.52 mmol) was added. The mixture was stirred at room temperature for 36 hours, and then the pH was adjusted to 7 with 1N hydrochloric acid. The mixture was concentrated to obtain, by the acidic preparation method, a

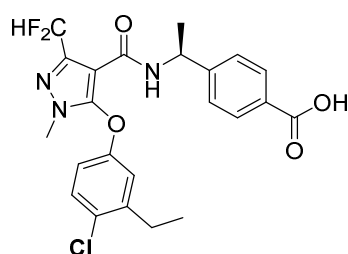
white solid of (S)-4-(1-(5-(3-(1,1-difluoroethyl)-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-10**) (116 mg, 31% yield).

[00302] LC-MS, M/Z (ESI):498.1 [M+H]⁺

5 **[00303]** ¹H NMR (400mHz, DMSO-d₆) δ12.10 (s, 1H), 8.19 (d, 1H), 7.75 (d, 2H), 7.39 (t, 1H), 7.26 (t, 1H), 7.15 (d, 2H), 7.13 (d, 1H), 7.12 (t, 1H), 4.90 (t, 1H), 3.75 (s, 3H), 2.01 (t, 3H), 1.22 (d, 3H).

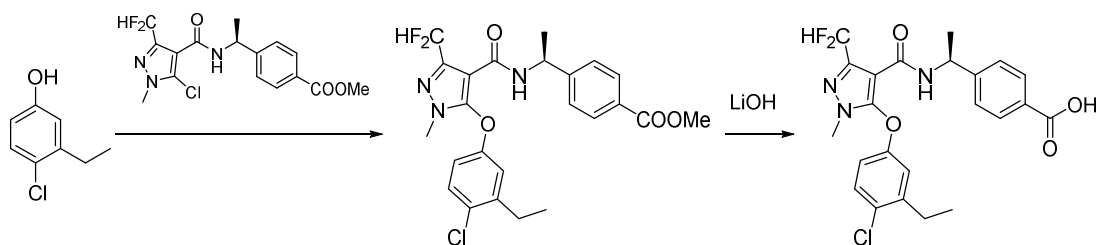
[00304] Example 11: Preparation of Compound I-11

10 **[00305]** (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-11**)



[00306] **I-11**

[00307] The synthesis scheme for **Compound I-11** is shown below:

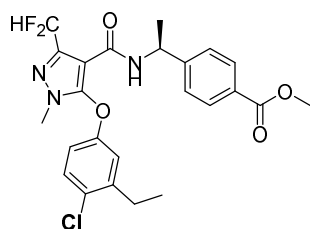


[00308] **I-11A**

I-11B

I-11

15 **[00309]** First step: methyl (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-11B**)



[00310] **I-11B**

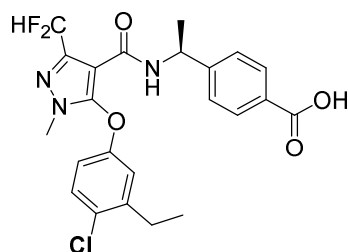
[00311] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (371 mg, 1.00 mmol) was added to dimethyl sulfoxide (3 mL) at room temperature; 4-chloro-3-ethylphenol (156 mg, 1.00 mmol) and

potassium hydroxide (112 mg, 2.00 mmol) were added; and the mixture was heated to 120°C and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (10 mL), extracted with ethyl acetate (10 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of methyl (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-11B**) (80 mg, yield 16%).

[00312] LC-MS, M/Z (ESI): 492.3 (M+1).

[00313] Second step: (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-11**)

[00314]



[00315] **I-11**

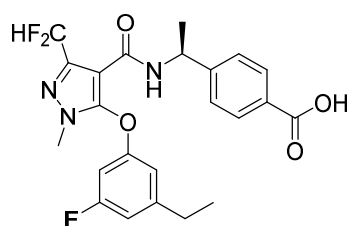
[00316] A starting material methyl (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-11B**) (80 mg, 0.16 mmol) was added to tetrahydrofuran (1 mL) at room temperature; and water (1 mL) and lithium hydroxide monohydrate (21 mg, 0.48 mmol) were added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of (S)-4-(1-(5-(4-chloro-3-ethylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-11**) (13.3 mg, 17% yield).

[00317] ¹H NMR (400m Hz, DMSO-d₆) δ12.8 (s, 1H), 8.04 (d, 1H), 7.73 (d, 2H), 7.41 (d, 1H), 7.11 (t, 1H), 7.10 (d, 2H), 7.08 (d, 1H), 6.82 (d, 1H), 4.91 (t, 1H), 3.73 (s, 3H), 2.67 (q, 2H), 1.24 (d, 3H), 1.11 (t, 3H).

[00318] LC-MS, M/Z (ESI): 478.3 (M+1)

[00319] Example 12: Preparation of Compound I-12

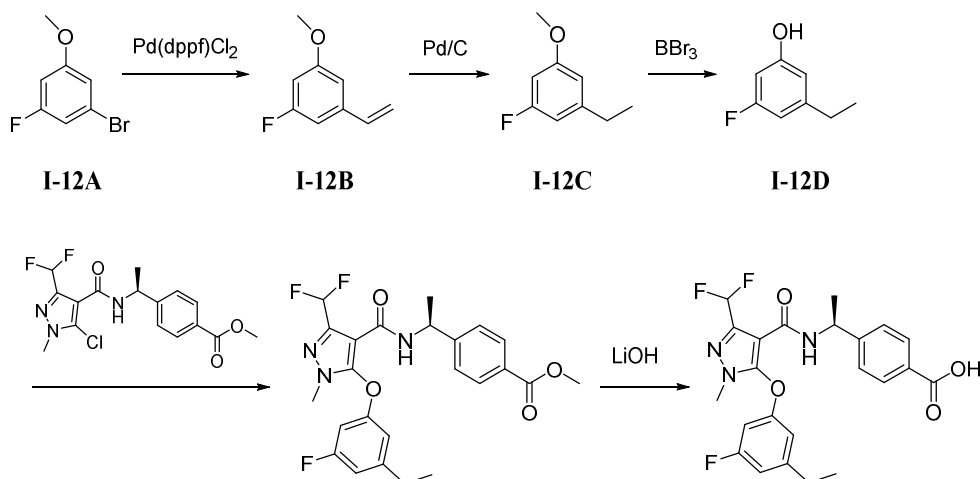
[00320] (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-12**)



[00321]

I-12

[00322] The synthesis scheme for **Compound I-12** is shown below:

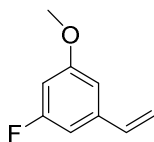


[00323]

I-12E

I-12

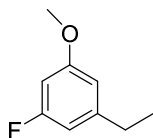
[00324] First step: 1-fluoro-3-methoxy-5-vinylbenzene (**Compound I-12B**)



5 [00325] **I-12B**

[00326] 1-bromo-3-fluoro-5-methoxybenzene (2.0 g, 9.7 mmol) was added to 1, 4-dioxane (50 mL) at room temperature; potassium vinylfluoroborate (1.6 g, 11.7 mmol); [1, 1-bis(diphenylphosphino) ferrocene] dichloropalladium (1.4 g, 1.96 mmol) and potassium carbonate (2.7 g, 19.6 mmol) were added; and the mixture was heated to 100°C under nitrogen protection and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (200 mL), extracted with dichloromethane (80 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (pure petroleum ether) to obtain 1-fluoro-3-methoxy-5-vinylbenzene (**Compound I-12B**) (600 mg, yield 40%).

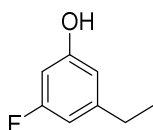
15 [00327] Second step: 1-ethyl-3-fluoro-5-methoxybenzene (**Compound I-12C**)



[00328] I-12C

[00329] 1-fluoro-3-methoxy-5-vinylbenzene (300 mg, 1.97 mmol) was added to methanol (20 mL) at room temperature and 10% palladium on carbon (200 mg) was added. Under the introduction of H₂, the mixture was stirred at room temperature for 16 hours. The mixture was
 5 filtered and washed with methanol (30 mL ×3), and the organic phases were combined and concentrated to obtain a colorless liquid crude product of 1-ethyl-3-fluoro-5-methoxybenzene (**Compound I-12C**) (200 mg, yield 66%).

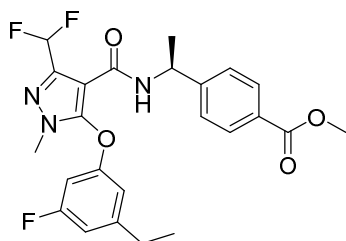
[00330] Third step: 3-ethyl-5-fluorophenol (**Compound I-12D**)



[00331] I-12D

[00332] 1-ethyl-3-fluoro-5-methoxybenzene (200 mg, 1.3 mmol) was added to dichloromethane (2 mL) at room temperature. The mixture was cooled to 0°C, and BBr₃ (1 mol/L in dichloromethane, 2 mL) was added. The mixture was naturally warmed to room temperature, and stirred at room temperature for 4 hours. The reaction was quenched by adding water (3 mL), and the mixture was extracted with dichloromethane (5 mL ×3) and separated.
 15 The organic phases were combined and dried over anhydrous sodium sulfate, filtered, and concentrated to obtain a colorless liquid crude product of 3-ethyl-5-fluorophenol (**Compound I-12D**) (180 mg, yield 99%).

[00333] Fourth step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-12E**)

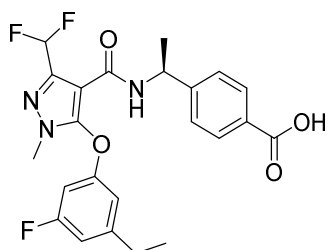


[00334] I-12E

[00335] A compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (250 mg, 0.67 mmol) was added to N, N-dimethylformamide (15 mL) and heated to 120°C; 3-ethyl-5-fluorophenol (180 mg, 1.28 mmol), potassium carbonate (270 mg, 1.96 mmol), cuprous iodide (50 mg, 0.26 mmol) and 1, 10-phenanthroline (90 mg, 0.50 mmol) were added. The mixture was stirred for 1 hour at room temperature. Then, the mixture was cooled to room temperature, diluted with water (30 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a colorless liquid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-12E**) (200 mg, yield 62%).

[00336] LC-MS, M/Z (ESI): 476.2 (M+1).

[00337] Fifth step: (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-12**)



[00338] I-12

[00339] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (200 mg, 0.42 mmol) was added to tetrahydrofuran (10 mL) and water (5 mL) at room temperature, and lithium hydroxide monohydrate (53 mg, 1.26 mmol) was added. The mixture was stirred at room temperature for 16 hours. The reaction mixture was concentrated to obtain a white solid of the compound (S)-

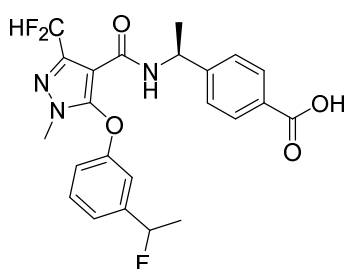
4-(1-(3-(difluoromethyl)-5-(3-ethyl-5-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (130 mg, 67% yield).

[00340] ¹H NMR (400m Hz, DMSO-d₆) δ12.8 (s, 1H), 8.10 (d, 1H), 7.74 (d, 2H), 7.16 (d, 2H), 7.11 (t, 1H), 6.93 (d, 1H), 6.72 (s, 1H), 6.69 (d, 1H), 4.92 (t, 1H), 3.73 (s, 3H), 2.58-2.54 (m, 2H), 1.25 (d, 3H), 1.13 (t, 3H).

[00341] LC-MS, M/Z (ESI): 462.1 (M+1)

[00342] Example 13: Preparation of Compound I-13

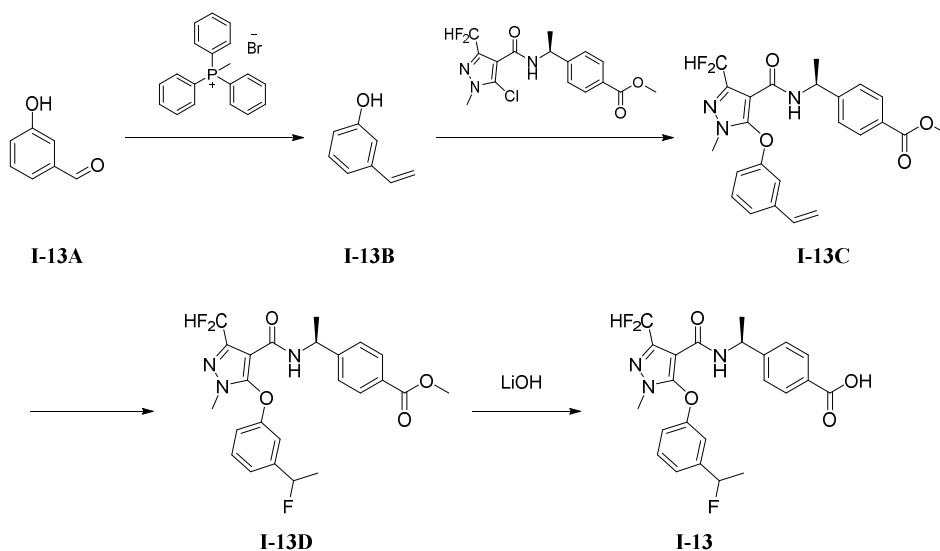
[00343] 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-13**)



I-13

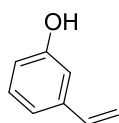
[00344]

[00345] The synthesis scheme for **Compound I-13** is shown below:



[00346]

[00347] First step: 3-vinylphenol (**Compound I-13B**)

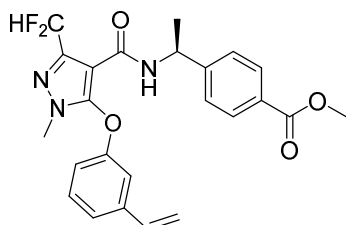


[00348] **I-13B**

[00349] A compound methyltriphenylphosphine bromide (7.30 g, 20.5 mmol) was added to

anhydrous tetrahydrofuran (40 mL) at 0°C, and potassium tert-butoxide (2.3 g, 20.5 mmol) was added. The mixture was stirred at low temperature for 2 hours. A solution of 3-hydroxybenzaldehyde (1.0 g, 8.20 mmol) and tetrahydrofuran (15 mL) was added dropwise, and after completion of addition, the mixture was warmed up naturally to room temperature and stirred for 16 hours. The mixture was diluted with water (100 mL), extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a yellow liquid of 3-vinylphenol (**Compound I-13B**) (500 mg, yield 49%).

[00350] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13C**)

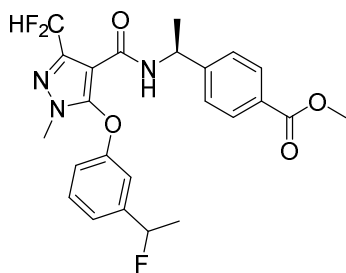


[00351] **I-13C**

[00352] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (600 mg, 1.62 mmol) was added to N, N-dimethylformamide (18 mL) at room temperature; and 3-vinylphenol (**Compound I-13B**) (388 mg, 3.23 mmol) and potassium hydroxide (181 mg, 3.23 mmol) were added. The mixture was heated to 120°C in the microwave and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (60 mL), extracted with ethyl acetate (60 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13C**) (200 mg, yield 27%).

[00353] LC-MS, M/Z (ESI): 456.2 (M+1).

[00354] Third step: methyl 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13D**)

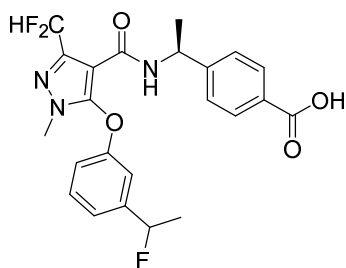


[00355] I-13D

[00356] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13C**) (100 mg, 0.22 mmol) was added to acetonitrile (25 mL); and a selective fluorinating reagent (233 mg, 0.66 mmol), palladium tetrakis(triphenylphosphonium) (25 mg, 0.02 mmol) and triethylsilane (40.0 mg, 0.33 mmol) were added at room temperature. The mixture was stirred for 16 hours under nitrogen protection at room temperature. Then, the mixture was diluted with water (60 mL), extracted with ethyl acetate (60 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated, and the residue purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 5: 1) to obtain a white solid of methyl 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13D**) (65.0 mg, yield 62%).

[00357] LC-MS, M/Z (ESI): 476.3 (M+1)

[00358] Fourth step: 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-13**)



[00359] I-13

[00360] A starting material methyl 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-13D**) (150 mg, 0.19 mmol) was added to tetrahydrofuran (24 mL); and water (12 mL) and methanol (12 mL), lithium hydroxide (15.0 mg, 0.36 mmol) were added at room temperature. The mixture was stirred for 4 hours at room temperature. The reaction mixture was concentrated

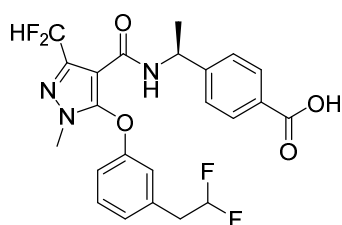
to obtain, by the acidic preparation method A, a white solid of 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-fluoroethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-13**) (50.0 mg, 32% yield).

[00361] ^1H NMR (400m Hz, DMSO- d_6) δ 12.8 (s, 1H), 8.03 (d, 1H), 7.72 (d, 2H), 7.45 (t, 1H), 7.25 (t, 1H), 7.11 (d, 2H), 6.98 (t, 1H), 6.94 (t, 1H), 5.78 (dd, 1H), 4.89 (m, 1H), 3.73 (s, 3H), 3.60 (s, 1H), 1.55 (dd, 3H), 1.24 (d, 3H).

[00362] LC-MS, M/Z (ESI): 462.3 (M+1)

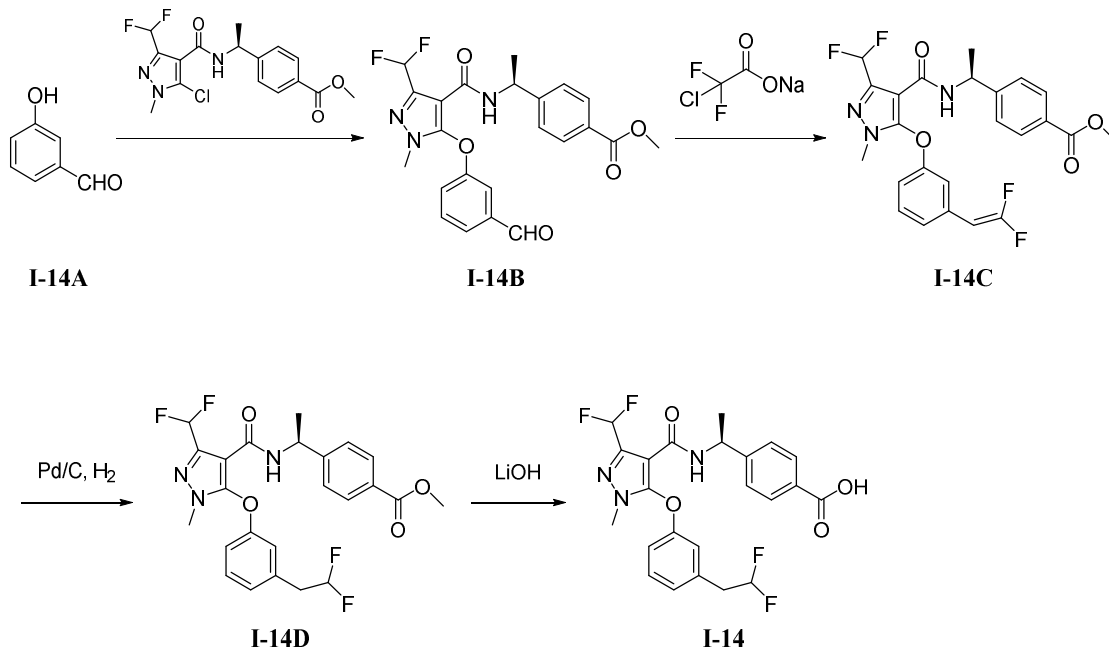
[00363] Example 14: Preparation of Compound I-14

[00364] (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-14**)



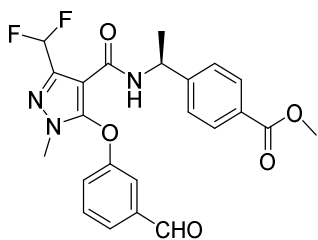
[00365] **I-14**

[00366] The synthesis scheme for **Compound I-14** is shown below:



[00367]

[00368] First step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14B**)

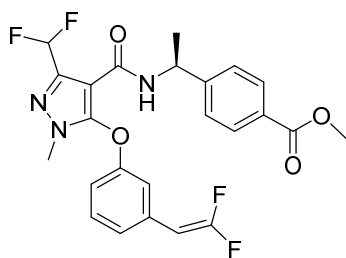


[00369] I-14B

[00370] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (574 mg, 1.54 mmol) was added to N, N-dimethylformamide (15 mL) at room temperature; and 3-hydroxybenzaldehyde (378 mg, 3.09 mmol), copper iodide (60 mg, 0.31 mmol), 1, 10-phenanthroline (114 mg, 0.62 mmol) and cesium carbonate (1.50 g, 4.62 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (50 mL), and the pH was adjusted to 4 with 1N hydrochloric acid. The mixture was extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a light yellow solid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14B**) (280 mg, yield 40%).

[00371] LC-MS, M/Z (ESI): 458.3 [M+H]⁺.

[00372] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-(2,2-difluorovinyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14C**)



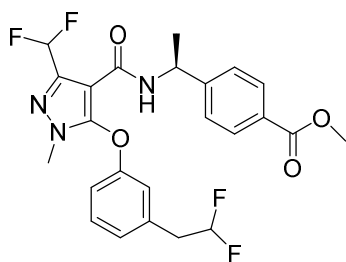
[00373] I-14C

[00374] The compound methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-formylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14B**) (280 mg, 0.61 mmol) was added to N, N-dimethylformamide (15 mL) at room temperature, and triphenylphosphine (200 mg, 0.76 mmol) and sodium difluorochloroacetate (140 mg, 0.92 mmol) were added, and the

mixture was heated to 100°C in the microwave under nitrogen protection and stirred for 1 h. The mixture was cooled to room temperature, diluted by addition of water (50 mL), and the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow solid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-(2,2-difluorovinyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14C**) (120 mg, yield 40%).

[00375] LC-MS, M/Z (ESI): 492.4 [M+H]⁺.

[00376] Third step: methyl (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14D**)

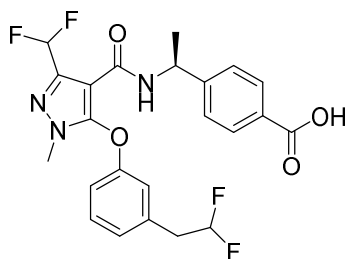


[00377] **I-14D**

[00378] The compound methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-(2,2-difluorovinyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14C**) (120 mg, 0.24 mmol) was added to methanol (20 mL) at room temperature; 10% palladium on carbon (20 mg) was added; and hydrogen was introduced. The mixture was stirred at room temperature for 16 hours. The mixture was filtered and concentrated to obtain a light yellow solid crude product methyl (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14D**) (120 mg, 99% yield).

[00379] LC-MS, M/Z (ESI): 494.4 [M+H]⁺.

[00380] Fourth step: (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-14**)



[00381] **I-14**

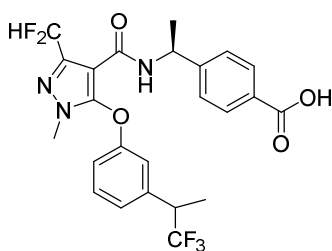
[00382] A starting material methyl (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-14D**) (120 mg, 0.24 mmol) was added to tetrahydrofuran (5 mL) and water (5 mL) at room temperature, and then lithium hydroxide (31 mg, 0.72 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-(3-(2,2-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-14**) (90 mg, 77% yield).

[00383] ^1H NMR (400 MHz, DMSO- d_6) δ 12.8 (s, 1H), 7.95 (d, 1H), 7.74 (d, 2H), 7.39 (t, 1H), 7.25 (t, 1H), 7.16 (d, 2H), 7.13 (d, 1H), 7.04 (d, 1H), 6.91 (t, 1H), 6.34 (dt, 1H), 4.91 (t, 1H), 3.71 (s, 3H), 3.22 (q, 2H), 1.22 (d, 3H).

[00384] LC-MS, M/Z (ESI): 480.4 $[\text{M}+\text{H}]^+$

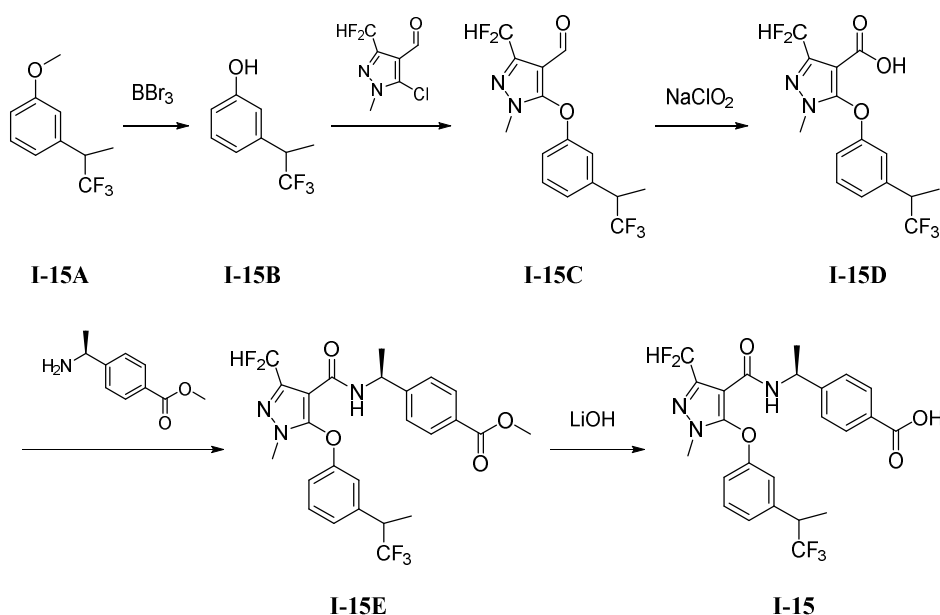
[00385] Example 15: Preparation of Compound I-15

[00386] 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-15**)



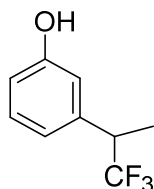
[00387] **I-15**

[00388] The synthesis scheme for **Compound I-15** is shown below:



[00389]

[00390] First step: 3-(1,1,1-trifluoroprop-2-yl)phenol (**Compound I-15B**)

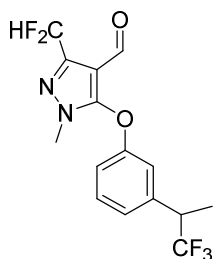


[00391] **I-15B**

[00392] A compound 1-methoxy-3-(1, 1, 1-trifluoroprop-2-yl) benzene (800 mg, 0.72 mmol)

5 was added to dichloromethane (1 mL) at room temperature, and then the mixture was cooled to -78°C, and then boron tribromide (1.50 g, 5.88 mmol) was added. The mixture was stirred at room temperature for 4 hours; and then the mixture was diluted with water (50 mL), extracted with dichloromethane (50 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated and the residue purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 5: 1) to obtain a light yellow liquid of 3-(1,1,1-trifluoroprop-2-yl)phenol (**Compound I-15B**) (640 mg, 86% yield).

[00393] Second step: 3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoroprop-2-yl)phenoxy)-1H-pyrazole-4-carbaldehyde (**Compound I-15C**)

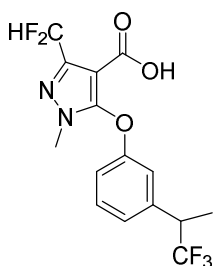


[00394] I-15C

[00395] The compound 5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (660 mg, 3.37 mmol) was added to N, N dimethylformamide (6 mL) at room temperature, and then 3-(1, 1, 1-trifluoropropan-2-yl) phenol (640 mg, 3.37 mmol) and potassium carbonate (930 mg, 6.72 mmol) were added. The mixture was heated to 100°C under nitrogen protection and stirred for 8 h; and then the mixture was cooled to room temperature, diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 5: 1) to obtain a light yellow liquid of 33-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carbaldehyde (**Compound I-15C**) (1.0 g, yield 18%).

[00396] LC-MS, M/Z (ESI): 349.2 [M+H]⁺.

[00397] Third step: 3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxylic acid (**Compound I-15D**)



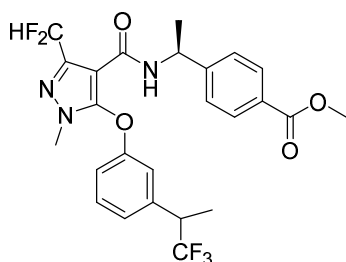
[00398] I-15D

[00399] A compound 3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carbaldehyde (800 mg, 2.30 mmol) was added to tert-butanol (8 mL) and water (8 mL) at room temperature. 2-methyl-2-butene (322 mg, 4.60 mmol), sodium chlorite (414 mg, 4.60 mmol) and sodium dihydrogen phosphate (617 mg, 5.06 mmol) were added. The mixture was stirred at room temperature for 8 h; and then the mixture was diluted with water (15 mL), the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl

acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a light yellow solid of crude 3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxylic acid (**Compound I-15D**) (850 mg, yield 100%).

5 **[00400]** LC-MS, M/Z (ESI): 365.1 [M+H]⁺.

[00401] Fourth step: methyl 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-15E**)

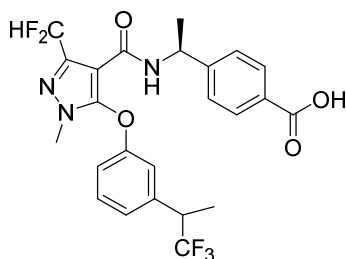


[00402] **I-15E**

10 **[00403]** A compound 3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxylic acid (850 mg, 2.34 mmol) was added to N, N-dimethylformamide (10 mL); and methyl (S)-4-(1-aminoethyl) benzoate (418 mg, 2.34 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (1.3 g, 3.50 mmol) and N, N-diisopropylethylamine (900 mg, 7.0 mmol) were added. The mixture was stirred at
15 room temperature for 16 hours, diluted with water (40 mL), extracted with ethyl acetate (40 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated and purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a light yellow liquid of methyl 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate
20 (**Compound I-15E**) (1.0 g, yield 81%).

[00404] LC-MS, M/Z (ESI): 526.3 [M+H]⁺.

[00405] Fifth step: 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-15**)



[00406] I-15

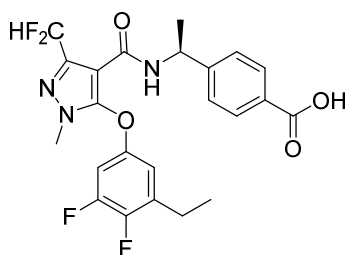
[00407] The starting material methyl 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-15E**) (1.0 g, 1.90 mmol) was added to tetrahydrofuran (10 mL); and water (2 mL) at room temperature; and lithium hydroxide (200 mg, 4.76 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 by 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparative method A, a white solid of 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(1,1,1-trifluoropropan-2-yl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-15**) (356 mg, 37% yield).

[00408] ¹H NMR (400m Hz, DMSO-d₆) δ12.8 (s, 1H), 7.97 (dd, 1H), 7.73 (d, 2H), 7.43 (dd, 1H), 7.28 (d, 1H), 7.25 (t, 1H), 7.13 (d, 2H), 7.11 (d, 1H), 6.99 (t, 1H), 4.90 (t, 1H), 3.89-3.85 (m, 1H), 3.73 (s, 3H), 1.41 (d, 3H), 1.21 (t, 3H).

[00409] LC-MS, M/Z (ESI): 512.3 [M+H]⁺

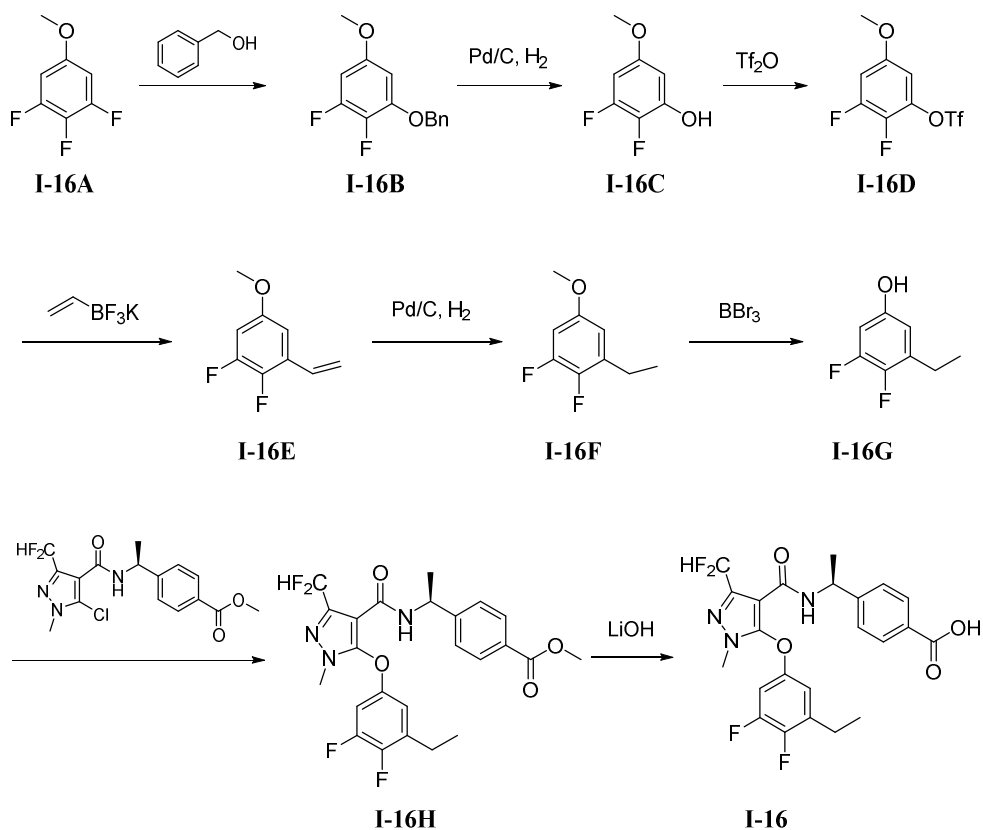
[00410] Example 16: Preparation of Compound I-16

[00411] (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-16**)



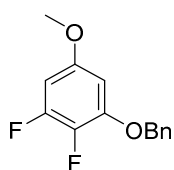
[00412] I-16

[00413] The synthesis scheme for **Compound I-16** is shown below:



[00414]

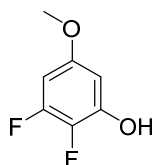
[00415] First step: 1-(benzyloxy)-2,3-difluoro-5-methoxybenzene (**Compound I-16B**)



[00416] **I-16B**

[00417] A Compound 1, 2, 3-trifluoro-5-methoxybenzene (8.5 g, 52.4 mmol) was added to
 5 toluene (100 mL) at room temperature; benzyl alcohol (11.3 g, 104 mmol) and potassium
 hydroxide (5.86 g, 104 mmol) were added; and the mixture was heated to 120°C and stirred for
 16 hours. Then, the mixture was cooled to room temperature, diluted with water (500 mL), and
 the pH was adjusted to 7-8 with 1N hydrochloric acid. The mixture was extracted with
 dichloromethane (100 mL × 3) and separated. The organic phases were combined and dried over
 10 anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified
 by a silica gel column (petroleum ether: ethyl acetate (V/V), 20: 1) to obtain a colorless liquid
 of 1-(benzyloxy)-2,3-difluoro-5-methoxybenzene (compound **I-16B**) (5.2 g, yield 40%).

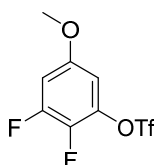
[00418] Second step: 2,3-difluoro-5-methoxyphenol (**Compound I-16C**)



[00419] I-16C

[00420] A compound 1-(benzyloxy)-2,3-difluoro-5-methoxybenzene (5.2 g, 20.8 mmol) was added to tetrahydrofuran (100 mL) at room temperature; 10% palladium on carbon (520 mg) was added; and hydrogen was introduced. The mixture was stirred at room temperature for 16 hours. Then, the mixture was filtrated, concentrated, and purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 5: 1) to obtain a colorless liquid crude product of 2,3-difluoro-5-methoxyphenol (**Compound I-16C**) (3.8 g, 100% yield).

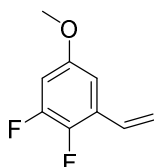
[00421] Third step: 2,3-difluoro-5-methoxyphenyl trifluoromethanesulfonate (**Compound I-16D**)



[00422] I-16D

[00423] A compound 2,3-difluoro-5-methoxyphenol (500 mg, 3.12 mmol) was added to dichloromethane (20 mL) at room temperature; and trifluoromethanesulfonic anhydride (881 mg, 3.12 mmol) and pyridine (493 mg, 6.25 mmol) were added. The mixture was stirred for 4 hours at room temperature; saturated sodium bicarbonate (20 mL) was added and then the mixture was separated and the organic phases were dried over anhydrous sodium sulfate, filtered and concentrated to obtain a colorless liquid crude product of 2,3-difluoro-5-methoxyphenyl trifluoromethanesulfonate (**Compound I-16D**) (600 mg, yield 66%).

[00424] Fourth step: 1,2-difluoro-5-methoxy-3-vinylbenzene (**Compound I-16E**)

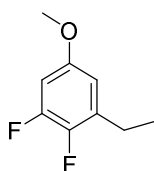


[00425] I-16E

[00426] A compound 2,3-difluoro-5-methoxyphenyl trifluoromethanesulfonate (600 mg, 2.05 mmol) was added to 1, 4-dioxane (50 mL) at room temperature; potassium vinyltrifluoroborate

(1.25 g, 9.37 mmol); potassium carbonate (1.30 g, 9.37 mmol) and dichloro [1, 1'-bis(diphenylphosphino) ferrocene] palladium (254 mg, 0.31 mmol) were added; and the mixture was heated to 90°C under nitrogen protection and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (200 mL), extracted with
5 dichloromethane (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 100: 1) to obtain a light yellow liquid of 1,2-difluoro-5-methoxy-3-vinylbenzene (**Compound I-16E**) (200 mg, yield 57%).

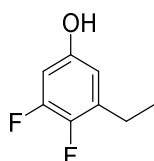
[00427] Fifth step: 1-ethyl-2,3-difluoro-5-methoxybenzene (**Compound I-16F**)



10 **[00428]** **I-16F**

[00429] A compound 1,2-difluoro-5-methoxy-3-vinylbenzene (130 mg, 0.82 mmol) was added to methanol (15 mL) at room temperature; 10% palladium on carbon (20 mg) was added; and hydrogen was introduced under stirring at room temperature for 16 hours. The mixture was filtered and concentrated to obtain 1-ethyl-2,3-difluoro-5-methoxybenzene (**Compound I-16F**)
15 (100 mg, 76% yield) a light yellow liquid.

[00430] Sixth step: 3-ethyl-4,5-difluorophenol (**Compound I-16G**)

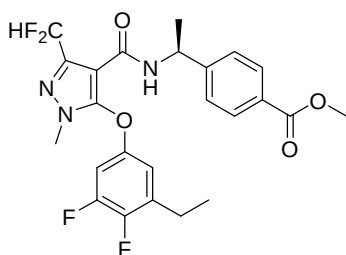


[00431] **I-16G**

[00432] A compound 1-ethyl-2,3-difluoro-5-methoxybenzene (100 mg, 0.58 mmol) was added to dichloromethane (10 mL) at room temperature; boron tribromide (726 mg, 2.90 mmol) was
20 added at 0°C under stirring for 3 h; and the mixture was diluted by addition of water (20 mL), extracted with dichloromethane (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a light yellow liquid crude product of 3-ethyl-4,5-difluorophenol (**Compound I-16G**) (60 mg, yield 66%).

[00433] Seventh step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy))-1-

methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-16H**)

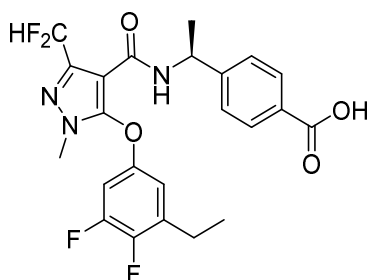


[00434] I-16H

[00435] A compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (600 mg, 1.61 mmol) was added to N, N-dimethylformamide (15 mL) at room temperature; and 3-ethyl-4, 5-difluorophenol (450 mg, 3.0 mmol) and potassium hydroxide (210 mg, 3.9 mmol) were added. The mixture was heated to 120°C in the microwave and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (50 mL), the pH was adjusted to 4 with 1N hydrochloric acid. The mixture was extracted with ethyl acetate (60 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated and the residue purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow solid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-16H**) (350 mg, yield 44%).

[00436] LC-MS, M/Z (ESI): 494.4 [M+H]⁺.

[00437] Eighth step: (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-16**)



[00438] I-16

[00439] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (325 mg, 0.66 mmol) was added to tetrahydrofuran (15 mL), methanol (3 mL) and water (3 mL) at room temperature; and lithium hydroxide (32 mg, 1.32 mmol) was added. The mixture was stirred at room

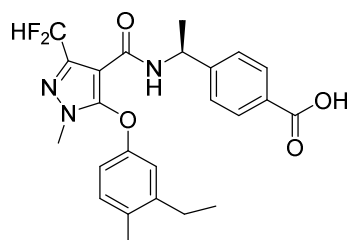
temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4,5-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (170 mg, 54% yield).

5 **[00440]** ^1H NMR (400m Hz, DMSO- d_6) δ 12.8 (s, 1H), 8.15 (d, 1H), 7.76 (d, 2H), 7.24 (t, 1H), 7.18 (d, 2H), 7.01 (t, 1H), 6.80 (d, 1H), 4.94 (t, 1H), 3.74 (s, 3H), 2.62 (q, 2H), 1.27 (d, 3H), 1.11 (t, 3H).

[00441] LC-MS, M/Z (ESI): 480.4 $[\text{M}+\text{H}]^+$

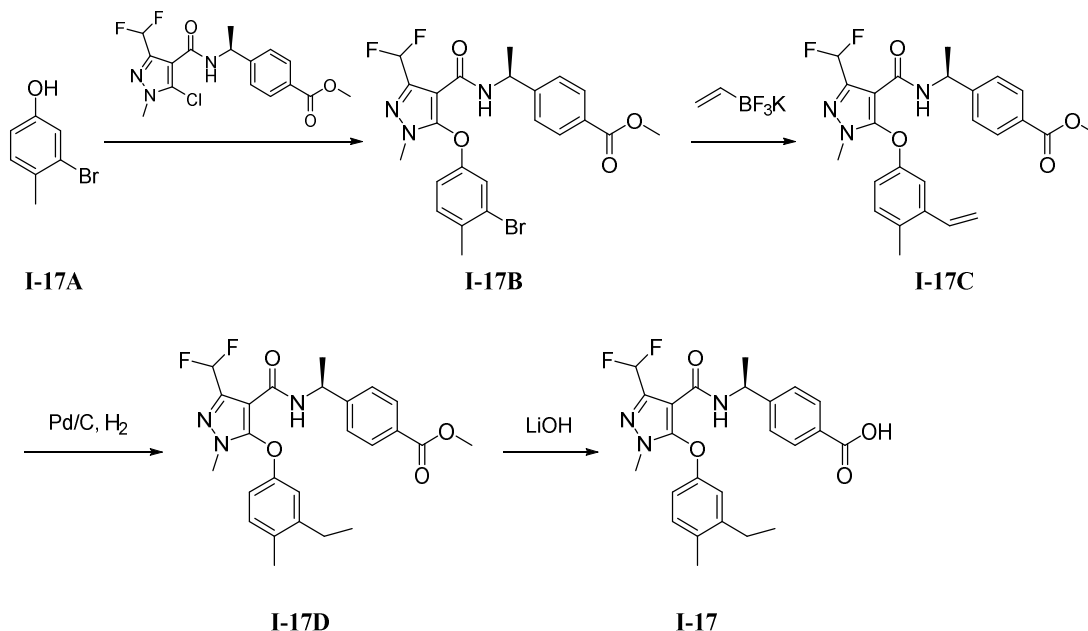
[00442] Example 17: Preparation of Compound I-17

10 **[00443]** (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-17**)



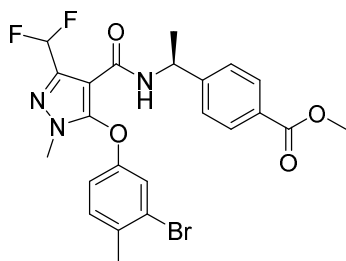
[00444] **I-17**

[00445] The synthesis scheme for **Compound I-17** is shown below:



[00446]

15 **[00447]** First step: methyl (S)-4-(1-(5-(3-bromo-4-methylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17B**)

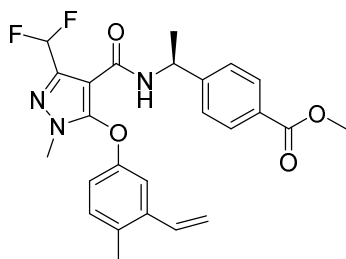


[00448] I-17B

[00449] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (500 mg, 1.35 mmol) was added to N, N-dimethylformamide (10 mL) at room temperature; 3-bromo-4-methylphenol (380 mg, 2.02 mmol), cuprous iodide (100 mg, 0.52 mmol), 1, 10-phenanthroline (180 mg, 1.0 mmol) and cesium carbonate (1.30 g, 4.05 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (50 mL), and the pH was adjusted to 4 with 1N hydrochloric acid. The mixture was extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a light yellow solid of methyl (S)-4-(1-(5-(3-bromo-4-methylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17B**) (270 mg, yield 38%).

[00450] LC-MS, M/Z (ESI): 522.3 [M+H]⁺.

[00451] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-methyl-3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17C**)



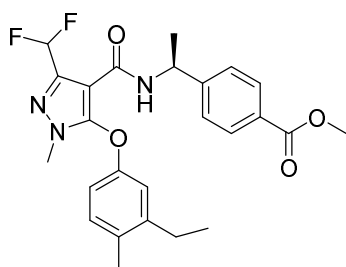
[00452] I-17C

[00453] The compound methyl (S)-4-(1-(5-(3-bromo-4-methylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17B**) (260 mg, 0.50 mmol) was added to 1, 4-dioxane (5 mL) at room temperature; dichloro[1, 1'-

bis(diphenylphosphino)ferrocene]palladium (73 mg, 0.10 mmol), potassium vinyltrifluoroborate (100 mg, 0.75 mmol), and potassium carbonate (140 mg, 1.0 mmol) were added; and the mixture was heated to 100°C under nitrogen protection and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (50 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated. The residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-methyl-3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17C**) (160 mg, yield 68%).

[00454] LC-MS, M/Z (ESI): 470.3 [M+H]⁺.

[00455] Third step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17D**)

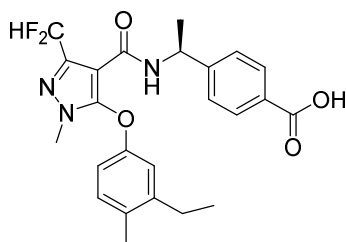


[00456] **I-17D**

[00457] The compound methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-methyl-3-vinylphenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (160 mg, 0.34 mmol) was added to methanol (10 mL) at room temperature and 10% palladium on carbon (20 mg) was added; and hydrogen was introduced under stirring for 16 hours at room temperature; and the mixture was filtered and concentrated to obtain a light yellow solid crude product of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17D**) (150 mg, yield 93%).

[00458] LC-MS, M/Z (ESI): 472.3 [M+H]⁺

[00459] Fourth step: (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-17**)



[00460] I-17

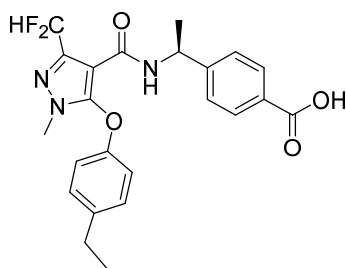
[00461] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-17D**) (150 mg, 0.32 mmol) was added to tetrahydrofuran (5 mL) and water (5 mL) at room temperature, and lithium hydroxide (24 mg, 1.0 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 by 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-methylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (85 mg, 58% yield).

[00462] ^1H NMR (400 MHz, DMSO- d_6) δ 12.7 (s, 1H), 7.90 (d, 1H), 7.69 (d, 2H), 7.25 (t, 1H), 7.13 (d, 1H), 7.11 (d, 2H), 6.85 (d, 1H), 6.64 (dd, 1H), 4.93 (t, 1H), 3.71 (s, 3H), 2.55 (q, 2H), 2.23 (s, 3H), 1.24 (d, 3H), 1.07 (t, 3H).

[00463] LC-MS, M/Z (ESI): 458.3 $[\text{M}+\text{H}]^+$

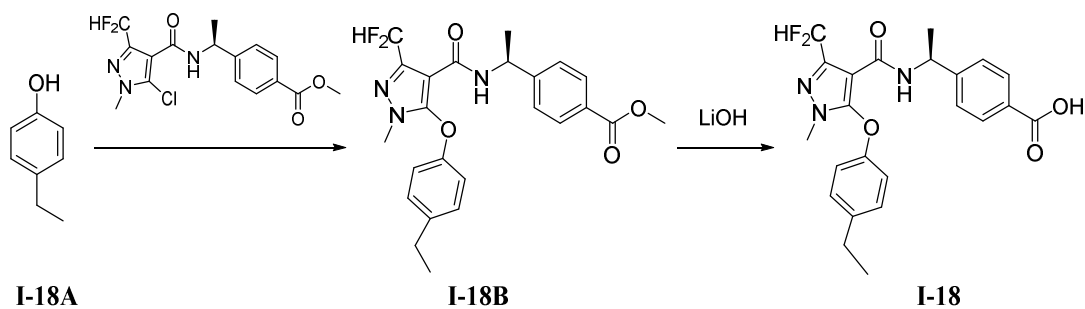
[00464] Example 18: Preparation of Compound I-18

[00465] (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl) benzoic acid (**Compound I-18**)

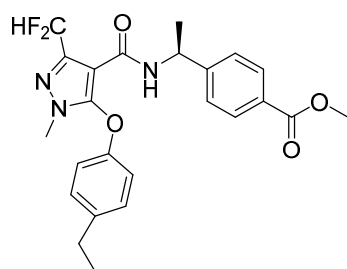


[00466] I-18

[00467] The synthesis scheme for **Compound I-18** is shown below:



[00469] First step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-18B**)



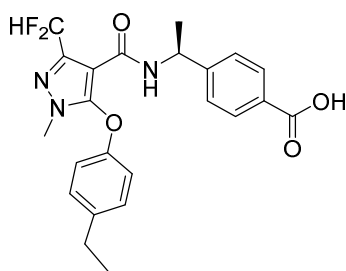
[00470] I-18B

5 **[00471]** The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (371 mg, 1.0 mmol) was added to dimethyl sulfoxide (3 mL) at room temperature; 4-ethylphenol (122 mg, 1.0 mmol) and potassium hydroxide (114 mg, 2.0 mmol) were added; and the mixture was heated to 120°C in the microwave and stirred for 20 min. Then, the mixture was cooled to room temperature and

10 diluted with water (20 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (20 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-18B**) (175 mg, yield 38%).

15 **[00472]** LC-MS, M/Z (ESI): 458.3 [M+H]⁺

[00473] Second step: (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-18**)



[00474] I-18

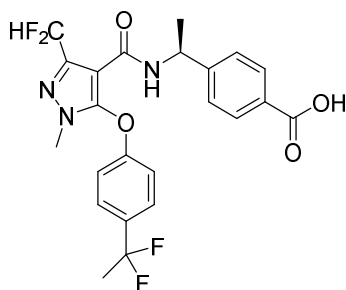
[00475] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-18B**) (175 mg, 0.38 mmol) was added to tetrahydrofuran (2 mL), methanol (2 mL) and water (1 mL) at room temperature; and lithium hydroxide (3 mg, 0.16 mmol) was added. The mixture was stirred for 12 hours at room temperature; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(4-ethylphenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-18**) (40 mg, 24% yield).

[00476] ^1H NMR (400MHz, DMSO- d_6) δ 12.7 (s, 1H), 7.95 (d, 1H), 7.71 (d, 2H), 7.24 (d, 2H), 7.21 (t, 1H), 7.11 (d, 2H), 6.92 (d, 2H), 4.91 (t, 1H), 3.72 (s, 3H), 2.62 (q, 2H), 1.24 (d, 3H), 1.19 (t, 3H).

[00477] LC-MS, M/Z (ESI): 444.3 $[\text{M}+\text{H}]^+$

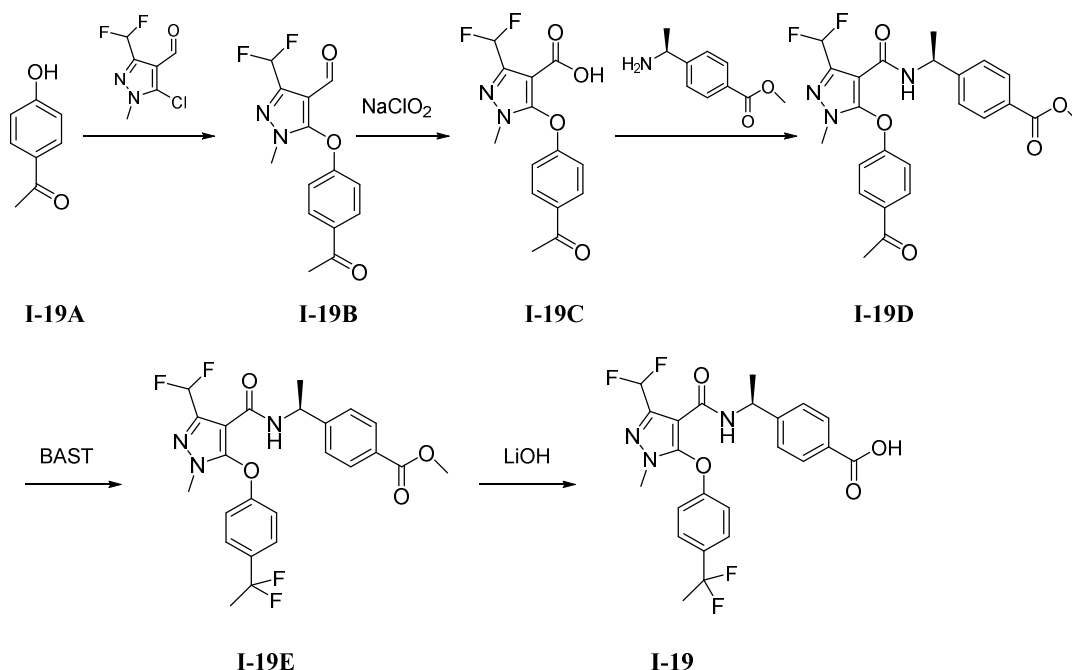
[00478] Example 19: Preparation of Compound I-19

[00479] (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-19**)



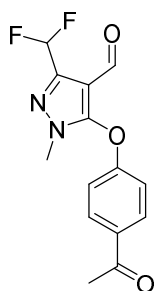
[00480] I-19

[00481] The synthesis scheme for **Compound I-19** is shown below:



[00482]

[00483] First step: 5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (**Compound I-19B**)



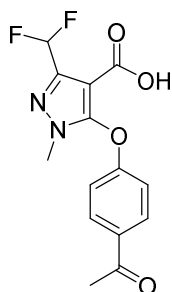
[00484] **I-19B**

- 5 [00485] A compound 5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (1.0 g, 5.15 mmol) was added to N, N-dimethylformamide (15 mL) at room temperature; 4-acetylphenol (1.05 g, 7.73 mmol), cuprous iodide (918 mg, 5.15 mmol), 1, 10-phenanthroline (1.05 g, 7.73 mmol) and potassium carbonate (1.42 g, 10.3 mmol) were added; and the mixture was heated to 100°C under nitrogen protection and stirred for 2 hours. Then, the mixture was
- 10 cooled to room temperature, diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a light yellow solid crude product of 5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (**Compound I-19B**) (270 mg, yield

18%).

[00486] LC-MS, M/Z (ESI): 295.1 [M+H]⁺

[00487] Second step: 5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxylic acid (**Compound I-19C**)

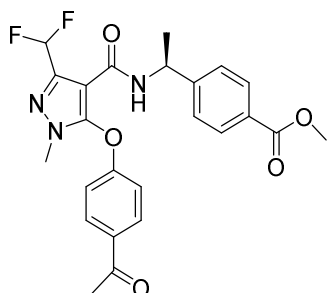


5 **[00488]** **I-19C**

[00489] The compound 5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (**Compound I-19B**) (400 mg, 1.36 mmol) was added to tert-butanol (9 mL) and water (3 mL) at room temperature; and 2-methyl-2-butene (476 mg, 6.80 mmol), sodium chlorite (369 mg, 4.08 mmol) and sodium dihydrogen phosphate (408 mg, 3.40 mmol) were added. The mixture was stirred for 14 hours at room temperature; the mixture was diluted by addition of water (15 mL), the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain crude 5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxylic acid (compound **I-19C**) (350 mg, yield 100%) a light yellow solid.

[00490] LC-MS, M/Z (ESI): 311.1 [M+H]⁺.

[00491] Third step: methyl (S)-4-(1-(5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19D**)



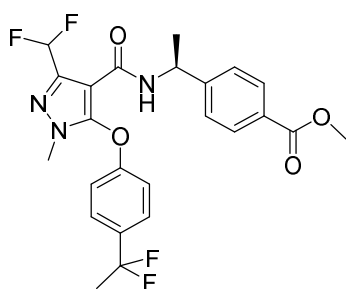
[00492] **I-19D**

20 **[00493]** A compound 3-(difluoromethyl)-5-(3-isopropylphenoxy)-1-methyl-1H-pyrazole-4-

carboxylic acid (350 mg, 1.13 mmol) was added to N, N-dimethylformamide (7 mL); methyl (S)-4-(1-aminoethyl) benzoate (242 mg, 1.36 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (650 mg, 1.71 mmol) and N, N-diisopropylethylamine (437 mg, 3.67 mmol) were added; and the mixture was stirred at room temperature for 16 hours, diluted with water (40 mL), extracted with ethyl acetate (40 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated, and purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow liquid of methyl (S)-4-(1-(5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19D**) (500 mg, yield 94%).

[00494] LC-MS, M/Z (ESI): 472.2 [M+H]⁺

[00495] Fourth step: methyl (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19E**)



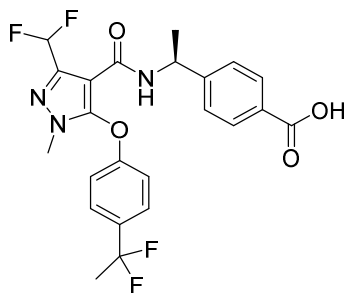
[00496] **I-19E**

[00497] The compound methyl (S)-4-(1-(5-(4-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19D**) (50 mg, 0.53 mmol) was added to bis(2-methoxyethyl)aminosulfur trifluoride (5 mL) at room temperature. The mixture was heated to 50°C and stirred for 48 h; then the mixture was cooled to room temperature, and added dropwise to a saturated solution of sodium bicarbonate (50 mL) and extracted with ethyl acetate (40 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a light yellow solid crude product of methyl (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19E**) (250 mg, yield 96%).

[00498] LC-MS, M/Z (ESI): 494.5 [M+H]⁺.

[00499] Fifth step: (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-

1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-19**)



[00500] **I-19**

[00501] A starting material methyl (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-19E**)

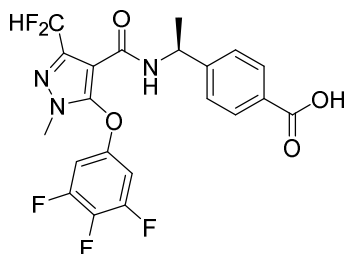
5 (250 mg, 0.51 mmol) was added to tetrahydrofuran (4 mL); and methanol (2 mL) and water (2 mL) at room temperature and lithium hydroxide (61 mg, 2.55 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-(4-(1,1-difluoroethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-19**) (19 mg, 8% yield).

[00502] ^1H NMR (400m Hz, DMSO- d_6) δ 12.8 (s, 1H), 8.11 (d, 1H), 7.71 (d, 2H), 7.60 (d, 2H), 7.25 (t, 1H), 7.12 (d, 2H), 7.08 (d, 2H), 4.90 (t, 1H), 3.73 (s, 3H), 2.01 (t, 3H), 1.23 (d, 3H).

[00503] LC-MS, M/Z (ESI): 480.5 $[\text{M}+\text{H}]^+$

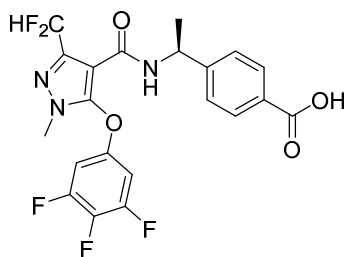
[00504] Example 20: Preparation of Compound I-20

15 **[00505]** (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3,4,5-trifluorophenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-20**)



[00506] **I-20**

[00507] The synthesis scheme for **Compound I-20** is shown below:



[00514] I-20

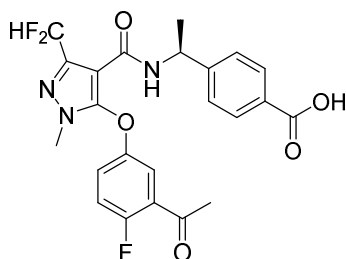
[00515] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3,4,5-trifluorophenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (80 mg, 0.16 mmol) was added to tetrahydrofuran (2 mL) and water (1 mL) at room temperature; lithium hydroxide (17 mg, 0.40 mmol) was added; and the mixture was heated to 50°C and stirred for 2 hours. Then, the mixture was cooled to room temperature; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3,4,5-trifluorophenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-20**) (7 mg, 9.0% yield).

[00516] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.24 (d, 1H), 7.78 (d, 2H), 7.24 (d, 2H), 7.22 (t, 1H), 7.09 (d, 1H), 7.02 (d, 1H), 4.96 (t, 1H), 3.73 (s, 3H), 1.30 (d, 3H).

[00517] LC-MS, M/Z (ESI): 470.1 [M+H]⁺

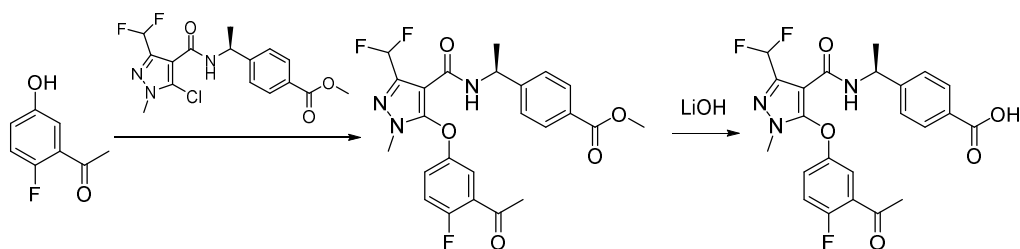
[00518] Example 21: Preparation of Compound I-21

[00519] (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-21**)



[00520] I-21

[00521] The synthesis scheme for **Compound I-21** is shown below:

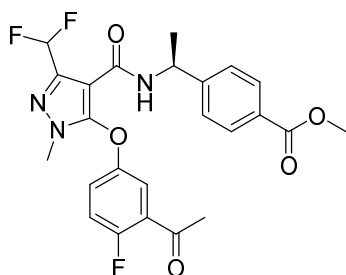


[00522] I-21A

I-21B

I-21

[00523] First step: methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-21B**)



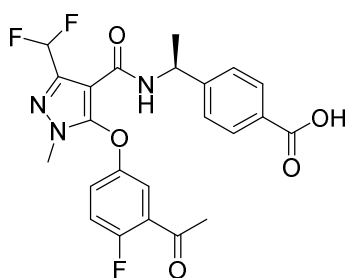
I-21B

[00524]

5 **[00525]** A compound 1-(2-fluoro-5-hydroxyphenyl) ethan-1-one (1.58 g, 10.8 mmol) was added to N, N-dimethylformamide (30 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (2.0 g, 5.4 mmol) and potassium hydroxide (450 mg, 8.1 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2
10 hours; the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a white solid of methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-21B**) (346 mg, yield 13%).

[00526] LC-MS, M/Z (ESI): 490.2 [M+H]⁺

[00527] Second step: (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-21**)



[00528] I-21

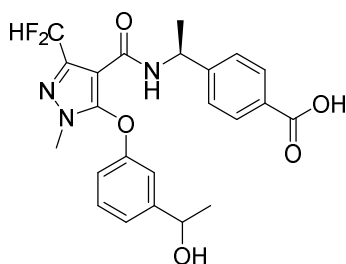
[00529] The starting material methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (compound **I-21B**) (65 mg, 0.13 mmol) was added to tetrahydrofuran (5 mL), methanol (5 mL) and water (1 mL) at room temperature; and lithium hydroxide (20 mg, 0.47 mmol) was added. The mixture was stirred at room temperature for 5 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-21**) (30 mg, 66% yield).

[00530] ^1H NMR (400MHz, DMSO- d_6) δ 12.8 (s, 1H), 8.18 (d, 1H), 7.72 (d, 2H), 7.38-7.32 (m, 2H), 7.29-7.24 (m, 1H), 7.26 (t, 1H), 7.16 (d, 2H), 4.91 (t, 1H), 3.73 (s, 3H), 2.54 (d, 3H), 1.24 (d, 3H).

[00531] LC-MS, M/Z (ESI): 476.1 $[\text{M}+\text{H}]^+$

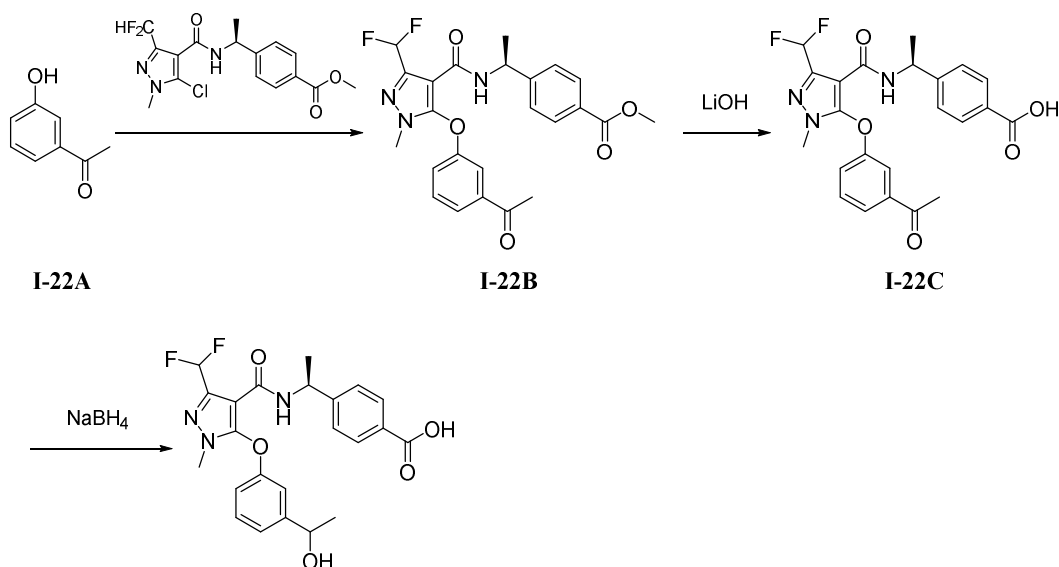
[00532] Example 22: Preparation of **Compound I-22**

[00533] 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22**)



[00534] I-22

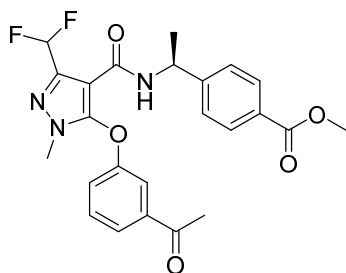
[00535] The synthesis scheme for **Compound I-22** is shown below:



[00536]

I-22

[00537] First step: methyl (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-22B**)



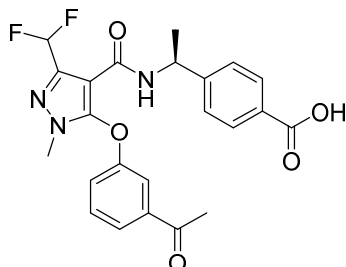
[00538]

I-22B

- 5 [00539] A compound 1-(3-hydroxyphenyl) ethan-1-one (275 mg, 2.02 mmol) was added to N, N-dimethylformamide (6 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (500 mg, 1.35 mmol), copper iodide (100 mg, 0.52 mmol), 1, 10-phenanthroline (180 mg, 1.0 mmol) and cesium carbonate (1.30 g, 4.05 mmol) were added. The mixture was heated to 120°C in the
- 10 microwave under nitrogen protection and stirred for 2 hours. Then the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V)
- 15 = 3: 1) to obtain a white solid of methyl (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-22B**) (300 mg, yield 47%).

[00540] LC-MS, M/Z (ESI): 472.3 [M+H]⁺

[00541] Second step: (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22C**)

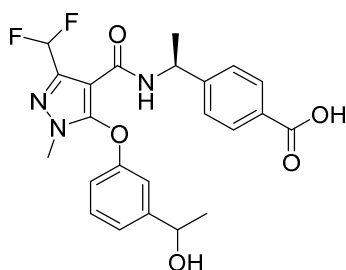


[00542] **I-22C**

5 [00543] The starting material methyl (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-22B**) (290 mg, 0.62 mmol) was added to tetrahydrofuran (5 mL) and water (5 mL) at room temperature; and lithium hydroxide (45 mg, 1.87 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was
10 concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22C**) (180 mg, 64% yield).

[00544] LC-MS, M/Z (ESI): 458.3 [M+H]⁺

[00545] Third step: 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22**)
15



[00546] **I-22**

[00547] The compound (S)-4-(1-(5-(3-acetylphenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22C**) (170 mg, 0.37 mmol) was added to methanol (15 mL) and water (15 mL) at room temperature; and sodium borohydride
20 (15 mg, 0.39 mmol) was added. The mixture was stirred for 2 hours at room temperature; and

the pH was adjusted to 4 with 1N hydrochloric acid and concentrated to obtain 4-((1S)-1-(3-(difluoromethyl)-5-(3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-22**) (168 mg, 98% yield) a white solid by the acidic preparation method A.

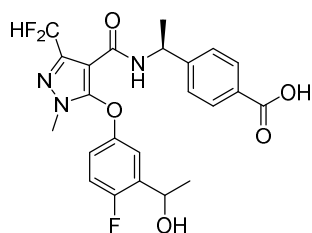
5 **[00548]** LC-MS, M/Z (ESI): 460.3 [M+H]⁺

[00549] ¹H NMR (400mHz, DMSO-d₆) δ12.7 (s, 1H), 7.92 (d, 1H), 7.73 (d, 2H), 7.36 (t, 1H), 7.26 (t, 1H), 7.18 (d, 1H), 7.13 (d, 2H), 7.11 (t, 1H), 6.83 (dd, 1H), 5.26 (d, 1H), 4.92 (t, 1H), 4.71 (s, 1H), 3.72 (s, 3H), 1.28 (dd, 3H), 1.22 (d, 3H).

[00550] LC-MS, M/Z (ESI): 478.1 [M+H]⁺

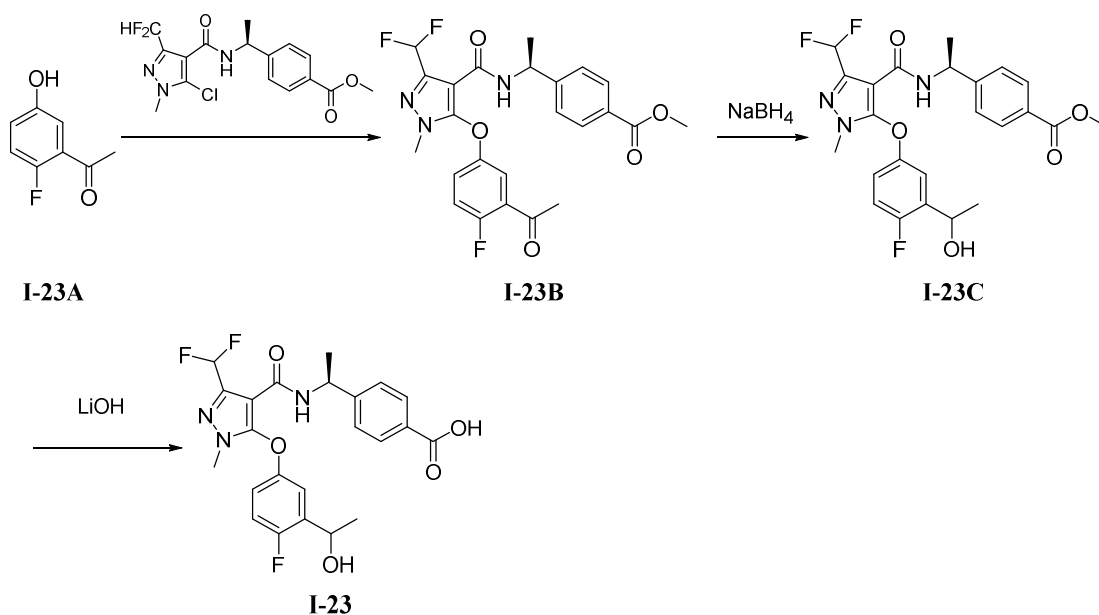
10 **[00551]** Example 23: Preparation of Compound I-23

[00552] 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-23**)



[00553] **I-23**

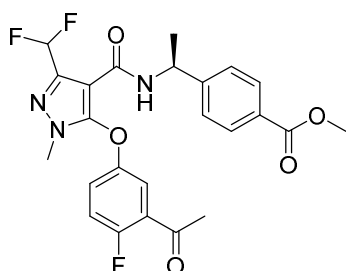
[00554] The synthesis scheme for **Compound I-23** is shown below:



15 **[00555]**

[00556] First step: methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-

methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23B**)



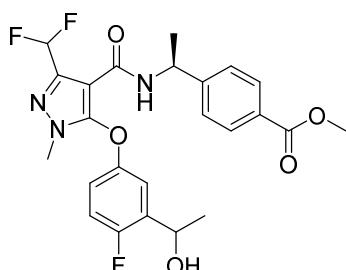
[00557] I-23B

[00558] A compound 1-(2-fluoro-5-hydroxyphenyl) ethan-1-one (1.58 g, 10.8 mmol) was added to N, N-dimethylformamide (30 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoat

(**Intermediate A**) (2.0 g, 5.4 mmol) and potassium hydroxide (450 mg, 8.1 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours; the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a white solid of methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23B**) (346 mg, 13% yield).

[00559] LC-MS, M/Z (ESI): 490.2 [M+H]⁺

[00560] Second step: methyl 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23C**)



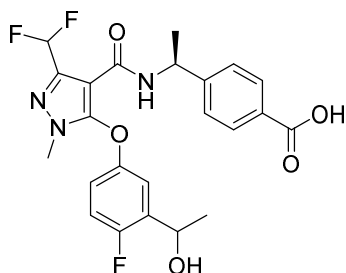
[00561] I-23C

[00562] The compound methyl (S)-4-(1-(5-(3-acetyl-4-fluorophenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23B**) (100 mg, 0.20 mmol)

was added to tetrahydrofuran (10 mL) at room temperature, cooled to 0°C; and sodium borohydride (23 mg, 0.60 mmol) was added. The mixture was stirred at room temperature for 3 h; the mixture was diluted by addition of water (100 mL), extracted with ethyl acetate (80 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of crude product of methyl 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23C**) (75 mg, yield 75%).

[00563] LC-MS, M/Z (ESI): 492.1 [M+H]⁺.

10 [00564] 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-23**)



[00565] **I-23**

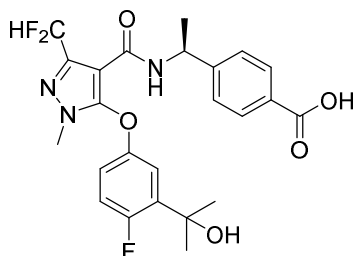
[00566] The starting material methyl 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-23C**) (75 mg, 0.15 mmol) was added to tetrahydrofuran (5 mL), methanol (5 mL) and water (1 mL) at room temperature; and lithium hydroxide monohydrate (20 mg, 0.47 mmol) was added. The mixture was stirred at room temperature for 5 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparative method A, a white solid of 4-((1S)-1-(3-(difluoromethyl)-5-(4-fluoro-3-(1-hydroxyethyl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-23**) (48 mg, 66% yield).

[00567] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.02 (d, 1H), 7.74 (d, 2H), 7.25 (t, 1H), 7.21-7.10 (m, 4 H), 6.90-6.81 (m, 1H), 4.96-4.86 (m, 2H), 3.72 (s, 3H), 3.67 (s, 1H), 1.28 (d, 3H), 1.24 (d, 3H).

25 [00568] LC-MS, M/Z (ESI): 478.1 [M+H]⁺

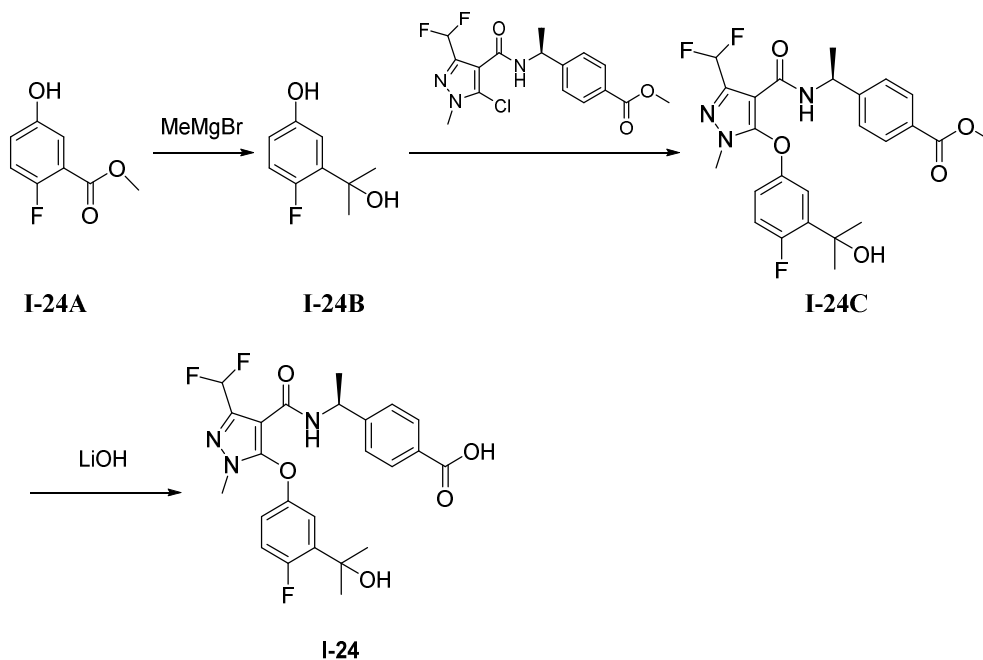
[00569] Example 24: Preparation of Compound I-24

[00570] (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-24**)



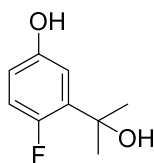
[00571] **I-24**

5 **[00572]** The synthesis scheme for **Compound I-24** is shown below:



[00573]

[00574] First step: 4-fluoro-3-(2-hydroxypropan-2-yl)phenol (**Compound I-24B**)

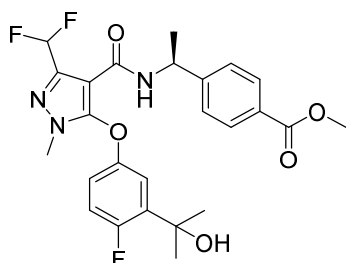


[00575] **I-24B**

10 **[00576]** A compound methyl 2-fluoro-5-hydroxybenzoate (340 mg, 2.0 mmol) was added to a solution of methyl magnesium bromide (1 mol/L) in tetrahydrofuran (5 mL) at room temperature and stirred for 16 hours; a saturated solution of ammonium chloride (10 mL) was added and extracted with dichloromethane (30 mL × 3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a

colorless liquid crude product of 4-fluoro-3-(2-hydroxypropan-2-yl)phenol (**Compound I-24B**) (340 mg, yield 100%).

[00577] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-24C**)

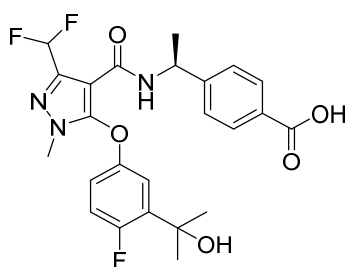


5 **[00578]** **I-24C**

[00579] The compound 4-fluoro-3-(2-hydroxypropan-2-yl)phenol (**Compound I-24B**) (340 mg, 2.0 mmol) was added to N, N-dimethylformamide (9 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (495 mg, 1.34 mmol), cuprous iodide (100 mg, 0.52 mmol), 1, 10-phenanthroline (180 mg, 1.0 mmol) and cesium carbonate (1.30 g, 4.05 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 2 hours. Then, the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, 15 filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-24C**) (120 mg, yield 32%).

[00580] LC-MS, M/Z (ESI): 506.3 [M+1]⁺

20 **[00581]** Third step: (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-24**)



[00582] I-24

[00583] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate

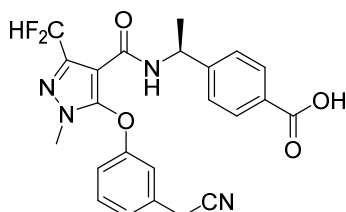
(**Compound I-24C**) (120 mg, 0.24 mmol) was added to tetrahydrofuran (5 mL), and water (1 mL) at room temperature; and lithium hydroxide (20 mg, 0.47 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(4-fluoro-3-(2-hydroxypropan-2-yl)phenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-24**) (22 mg, 19% yield).

[00584] ^1H NMR (400MHz, DMSO- d_6) δ 12.8 (s, 1H), 7.96 (d, 1H), 7.73 (d, 2H), 7.33 (t, 1H), 7.26 (t, 1H), 7.17 (d, 2H), 7.14 (t, 1H), 6.85 (t, 1H), 4.92 (t, 1H), 3.73 (s, 3H), 3.72 (s, 1H), 1.44 (s, 6H), 1.25 (d, 3H).

[00585] LC-MS, M/Z (ESI): 492.3 $[\text{M}+1]^+$

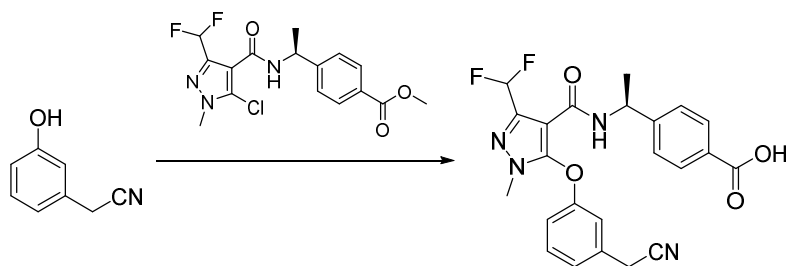
[00586] Example 25: Preparation of Compound I-25

[00587] (S)-4-(1-(5-(3-(cyanomethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-25**)



[00588] I-25

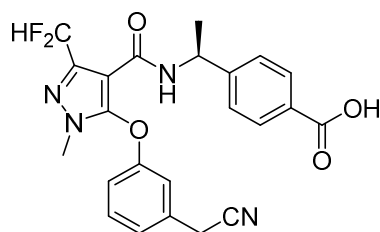
[00589] The synthesis scheme for **Compound I-25** is shown below:



[00590] I-25A

I-25

[00591] First step: (S)-4-(1-(5-(3-(cyanomethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-25**)



I-25

[00592]

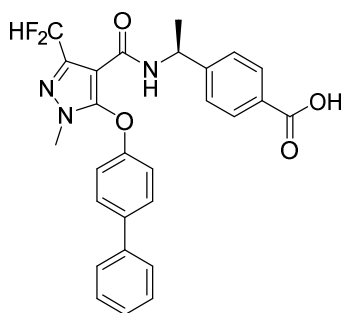
5 **[00593]** A compound 2-(3-hydroxyphenyl) acetonitrile (72 mg, 0.40 mmol) was added to N, N dimethylformamide (2 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (100 mg, 0.27 mmol) and potassium hydroxide (30 mg, 0.53 mmol) were added. The mixture was heated to 170°C in the microwave under nitrogen protection and stirred for 5 min; and then, the
10 mixture was cooled to room temperature and added with water (30 mL). The mixture was stirred for 0.5 hour; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-(3-(cyanomethyl)phenoxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (10 mg, yield 8%).

15 **[00594]** ¹H NMR (400MHz, CDCl₃) δ12.7 (s, 1H), 7.93 (d, 2H), 7.38 (t, 1H), 7.25 (t, 1H), 7.19 (d, 1H), 7.16 (d, 2H), 6.94 (s, 1H), 6.82 (t, 1H), 6.35 (d, 1H), 5.18 (t, 1H), 3.73 (s, 2H), 3.70 (s, 3H), 1.41 (d, 3H).

[00595] LC-MS, M/Z (ESI): 455.5 [M+H]⁺

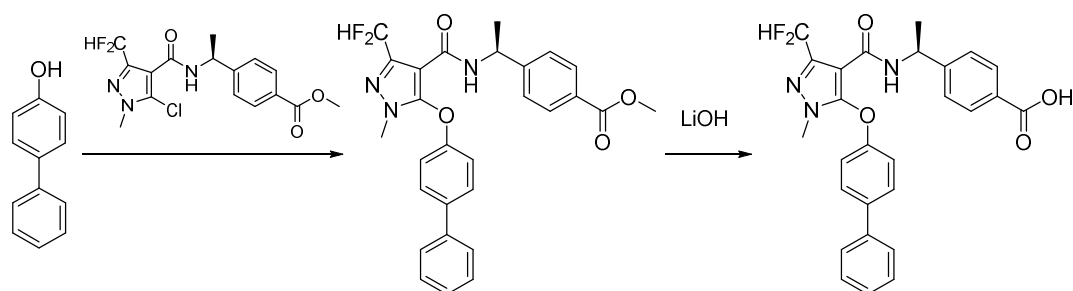
[00596] Example 26: Preparation of Compound I-26

20 **[00597]** (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-26**)



[00598] **I-26**

[00599] The synthesis scheme for **Compound I-26** is shown below:

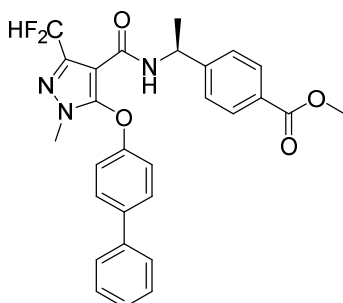


[00600] **I-26A**

I-26B

I-26

[00601] First step: methyl (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-26B**)



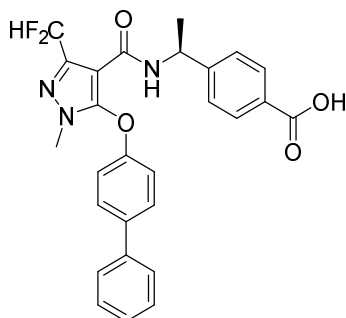
[00602] **I-26B**

[00603] A compound [1, 1'-biphenyl]-4-ol (458 mg, 2.70 mmol) was added to N, N-dimethylformamide (10 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (500 mg, 1.35 mmol) and potassium hydroxide (227 mg, 4.05 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 1 hour. Then, the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column

(petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a white solid crude product of methyl (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-26B**) (500 mg, yield 73%).

[00604] LC-MS, M/Z (ESI): 506.4 [M+H]⁺

- 5 **[00605]** Second step: (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-26**)



[00606] **I-26**

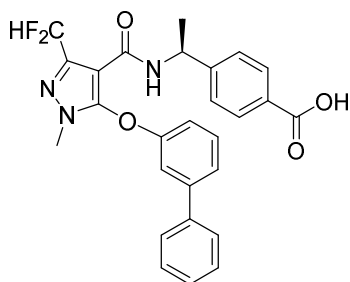
[00607] A starting material methyl (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-26B**) (450 mg, 0.89 mmol) was added to tetrahydrofuran (10 mL), methanol (5 mL) and water (4 mL) at room temperature; and lithium hydroxide (108 mg, 4.5 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-([1,1'-biphenyl]-4-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-26**) (70 mg, 16% yield).

[00608] ¹H NMR (400MHz, DMSO-d₆) δ 12.8 (s, 1H), 8.08 (d, 1H), 7.71 (d, 2H), 7.69 (d, 2H), 7.63 (d, 2H), 7.48 (t, 2H), 7.38 (t, 1H), 7.26 (t, 1H), 7.15 (d, 2H), 7.12 (d, 2H), 4.92 (t, 1H), 3.76 (s, 3H), 1.25 (d, 3H).

[00609] LC-MS, M/Z (ESI): 492.4 [M+H]⁺

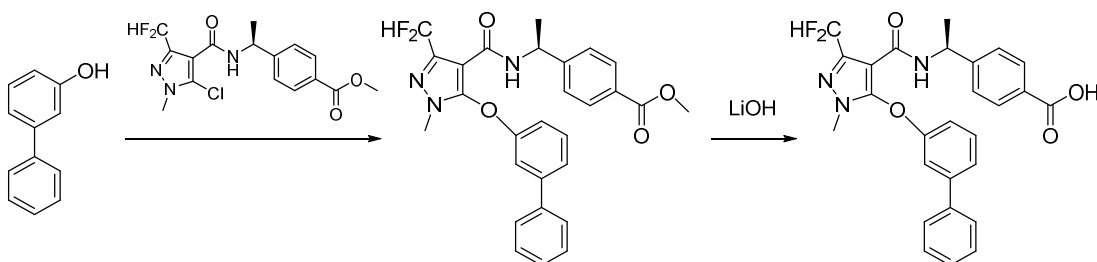
- 20 **[00610]** Example 27: Preparation of Compound I-27

[00611] (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-27**)



[00612] **I-27**

[00613] The synthesis scheme for **Compound I-27** is shown below:

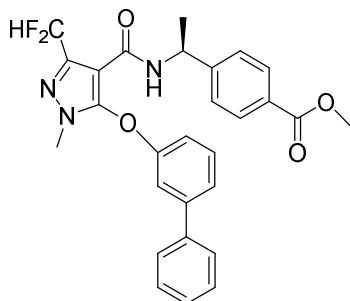


[00614] **I-27A**

I-27B

I-27

- 5 **[00615]** First step: methyl (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-27B**)



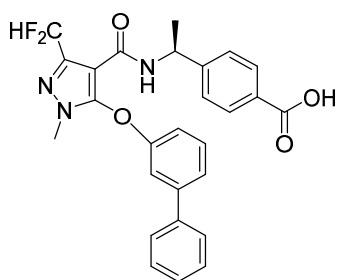
[00616] **I-27B**

- 10 **[00617]** A compound [1, 1'-biphenyl]-3-ol (458 mg, 2.70 mmol) was added to N, N-dimethylformamide (10 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (500 mg, 1.35 mmol) and potassium hydroxide (227 mg, 4.05 mmol) were added. The mixture was heated to 120°C in the microwave under nitrogen protection and stirred for 1 hour. Then, the mixture was cooled to room temperature and added with water (30 mL); the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL ×3)
- 15 and separated. The organic phases were combined, dried over anhydrous sodium sulfate,

filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a white solid crude product of methyl (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-27B**) (450 mg, yield 66%).

5 **[00618]** LC-MS, M/Z (ESI): 506.4 [M+H]⁺

[00619] Second step: (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-27**)



[00620] **I-27**

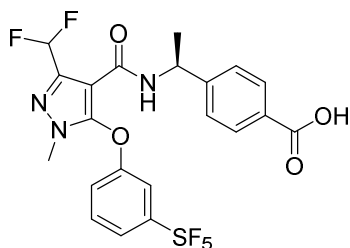
[00621] The starting material methyl (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-27B**) (400 mg, 0.79 mmol) was added to tetrahydrofuran (10 mL), methanol (5 mL) and water (5 mL) at room temperature; and lithium hydroxide (96 mg, 4.0 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(5-([1,1'-biphenyl]-3-yloxy)-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-27**) (115 mg, 30% yield).

[00622] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.07 (d, 1H), 7.68 (d, 2H), 7.62 (d, 2H), 7.50-7.44 (m, 4 H), 7.41 (d, 1H), 7.31 (s, 1H), 7.25-6.96 (m, 4 H), 4.89 (t, 1H), 3.77 (s, 3H), 1.19 (d, 3H).

20 **[00623]** LC-MS, M/Z (ESI): 492.4 [M+H]⁺

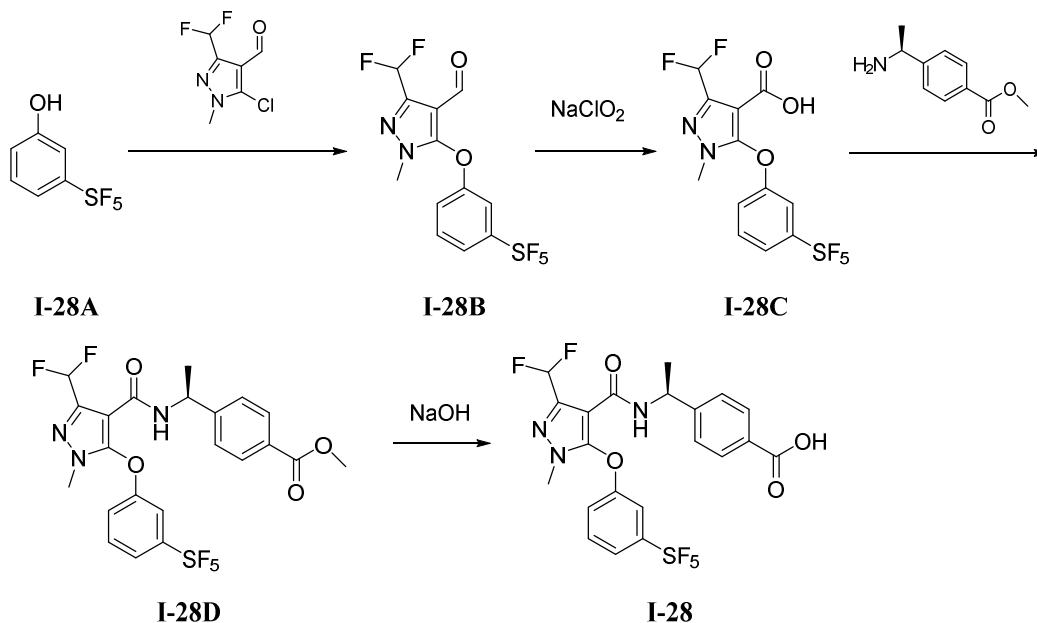
[00624] Example 28: Preparation of Compound I-28

[00625] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro-λ⁶-sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-28**)



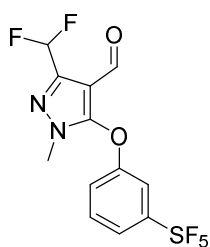
[00626] **I-28**

[00627] The synthesis scheme for **Compound I-28** is shown below:



[00628] **I-28D**

[00629] First step: 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carbaldehyde (**Compound I-28B**)



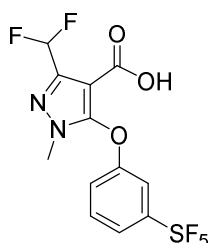
[00630] **I-28B**

[00631] A compound 5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (500 mg, 2.57 mmol) was added to N, N-dimethylformamide (5 mL) at room temperature; 3-(pentafluoro- λ^6 -sulfanyl) phenol (386 mg, 2.80 mmol) and potassium hydroxide (216 mg, 3.85 mmol) were added; and the mixture was heated to 150°C and stirred for 4 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), the pH was adjusted to 4

with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (15 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow liquid crude product of 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carbaldehyde (**Compound I-28B**) (750 mg, 98% yield).

[00632] LC-MS, M/Z (ESI): 379.1 [M+H]⁺

[00633] Second step: 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxylic acid (**Compound I-28C**)

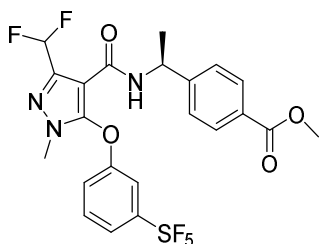


[00634] **I-28C**

[00635] The compound 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carbaldehyde (**Compound I-28B**) (750 mg, 2.55 mmol) was added to tert-butanol (6 mL) and water (7 mL) at room temperature; and 2-methyl-2-butene (355 mg, 5.07 mmol), sodium chlorite (456 mg, 5.07 mmol) and sodium dihydrogen phosphate (669 mg, 5.57 mmol) were added. The mixture was stirred for 14 hours at room temperature; the mixture was diluted with water (15 mL) and the pH was adjusted to 4 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a light yellow solid crude product of 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxylic acid (**Compound I-28C**) (800 mg, yield 100%).

[00636] LC-MS, M/Z (ESI): 395.1 [M+H]⁺

[00637] Third step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-28D**)

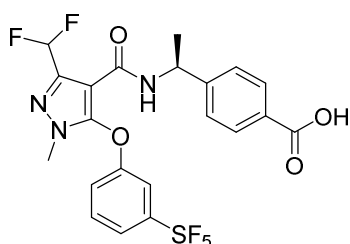


[00638] I-28D

[00639] The compound 3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxylic acid (compound **I-28C**) (800 mg, 2.58 mmol) was added to dichloromethane (20 mL); and methyl (S)-4-(1-aminoethyl) benzoate (459 mg, 2.56 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (1.40 g, 3.68 mmol) and N, N-diisopropylethylamine (991 mg, 7.68 mmol) were added. The mixture was stirred at room temperature for 16 hours, diluted with dichloromethane (40 mL), washed with water (20 mL $\times$ 3) and separated. The organic phases were dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a light yellow liquid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-28D**) (720 mg, yield 59%).

[00640] LC-MS, M/Z (ESI): 556.1 [M+H]⁺

[00641] Fourth step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-28**)



[00642] I-28

[00643] The starting material (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-28D**) (440 mg, 0.93 mmol) was added to methanol (10 mL) and water (1 mL) at room temperature; and sodium hydroxide (93 mg, 2.32 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was

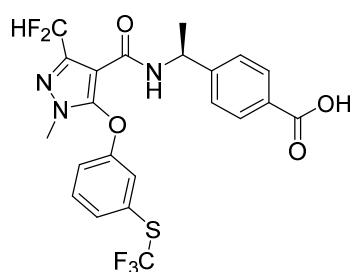
concentrated to obtain a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(pentafluoro- λ^6 -sulfanyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-28**) (93 mg, 22% yield).

[00644] ^1H NMR (400m Hz, DMSO- d_6) δ 12.7 (s, 1H), 8.18 (d, 1H), 7.74 (d, 2H), 7.73 (d, 1H), 7.71 (d, 1H), 7.65 (t, 1H), 7.26 (d, 1H), 7.23 (t, 1H), 7.21 (d, 2H), 4.89 (t, 1H), 3.78 (s, 3H), 1.11 (d, 3H).

[00645] LC-MS, M/Z (ESI): 542.1 $[\text{M}+\text{H}]^+$

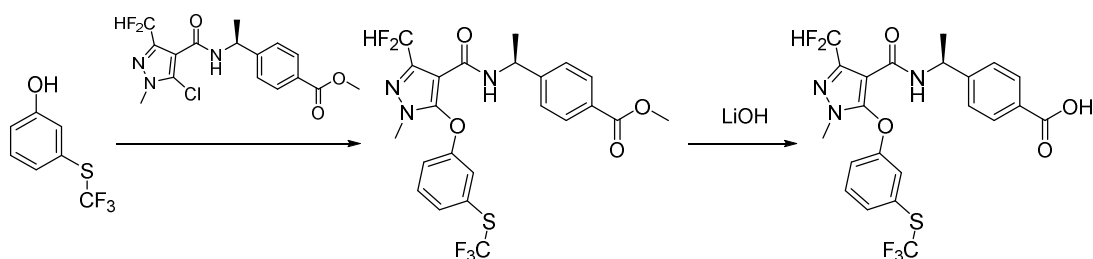
[00646] Example 29: Preparation of Compound I-29

[00647] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-29**)



[00648] **I-29**

[00649] The synthesis scheme for **Compound I-29** is shown below:

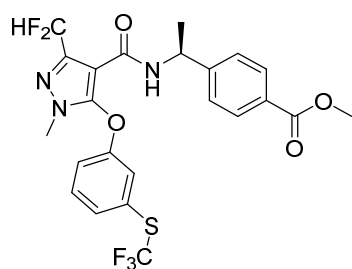


[00650] **I-29A**

I-29B

I-29

[00651] First step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-29B**)

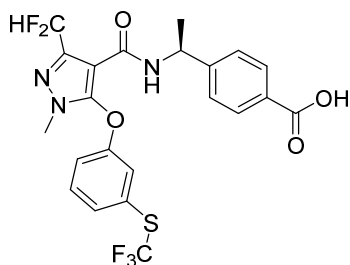


[00652] **I-29B**

[00653] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (186 mg, 0.50 mmol) was added to N, N-dimethylformamide (5 mL) at room temperature; 3-((trifluoromethyl) sulfanyl) phenol (120 mg, 0.60 mmol) and potassium hydroxide (45 mg, 0.80 mmol) were added; and the mixture was heated to 120°C and stirred for 12 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (15 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-29B**) (20 mg, yield 7.5%).

[00654] LC-MS, M/Z (ESI): 530.2 [M+H]⁺

[00655] Second step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-29**)



[00656] **I-29**

[00657] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (compound **I-29B**) (20 mg, 0.04 mmol) was added to tetrahydrofuran (2 mL), methanol (2 mL) and water (1 mL) at room temperature; lithium hydroxide (3 mg, 0.16 mmol) was added; and the mixture was heated to 50°C and stirred for 4 hours. Then, the mixture was cooled to room temperature and the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparative method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-((trifluoromethyl)thio)phenoxy)-1H-pyrazole-4-

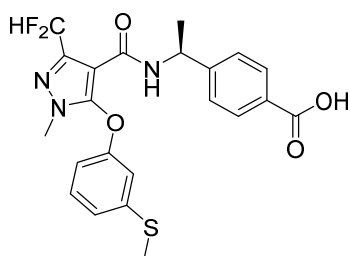
carboxamido)ethyl)benzoic acid (**Compound I-29**) (14 mg, 72% yield).

[00658] ^1H NMR (400 MHz, DMSO- d_6) δ 12.8 (s, 1H), 8.16 (d, 1H), 7.74 (d, 2H), 7.57 (t, 2H), 7.39 (s, 1H), 7.22-6.69 (m, 4 H), 4.91-4.84 (m, 1H), 3.74 (s, 3H), 1.21 (d, 3H).

[00659] LC-MS, M/Z (ESI): 516.2 $[\text{M}+\text{H}]^+$

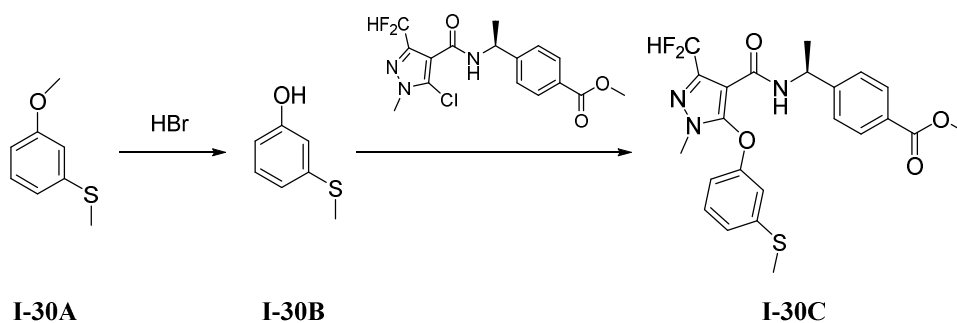
5 **[00660]** Example 30: Preparation of Compound I-30

[00661] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-30**)



[00662] **I-30**

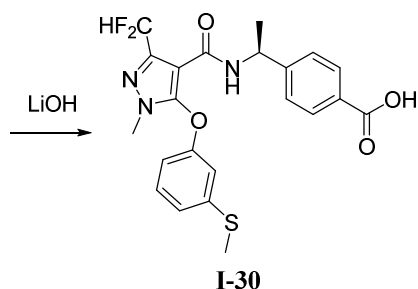
[00663] The synthesis scheme for **Compound I-30** is shown below:



I-30A

I-30B

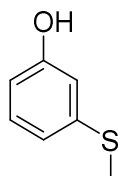
I-30C



I-30

10 **[00664]**

[00665] First step: 3-(methylthio)phenol (**Compound I-30B**)

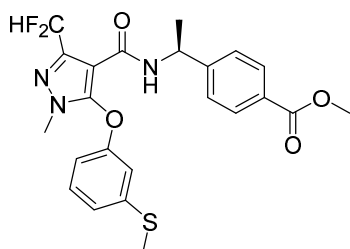


[00666] **I-30B**

[00667] A compound (3-methoxyphenyl) (methyl) sulfide (4.0 g, 26 mmol) was added to a

solution of 30% hydrobromic acid in acetic acid (12 mL) and a solution of 48% hydrobromic acid in water (3 mL) at room temperature; the mixture was heated to reflux under N₂ protection and stirred for 6 hours; and then the mixture was cooled to room temperature, diluted with water (100 mL), extracted with diethyl ether (100 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered, and concentrate to obtain a colorless liquid crude product of 3-(methylthio)phenol (**Compound I-30B**) (3.8 g, yield 100%).

[00668] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-30C**)

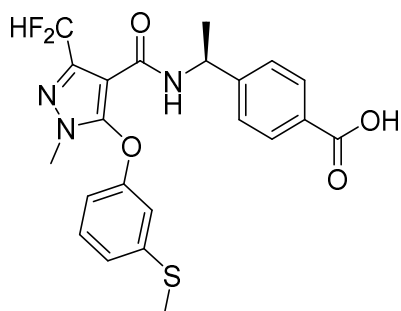


[00669] I-30C

[00670] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (700 mg, 1.89 mmol) was added to N, N-dimethylformamide (5 mL) at room temperature; and 3-(methylthio) phenol (528 mg, 3.77 mmol) and potassium hydroxide (318 mg, 5.67 mmol) were added. The mixture was heated to 150°C in the microwave and stirred for 2 hours. Then, the mixture was cooled to room temperature and diluted with water (20 mL), the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-30C**) (800 mg, 89% yield).

[00671] LC-MS, M/Z (ESI): 476.5 [M+H]⁺

[00672] Third step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-30**)



[00673] I-30

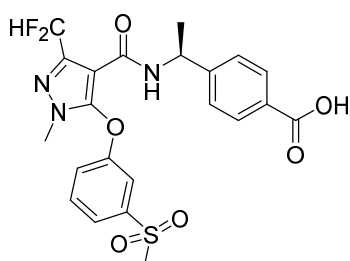
[00674] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-30C**) (800 mg, 1.68 mmol) was added to tetrahydrofuran (10 mL), methanol (10 mL) and water (2 mL) at room temperature; lithium hydroxide (121 mg, 5.0 mmol) was added; and the mixture was heated to 50°C and stirred for 4 hours. Then, the mixture was cooled to room temperature and the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-30**) (165 mg, yield 21%).

[00675] ¹H NMR (400MHz, DMSO-d₆) δ 12.8 (s, 1H), 8.03 (d, 1H), 7.74 (d, 2H), 7.34 (t, 2H), 7.14 (d, 2H), 7.11 (t, 1H), 6.97 (t, 1H), 6.70 (d, 1H), 4.91 (t, 1H), 3.73 (s, 3H), 2.45 (s, 3H), 1.24 (d, 3H).

[00676] LC-MS, M/Z (ESI): 462.5 [M+H]⁺

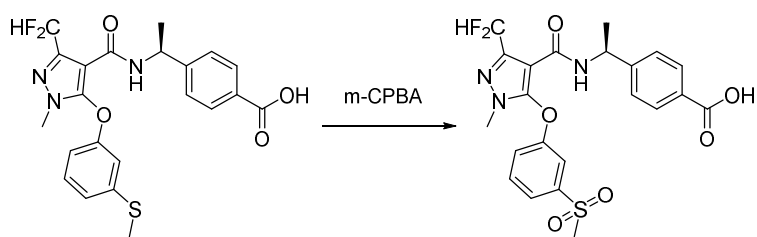
[00677] Example 31: Preparation of Compound I-31

[00678] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylsulfonyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-31**)



[00679] I-31

[00680] The synthesis scheme for **Compound I-31** is shown below:



[00681]

I-30

I-31

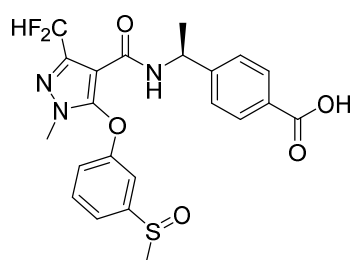
[00682] The starting material (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-30**) (80 mg, 0.17 mmol) was added to dichloromethane (5 mL); and m-chloroperoxybenzoic acid (83 mg, 0.48 mmol) was added at room temperature and stirred for 3 hours at room temperature. The reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylsulfonyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-31**) (64 mg, 75% yield).

[00683] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.21 (d, 1H), 7.77 (d, 2H), 7.75 (d, 1H), 7.70 (t, 1H), 7.59 (t, 1H), 7.37 (d, 1H), 7.25 (t, 1H), 7.15 (d, 2H), 4.87 (t, 1H), 3.77 (s, 3H), 3.19 (s, 3H), 1.19 (d, 3H).

[00684] LC-MS, M/Z (ESI): 494.1 [M+H]⁺

[00685] Example 32: Preparation of Compound I-32

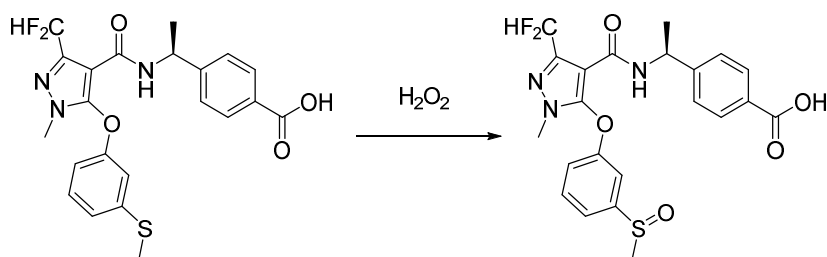
[00686] 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(methylsulfinyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-32**)



[00687]

I-32

[00688] The synthesis scheme for **Compound I-32** is shown below:



[00689]

I-30

I-32

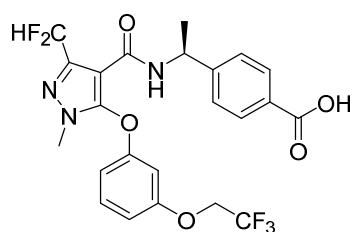
[00690] The starting material (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(methylthio)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-30**) (60 mg, 0.13 mmol) was added to tetrahydrofuran (5 mL); 30% hydrogen peroxide (1 mL) was added; and the mixture was heated to 60°C and stirred for 7 h at room temperature. Then, the mixture was cooled to room temperature and was concentrated to prepare, by the acidic preparation method A, a white solid of 4-((1S)-1-(3-(difluoromethyl)-1-methyl-5-(3-(methylsulfinyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-32**) (35 mg, yield 75%).

[00691] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.15 (d, 1H), 7.73 (d, 2H), 7.61 (d, 1H), 7.50 (t, 1H), 7.36 (s, 1H), 7.25 (t, 1H), 7.15 (d, 1H), 7.13 (d, 2H), 4.87 (t, 1H), 3.75 (s, 3H), 2.69 (s, 3H), 1.20 (d, 3H).

[00692] LC-MS, M/Z (ESI): 478.2 [M+H]⁺

[00693] Example 33: Preparation of **Compound I-33**

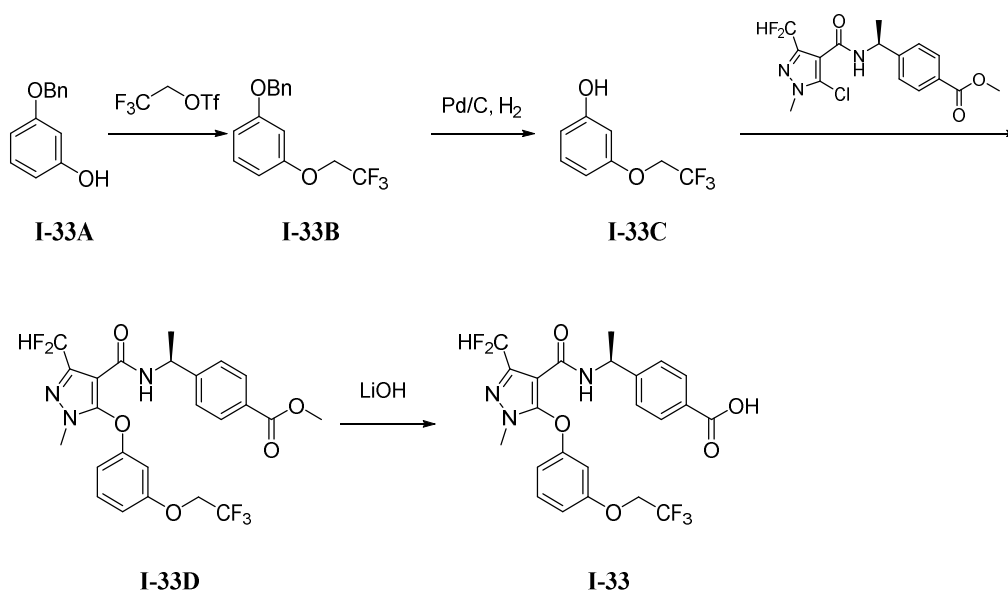
[00694] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-33**)



[00695]

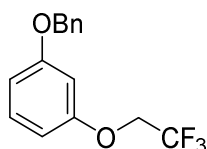
I-33

[00696] The synthesis scheme for **Compound I-33** is shown below:



[00697]

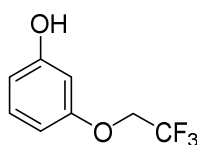
[00698] First step: 1-(benzyloxy)-3-(2,2,2-trifluoroethoxy)benzene (**Compound I-33B**)



[00699] **I-33B**

[00700] A compound 3-(benzyloxy) phenol (600 mg, 3.0 mmol) was added to N, N-dimethylformamide (10 mL) at room temperature; trifluoro-2, 2, 2-trifluoroethane-1-sulfonate (840 mg, 3.6 mmol) and potassium carbonate (1.24 g, 9.0 mmol) were added; and the mixture was heated to 60°C and stirred for 16 hours. Then, the mixture was cooled to room temperature, diluted with water (30 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered, and concentrated; the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a colorless liquid of 1-(benzyloxy)-3-(2,2,2-trifluoroethoxy)benzene (**Compound I-33B**) (380 mg, yield 45%).

[00701] Second step: 3-(2,2,2-trifluoroethoxy)phenol (**Compound I-33C**)

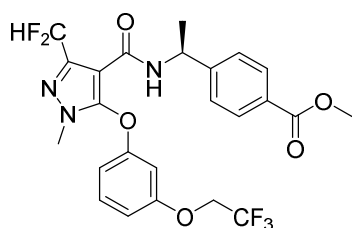


[00702] **I-33C**

[00703] The compound 1-(benzyloxy)-3-(2,2,2-trifluoroethoxy)benzene (**Compound I-33B**)

(520 mg, 1.84 mmol) was added to ethanol (10 mL) at room temperature; 10% palladium on carbon (50 mg) was added; and a drop of formic acid was added. Then, hydrogen was introduced, and the mixture was stirred for 12 hours; and the mixture was diluted with water (20 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of the captioned compound, i.e., 3-(2,2,2-trifluoroethoxy)phenol (**Compound I-33C**) (210 mg, yield 59%).

[00704] Third step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-33D**)



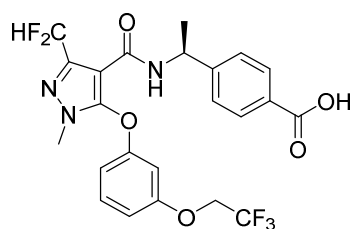
[00705] **I-33D**

[00706] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (371 mg, 1.0 mmol) was added to N, N-dimethylformamide (5 mL) at room temperature; 3-(2,2,2-trifluoroethoxy)phenol (**Compound I-33C**) (210 mg, 1.1 mmol), copper iodide (191 mg, 1.0 mmol), cesium carbonate (1.0 g, 3.1 mmol), and 1, 10-phenanthroline (72 mg, 0.40 mmol) were added; and the mixture was heated to 120°C in the microwave and stirred for 2 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-33D**) (120 mg, yield 8%).

[00707] LC-MS, M/Z (ESI): 528.2 [M+H]⁺

[00708] Fourth step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-

trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-33**)



[00709] **I-33**

[00710] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-33D**)

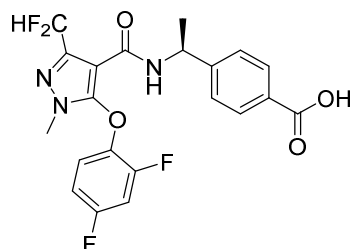
5 (120 mg, 0.23 mmol) was added to tetrahydrofuran (5 mL), methanol (5 mL) and water (1 mL) at room temperature; and lithium hydroxide (20 mg, 0.83 mmol) was added. The mixture was stirred for 16 hours at room temperature; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(2,2,2-trifluoroethoxy)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-33**) (45 mg, 72% yield).

[00711] ¹H NMR (400MHz, DMSO-d₆) δ12.8 (s, 1H), 8.02 (d, 1H), 7.74 (d, 2H), 7.38 (t, 1H), 7.16 (t, 1H), 7.14 (d, 2H), 6.93 (d, 1H), 6.76 (t, 1H), 6.65 (d, 1H), 4.92 (t, 1H), 4.78 (dd, 2H), 3.72 (s, 3H), 1.25 (d, 3H).

[00712] LC-MS, M/Z (ESI): 514.2 [M+H]⁺

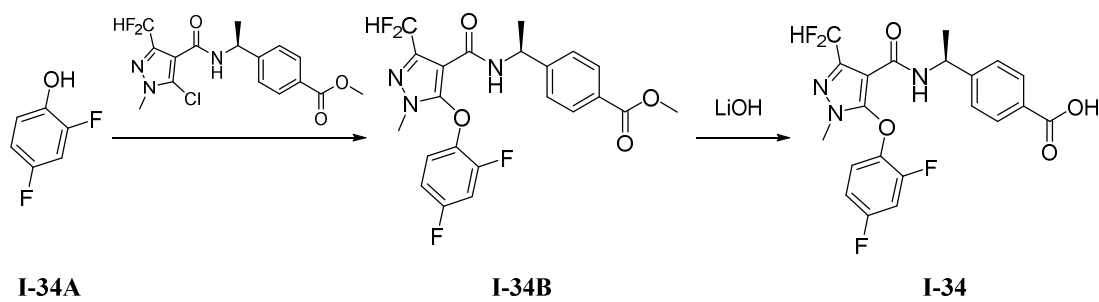
15 **[00713]** Example 34: Preparation of **Compound I-34**

[00714] (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-34**)

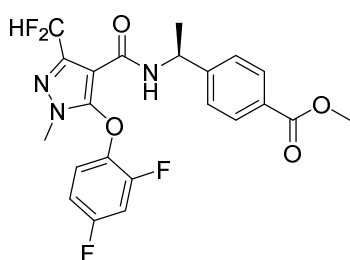


[00715] **I-34**

[00716] The synthesis scheme for **Compound I-34** is shown below:



[00718] First step: methyl (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-34B**)

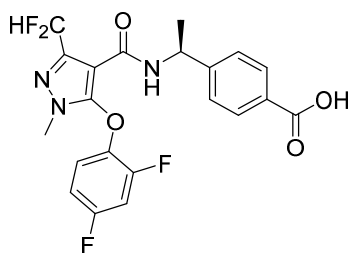


5 **[00719] I-34B**

[00720] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (370 mg, 1.0 mmol) was added to N, N-dimethylformamide (10 mL) at room temperature; 2, 4-difluorophenol (195 mg, 1.5 mmol) and potassium hydroxide (168 mg, 3.0 mmol) were added; and the mixture was heated to 120°C and stirred for 6 hours. Then, the mixture was cooled to room temperature and diluted with water (50 mL), the pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (60 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of methyl (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-34B**) (160 mg, yield 34%).

[00721] LC-MS, M/Z (ESI): 466.1 [M+H]⁺

[00722] Second step: (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-34**)



[00723] I-34

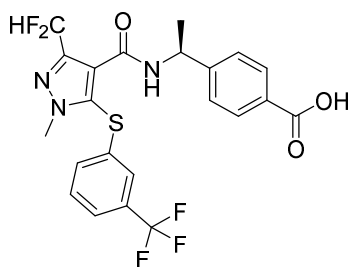
[00724] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-34B**) (160 mg, 0.34 mmol) was added to tetrahydrofuran (4 mL) and water (2 mL) at room temperature; lithium hydroxide (32 mg, 0.76 mmol) was added; and the mixture was heated to 50°C and stirred for 2 hours. Then, the mixture was cooled to room temperature and the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(2,4-difluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-34**) (11 mg, yield 7.0%).

[00725] ¹H NMR (400mHz, DMSO-d₆) δ12.7 (s, 1H), 8.28 (d, 1H), 7.78 (d, 2H), 7.45 (t, 1H), 7.21 (d, 2H), 7.17 (t, 1H), 7.04 (d, 1H), 7.02 (d, 1H), 4.88 (t, 1H), 3.76 (s, 3H), 1.26 (d, 3H).

[00726] LC-MS, M/Z (ESI): 452.1 [M+H]⁺

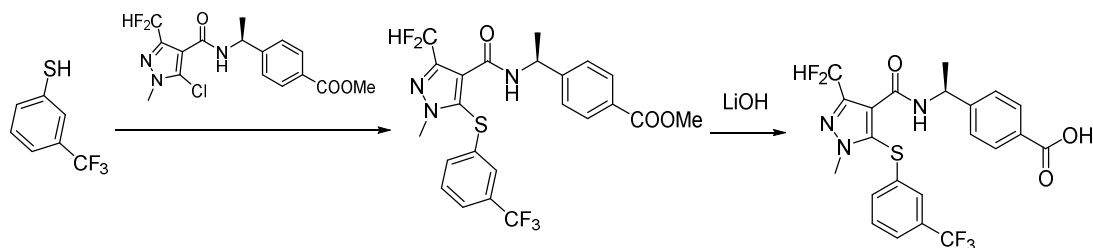
[00727] Example 35: Preparation of **Compound I-35**

[00728] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-



[00729] I-35

[00730] The synthesis scheme for **Compound I-35** is shown below:

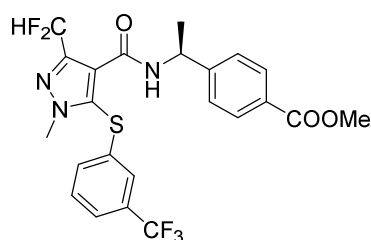


[00731] I-35A

I-35B

I-35

[00732] First step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-35B**)

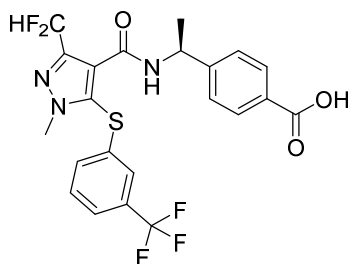


[00733] I-35B

5 **[00734]** A compound 3-(trifluoromethyl) benzenethiol (215 mg, 1.20 mmol) was added to N, N-dimethylformamide (20 mL) at room temperature; and sodium hydride (80 mg, 2.00 mmol) was added. The mixture was stirred for 0.5 hour; methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (375 mg, 1.00 mmol) was added; and the mixture was heated to 120°C and stirred for 8 h. Then, the mixture was
10 cooled to room temperature, diluted with water (100 mL), extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; the residue purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-carboxamido)ethyl)benzoate
15 (**Compound I-35B**) (350 mg, yield 67%).

[00735] LC-MS, M/Z (ESI): 514.2 (M+1).

[00736] Second step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-35**)



[00737] I-35

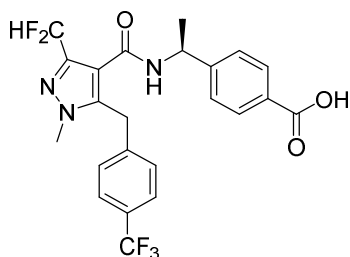
[00738] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-35B**) (220 mg, 0.43 mmol) was added to tetrahydrofuran (5 mL) at room temperature; water (5 mL),
 5 methanol (5 mL), and lithium hydroxide (30 mg, 1.25 mmol) were added; and the mixture was heated to 50°C and stirred for 18 hours. The reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((3-(trifluoromethyl)phenyl)thio)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-35**) (105 mg, 49% yield).

10 **[00739]** ¹H NMR (400m Hz, DMSO-d₆) δ12.8 (s, 1H), 8.68 (d, 1H), 7.79 (d, 2H), 7.63 (d, 1H), 7.58 (t, 2H), 7.39 (d, 2H), 7.27 (t, 1H), 7.14 (t, 1H), 5.10-5.03 (m, 1H), 3.89 (s, 3H), 1.38 (d, 3H).

[00740] LC-MS, M/Z (ESI): 500.2 (M+1)

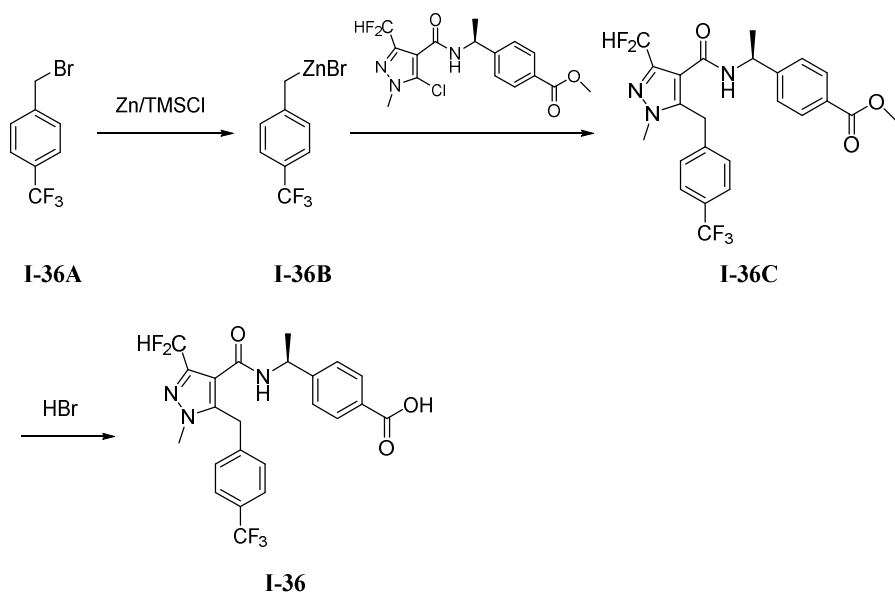
[00741] Example 36: Preparation of **Compound I-36**

15 **[00742]** (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-36**)



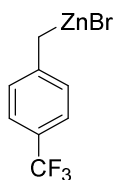
[00743] I-36

[00744] The synthesis scheme for **Compound I-36** is shown below:



[00745]

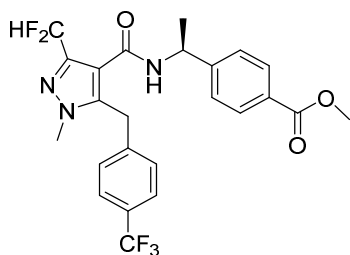
[00746] First step: (4-(trifluoromethyl)benzyl)zinc(II) bromide (**Compound I-36B**)



[00747] **I-36B**

[00748] Zinc powder (2.60 g, 40.0 mmol) was added to anhydrous tetrahydrofuran (15 mL) at room temperature; 1, 2-dibromoethane (0.02 mL) was added under nitrogen protection; and the mixture was heated to 60°C. Then, chlorotrimethylsilane (0.02 mL) was added, and the mixture was stirred for 15 min and cooled to 0°C. A solution of 4-(trifluoromethyl) benzyl bromide (4.80 g, 20.0 mmol) in tetrahydrofuran (5 mL) was added dropwise; and then the mixture was heated to 60°C and stirred for 1 hour. The mixture was cooled to room temperature to obtain a solution (1 mol/L) of (4-(trifluoromethyl)benzyl)zinc(II) bromide (**Compound I-36B**) in tetrahydrofuran (20 mL).

[00749] Second step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-36C**)

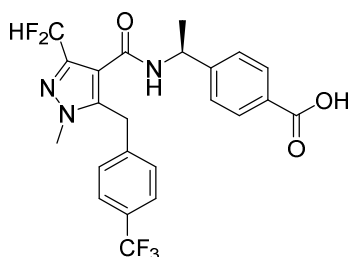


[00750] I-36C

[00751] The compound (4-(trifluoromethyl) benzyl) zinc bromide in tetrahydrofuran (**Compound I-36B**) (2.70 mL, 2.70 mmol) was added to N, N-dimethylformamide (15 mL) at room temperature; and methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (500 mg, 1.35 mmol), 4, 4'-di-tert-butyl-2, 2'-bipyridine (35 mg, 0.13 mmol) and nickel chloride glyme complex (30 mg, 0.13 mmol) were added. The mixture was heated to 100°C in the microwave under nitrogen protection and stirred for 1 hour; the mixture was cooled to room temperature and added with water (50 mL); and the mixture was extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-36C**) (180 mg, yield 27%).

[00752] LC-MS, M/Z (ESI): 496.3 [M+1]⁺.

[00753] Third step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-36**)



[00754] I-36

[00755] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-36C**) (150 mg, 0.30 mmol) was added to 48% hydrobrominated acetic acid solution (3 mL) at room

temperature, and the mixture was heated to 65°C and stirred for 16 hours. The mixture was then cooled to room temperature, and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(4-(trifluoromethyl)benzyl)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-36**)

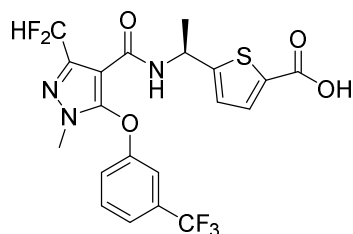
5 (90 mg, yield 62%).

[00756] ¹H NMR (400mHz, DMSO-d₆) δ12.8 (s, 1H), 8.47 (d, 1H), 7.86 (d, 2H), 7.62 (d, 2H), 7.41 (d, 2H), 7.33 (d, 2H), 7.23 (t, 1H), 5.12 (t, 1H), 4.37 (dd, 2H), 3.77 (s, 3H), 1.41 (s, 3H).

[00757] LC-MS, M/Z (ESI): 482.3 [M+1]⁺

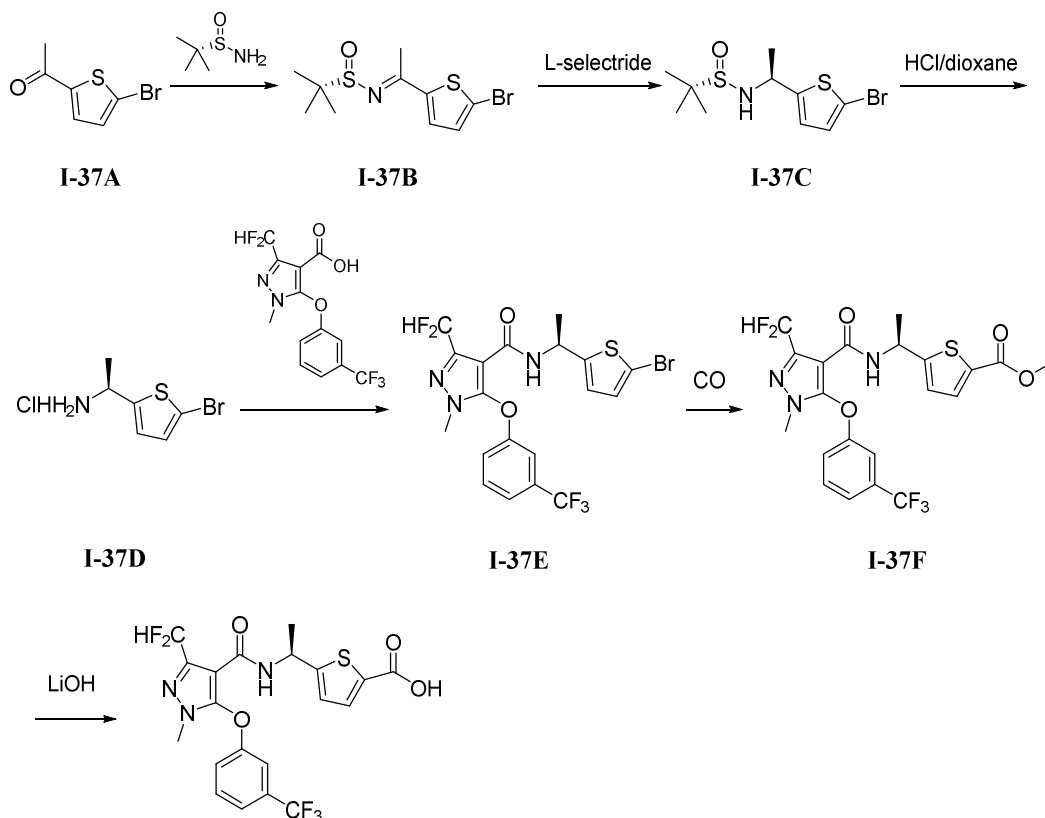
[00758] Example 37: Preparation of Compound I-37

10 **[00759]** (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylic acid (**Compound I-37**)



[00760] **I-37**

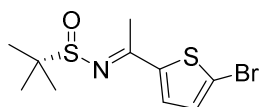
[00761] The synthesis scheme for **Compound I-37** is shown below:



[00762]

I-37

[00763] First step: (R, E)-N-(1-(5-bromothiophen-2-yl)ethylidene)-2-methylpropane-2-sulfinamide (**Compound I-37B**)



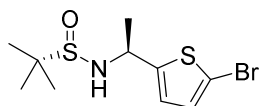
[00764]

I-37B

5 [00765] A compound 1-(5-bromothiophen-2-yl) ethan-1-one (2.05 g, 10.0 mmol) was added to dichloromethane (50 mL) at room temperature; (R)-tert-butylsulfinyl (1.8 g, 15.0 mmol) and tetraethyl titanate (5.70 g, 25.0 mmol) were added; and the mixture was heated to 50°C, stirred for 20 hours, and added with water (80 mL). The mixture was extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain a white solid of (R, E)-N-(1-(5-bromothiophen-2-yl)ethylidene)-2-methylpropane-2-sulfinamide (**Compound I-37B**) (500 mg, yield 16%).

[00766] Second step: (R)-N-((S)-1-(5-bromothiophen-2-yl)ethyl)-2-methylpropane-2-

sulfinamide (**Compound I-37C**)

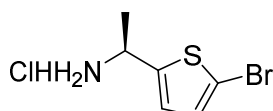


[00767] I-37C

[00768] The compound (R, E)-N-((S)-1-(5-bromothiophen-2-yl)ethylidene)-2-methylpropane-2-sulfinamide (**Compound I-37B**) (500 mg, 1.62 mmol) was added to tetrahydrofuran (10 mL) at room temperature; the mixture was cooled to -78°C, and added with 1 mol/L solution of lithium tri-sec-butylborohydride in tetrahydrofuran (2.4 mL, 2.4 mmol); and then the mixture was warmed naturally to room temperature and stirred for 16 hours. The pH was adjusted to 4 with 1N hydrochloric acid, and the mixture was diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid of (R)-N-((S)-1-(5-bromothiophen-2-yl)ethyl)-2-methylpropane-2-sulfinamide (**Compound I-37C**) (320 mg, yield 64%).

[00769] LC-MS, M/Z (ESI): 310.2 [M+H]⁺.

[00770] Third step: (S)-1-(5-bromothiophen-2-yl)ethan-1-amine hydrochloride (**Compound I-37D**)

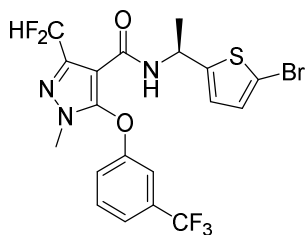


[00771] I-37D

[00772] The starting material (R)-N-((S)-1-(5-bromothiophen-2-yl)ethyl)-2-methylpropane-2-sulfinamide (**Compound I-37C**) (310 mg, 1.0 mmol) was added to methanol (10 mL) at room temperature, and 4 mol/L solution of dioxane hydrochloride (0.5 mL, 2.0 mmol) was added. The mixture was stirred at room temperature for 16 hours; the reaction mixture was concentrated and diethyl ether (10 mL) was added; the mixture was filtered to obtain (S)-1-(5-bromothiophen-2-yl)ethan-1-amine hydrochloride (60097D) (190 mg, yield 78%) as a white solid.

[00773] Fourth step: (S)-N-((1-(5-bromothiophen-2-yl)ethyl)-3-(difluoromethyl)-1-methyl-5-

(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamide (**Compound I-37E**)

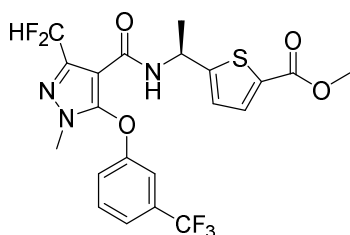


[00774] I-37E

[00775] A compound 3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl) phenoxy)-1H-pyrazole-4-carboxylic acid (140 mg, 0.41 mmol) was added to dichloromethane (5 mL) and N, N-dimethylformamide (5 mL) at room temperature; (S)-1-(5-bromothiophen-2-yl)ethan-1-amine hydrochloride (**Compound I-37D**) (118 mg, 0.49 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N-tetramethyluronium hexafluorophosphate (239 mg, 0.63 mmol) and N, N-diisopropylethylamine (135 mg, 1.05 mmol) were added. The mixture was stirred for 16 hours at room temperature. Then, the mixture was diluted with water (50 mL), extracted with ethyl acetate (50 mL $\times$ 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; and the residue was purified by thin-layer silica gel separation (petroleum ether: ethyl acetate (V/V) = 4: 1) to obtain a white solid crude product of (S)-N-(1-(5-bromothiophen-2-yl)ethyl)-3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamide (**Compound I-37E**) (170 mg, yield 78%).

[00776] LC-MS, M/Z (ESI): 524.2 [M+H]⁺

[00777] Fifth step: methyl (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylate (**Compound I-37F**)



[00778] I-37F

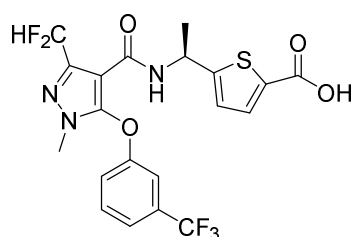
[00779] A compound (S)-N-(1-(5-bromothiophen-2-yl)ethyl)-3-(difluoromethyl)-1-methyl-5-

(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamide (**Compound I-37E**) (170 mg, 0.32 mmol) was added to methanol (5 mL) at room temperature; triethylamine (320 mg, 3.20 mmol) and [1, 1'-bis(diphenylphosphino) ferrocene] dichloropalladium (53 mg, 0.06 mmol) were added; carbon monoxide was introduced; and the mixture was heated to 120°C and stirred for 48 h.

Then, mixture was diluted with water (20 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a white solid of methyl (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylate (**Compound I-37F**) (140 mg, yield 86%).

[00780] LC-MS, M/Z (ESI): 504.5 [M+H]⁺

[00781] Sixth step: (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylic acid (**Compound I-37**)



[00782] **I-37**

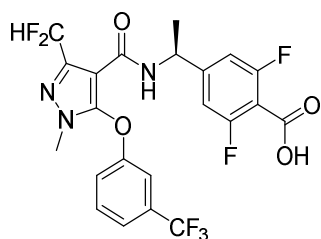
The starting material methyl (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylate (**Compound I-37F**) (140 mg, 0.28 mmol) was added to tetrahydrofuran (5 mL) and water (5 mL) at room temperature, and lithium hydroxide (20 mg, 0.47 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, (S)-5-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)thiophene-2-carboxylic acid (**Compound I-37**) (110 mg, yield 81%) as a white solid.

[00784] ¹H NMR (400MHz, DMSO-d₆) δ12.7 (s, 1H), 8.32 (d, 1H), 7.62 (t, 1H), 7.54 (t, 1H), 7.42 (d, 2H), 7.28 (t, 1H), 7.23 (t, 1H), 6.75 (d, 1H), 5.51-5.04 (m, 1H), 3.76 (s, 3H), 1.27 (d, 3H).

[00785] LC-MS, M/Z (ESI): 490.5 [M+H]⁺

[00786] Example 38: Preparation of Compound I-38

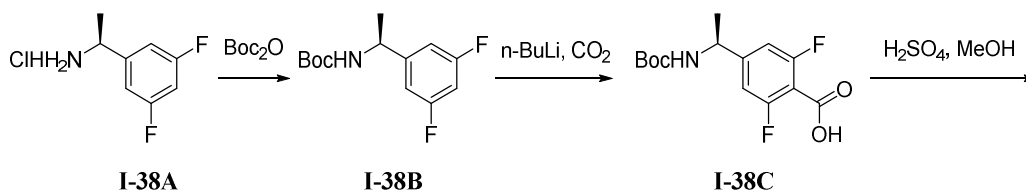
[00787] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoic acid (**Compound I-38**)



I-38

5 [00788]

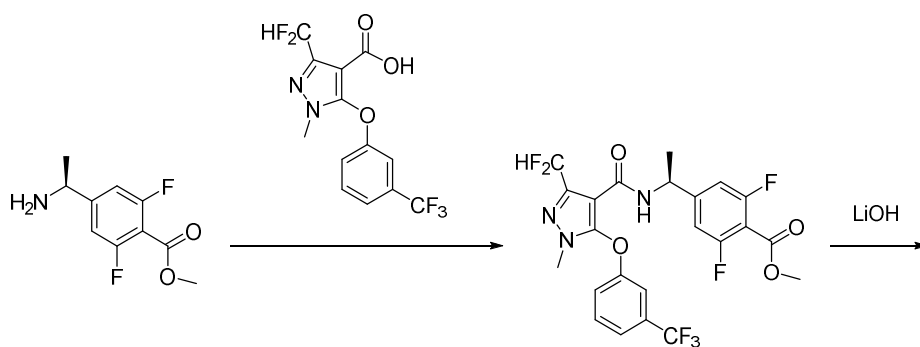
[00789] The synthesis scheme for **Compound I-38** is shown below:



I-38A

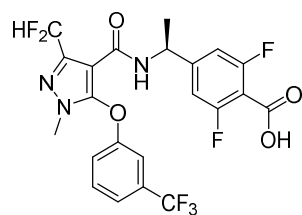
I-38B

I-38C



I-38D

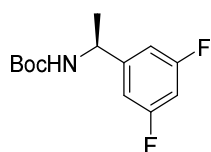
I-38E



I-38

[00790]

[00791] First step: tert-butyl (S)-1-(3,5-difluorophenyl)ethylcarbamate (**Compound I-38B**)

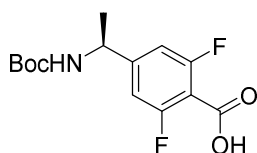


I-38B

[00792]

[00793] A compound (S)-1-(3, 5-difluorophenyl) ethan-1-aminohydrochloride (2.0 g, 10.3 mmol) was added to dichloromethane (50 mL) at room temperature; di-tert-butyl dicarbonate (4.5 g, 20.7 mmol) and triethylamine (3.1 g, 31.0 mmol) were added. The mixture was stirred for 16 hours at room temperature; then the reaction mixture was concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 10: 1) to obtain tert-butyl (S)-1-(3,5-difluorophenyl)ethyl)carbamate (**Compound I-38B**) (2.2 g, 72% yield) as a white solid.

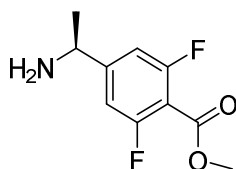
[00794] Second step: (S)-4-(1-((tert-butoxycarbonyl)amino)ethyl)-2,6-difluorobenzoic acid (**Compound I-38C**)



I-38C

[00796] A compound (S)-tert-butyl (1-(3, 5-difluorophenyl)ethyl)carbamate (500 mg, 2.50 mmol) was added to tetrahydrofuran (10 mL) at room temperature. The mixture was cooled to -78°C, and added with 2.5 mol/L solution of butyllithium in THF (2.0 mL, 5.0 mmol). The mixture was stirred for 2 hours; carbon dioxide was introduced at low temperature. The mixture was warmed up naturally to room temperature and stirred for 16 hours; the pH was adjusted to 2 with 1N hydrochloric acid; and the mixture was diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of (S)-4-(1-((tert-butoxycarbonyl)amino)ethyl)-2,6-difluorobenzoic acid (**Compound I-38C**) (550 mg, yield 90%).

[00797] Third step: methyl (S)-4-(1-aminoethyl)-2,6-difluorobenzoate (**Compound I-38D**)

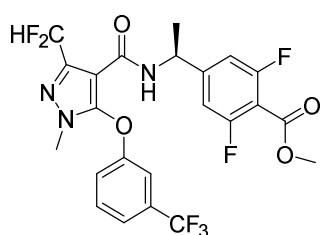


I-38D

[00799] The starting material (S)-4-(1-((tert-butoxycarbonyl)amino)ethyl)-2,6-difluorobenzoic acid (**Compound I-38C**) (500 mg, 2.0 mmol) was added to methanol (100 mL)

at room temperature and added with concentrated sulfuric acid (20 mL). The mixture was heated to 80°C and stirred for 6 hours; and then the mixture was diluted by addition of water (100 mL), extracted with ethyl acetate (100 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of methyl (S)-4-(1-aminoethyl)-2,6-difluorobenzoate (**Compound I-38D**) (1.2 g, yield 100%).

[00800] Fourth step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoate (**Compound I-38E**)

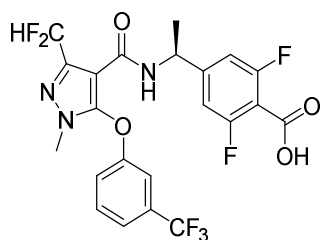


[00801] **I-38E**

[00802] A compound 3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl) phenoxy)-1H-pyrazole-4-carboxylic acid (100 mg, 0.30 mmol) was added to dichloromethane (8 mL) and N, N-dimethylformamide (2 mL) at room temperature; methyl (S)-4-(1-aminoethyl)-2,6-difluorobenzoate (**Compound I-38D**) (78 mg, 0.36 mmol) was added; and then O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (171 mg, 0.45 mmol) and N, N-diisopropylethylamine (58 mg, 0.45 mmol) were added. The mixture was stirred for 16 hours at room temperature, and the mixture was diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated; the residue was separated and purified by a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoate (**Compound I-38E**) (82 mg, yield 52%).

[00803] LC-MS, M/Z (ESI): 534.3 [M+H]⁺

[00804] Fifth step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoic acid (**Compound I-38**)



[00805] I-38

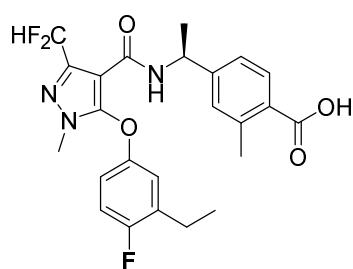
[00806] A starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoate (82 mg, 0.15 mmol) was added to tetrahydrofuran (2 mL) and water (4 mL) at room temperature, and lithium hydroxide (18.9 mg, 0.45 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 4 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)ethyl)-2,6-difluorobenzoic acid (**Compound I-38**) (16 mg, 20% yield).

[00807] ^1H NMR (400MHz, DMSO- d_6) δ 12.8 (s, 1H), 8.12 (d, 1H), 7.61 (t, 1H), 7.56 (t, 1H), 7.45 (s, 1H), 7.27 (t, 1H), 7.25 (t, 1H), 6.62 (d, 2H), 4.76 (t, 1H), 3.76 (s, 3H), 1.13 (d, 3H).

[00808] LC-MS, M/Z (ESI): 520.3 $[\text{M}+\text{H}]^+$

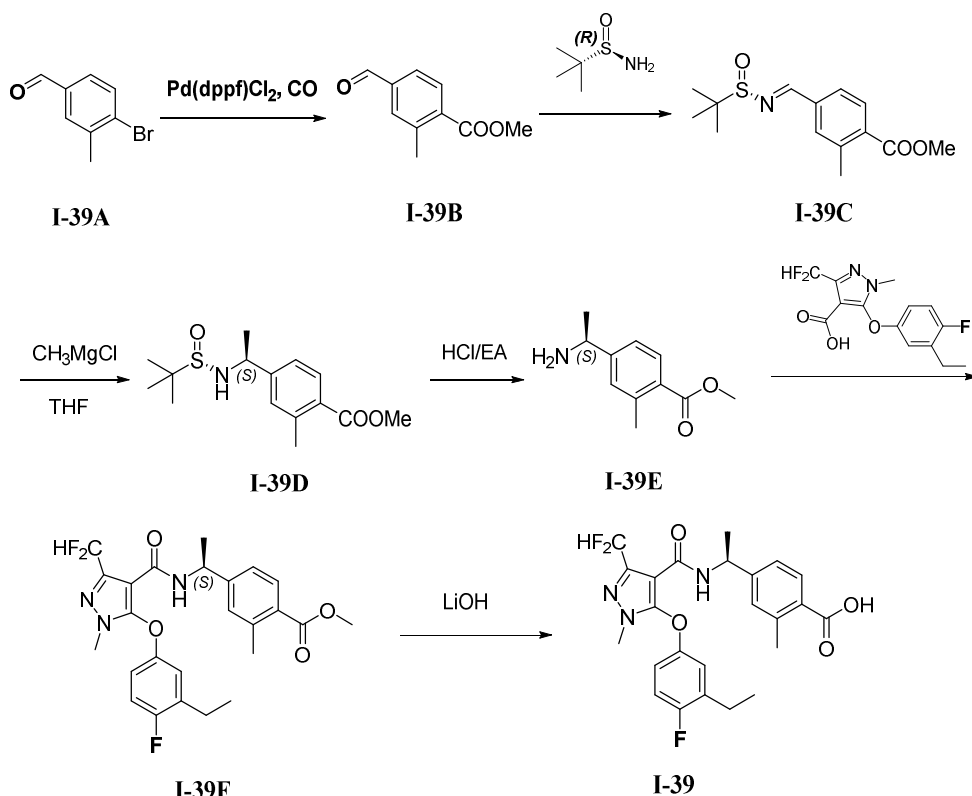
[00809] Example 39: Preparation of **Compound I-39**

[00810] (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)-2-methylbenzoic acid (**Compound I-39**)



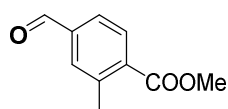
[00811] I-39

[00812] The synthesis scheme for **Compound I-39** is shown below:



[00813]

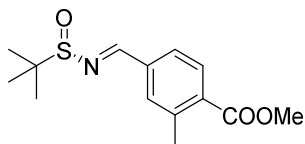
[00814] First step: synthesis of methyl 4-formyl-2-methylbenzoate (**Compound I-39B**)



[00815] **I-39B**

[00816] 4-bromo-3-methylbenzaldehyde (**Compound I-39A**) (10.0 g, 50.2 mmol) and triethylamine (15.3 g, 151 mmol) were added to methanol (200 mL); and 1, 1-bis(diphenylphosphonium) ferrocene palladium chloride (2.94 g, 4.02 mmol) was added at room temperature. The mixture was replaced by carbon monoxide for 3 times, and then reacted for 15 hours at 65°C under the pressure of 50 Psi; and then the mixture was cooled to room temperature and filtered through celite. The filter cake was washed with ethyl acetate 3 times and the filtrate was concentrated to obtain the crude product, which was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 100: 1-5: 1) to obtain the compound methyl 4-formyl-2-methylbenzoate (**Compound I-39B**) (8.0 g, yield 89%).

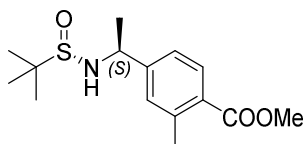
[00817] Second step: synthesis of methyl (S,E)-4-(((tert-butylsulfinyl)imino)methyl)-2-methylbenzoate (**Compound I-39C**)



[00818] I-39C

[00819] Methyl 4-formyl-2-methylbenzoate (**Compound I-39B**) (8.0 g, 44.9 mmol) and R-(+)-tert-butylsulfonamide (6.53 g, 53.9 mmol) and cesium carbonate (17.6 g, 53.9 mmol) were added to dichloromethane (200 mL). The mixture was reacted at 50°C for 15 h, added with
 5 water (200 mL) and dichloromethane (200 mL), and the aqueous phase was extracted with dichloromethane (200 mL ×3), combined and washed with brine (100 mL), dried over sodium sulfate, filtered, and spin-dried to obtain a crude product. The crude product was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 100: 1-1: 1) to obtain a compound methyl (S, E)-4-(((tert-butylsulfinyl)imino)methyl)-2-methylbenzoate (**Compound I-39C**) (10.0 g, 79% yield).
 10

[00820] Third step: synthesis of methyl 4-((S)-1-(((S)-tert-butylsulfinyl)amino)ethyl)-2-methylbenzoate (**Compound I-39D**)

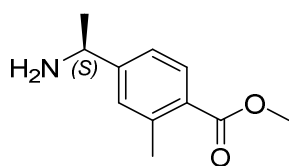


[00821] I-39D

[00822] Methyl (S, E)-4-(((tert-butylsulfinyl)imino)methyl)-2-methylbenzoate (**Compound I-39C**) (1.0 g, 3.55 mmol) was added to tetrahydrofuran (20 mL); and the mixture was cooled to -10°C. Then, methyl magnesium chloride (7.1 mL, 21.3 mmol, 3 mol/L) was slowly added dropwise and reacted at room temperature for 15 hours; water (50 mL) and ethyl acetate (100 mL) were added to the reaction. The aqueous phase was extracted with ethyl acetate (100 mL ×3); the combined extract liquor was washed with brine (100 mL), dried over sodium sulfate,
 15 filtered and spin-dried to obtain a crude product. The crude product was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 20: 1-1: 1) to obtain a compound methyl 4-((S)-1-(((S)-tert-butylsulfinyl)amino)ethyl)-2-methylbenzoate (**Compound I-39D**) (0.7 g, yield 66%).
 20

[00823] Fourth step: synthesis of methyl (S)-4-(1-aminoethyl)-2-methylbenzoate (**Compound**

I-39E)

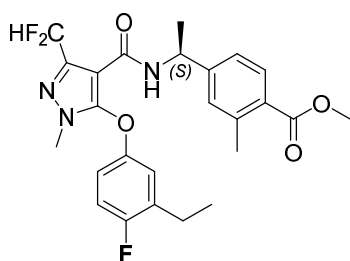


[00824] I-39E

[00825] Methyl 4-((S)-1-(((S)-tert-butylsulfinyl)amino)ethyl)-2-methylbenzoate (**Compound I-39D**) (0.7 g, 2.35 mmol) was added to ethyl acetate (10 mL); a solution (10 mL) of ethyl acetate/hydrogen chloride was added at room temperature. The reaction mixture was reacted at room temperature for 2 hours, and was spin-dried to obtain methyl (S)-4-(1-aminoethyl)-2-methylbenzoate (**Compound I-39E**) (0.4 g, yield 88%).

[00826] Fifth step: synthesis of methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)-2-methylbenzoate (**Compound**

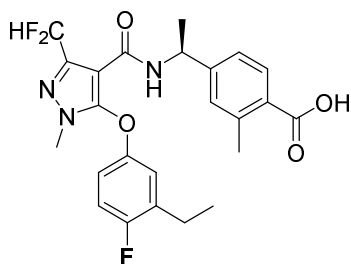
I-39F)



[00827] I-39F

[00828] Methyl (S)-4-(1-aminoethyl)-2-methylbenzoate (**Compound I-39E**) (0.1 g, 0.52 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (0.24 g, 0.62 mmol), N, N-diisopropylethylamine (0.20 g, 1.55 mmol), and 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxylic acid (0.20 g, 0.62 mmol) were added to N, N-dimethylformamide (5 mL) and reacted at room temperature for 12 hours. Then, the reaction mixture was added with water (50 mL) and ethyl acetate (100 mL), and extracted with ethyl acetate (100 mL x 4). The combined extract liquor was washed with brine (50 mL), dried over sodium sulfate, filtered, and spin-dried to obtain a crude product. The crude product was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 20: 1-1: 1) to obtain methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)-2-methylbenzoate (**Compound I-39F**) (0.15 g, yield 59%).

[00829] Sixth step: synthesis of (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)-2-methylbenzoic acid (**Compound I-39**)



[00830]

[00831] Methyl (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-

pyrazole-4-carboxamido)ethyl)-2-methylbenzoate (**Compound I-39F**) (0.15 g, 0.31 mmol) was added to tetrahydrofuran (5 mL), methanol (5 mL) and water (1 mL), and lithium hydroxide (0.03 g, 1.23 mmol) was added at room temperature. The reaction mixture reacted at room temperature for 2 hours, then the mixture was spin-dried. The crude product was added with water (50 mL) and ethyl acetate (100 mL), then the pH was adjusted to 7 with 1N hydrochloric acid, and the mixture was extracted with ethyl acetate (100 mL x 2). The organic layers were combined and washed with brine (50 mL), and the organic phase was dried over sodium sulfate and concentrated to obtain a crude product. The crude product was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a white solid of (S)-4-(1-(3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-

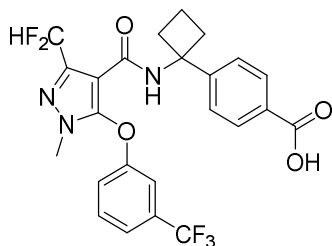
carboxamido)ethyl)-2-methylbenzoic acid (**Compound I-39**) (83 mg, 57% yield).

[00832] ^1H NMR (400 MHz, DMSO) δ 7.94 (d, J = 7.6 Hz, 1H), 7.59 (d, J = 8.0 Hz, 1H), 7.10 (dd, J = 7.6, 6.4 Hz, 2H), 6.99 – 6.94 (m, 2H), 6.89 (d, J = 8.2 Hz, 1H), 6.80 – 6.75 (m, 1H), 4.84 – 4.79 (m, 1H), 3.69 (s, 3H), 2.36 (s, 3H), 2.53 (dd, J = 14.8, 7.4 Hz, 2H) 1.19 (d, J = 7.0 Hz, 3H), 1.06 (t, J = 7.6 Hz, 3H).

[00833] LC-MS, M/Z (ESI): 476.3 $[\text{M}+\text{H}]^+$

[00834] Example 40: Preparation of Compound I-40

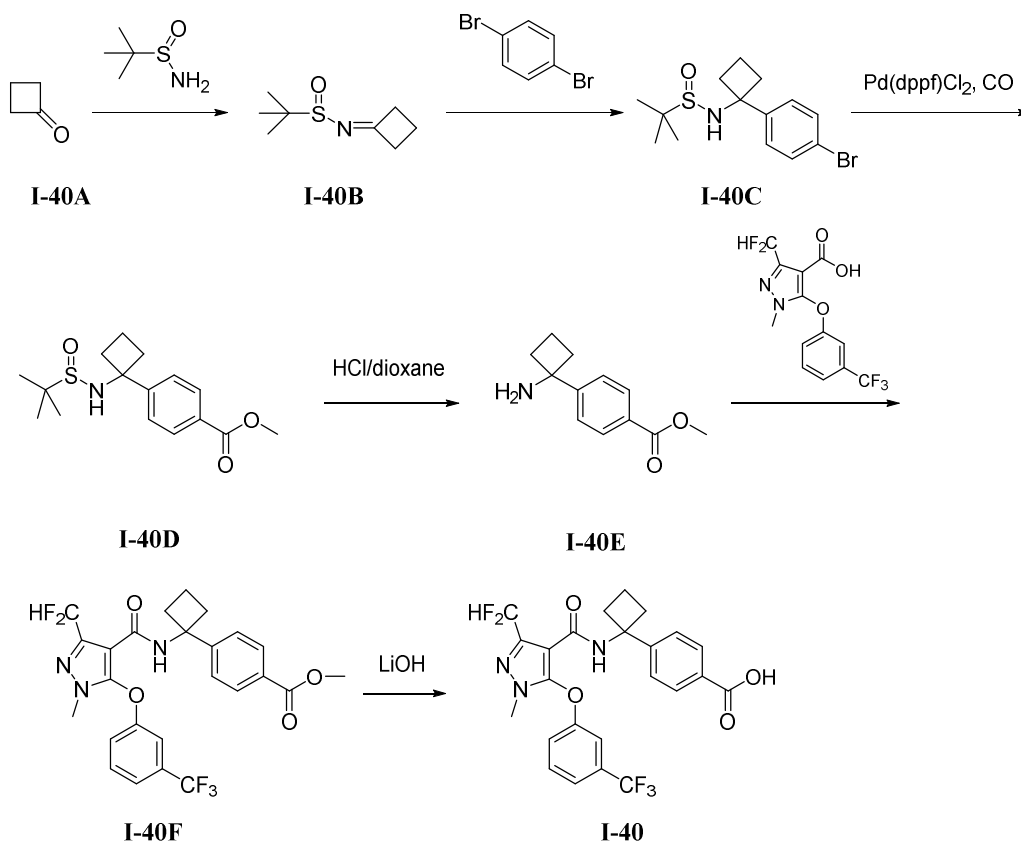
[00835] 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoic acid (**Compound I-40**)



[00836]

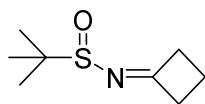
I-40

[00837] The synthesis scheme for **Compound I-40** is shown below:



[00838]

[00839] First step: N-cyclobutylidene-2-methylpropane-2-sulfinamide (**Compound I-40B**)



5

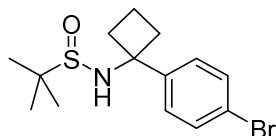
[00840] **I-40B**

[00841] The compound cyclobutanone (5.0 g, 71.4 mmol) was added to dichloromethane (100 mL) at room temperature; tert-butylsulfinamide (10.3 g, 85.6 mmol) and tetraethyl titanate (32.6 g, 143 mmol) were added; and the mixture was heated to 40°C and stirred for 24 hours. The mixture was cooled to room temperature and added with water (150 mL), and the mixture was extracted with ethyl acetate (100 mL × 3) and separated. The organic phases were combined,

10

dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a light yellow liquid of N-cyclobutylidene-2-methylpropane-2-sulfinamide (**Compound I-40B**) (6.5 g, yield 53%).

- 5 **[00842]** Second step: N-(1-(4-bromophenyl)cyclobutyl)-2-methylpropane-2-sulfinamide (**Compound I-40C**)

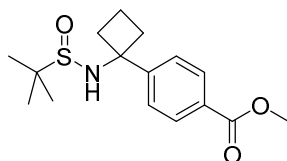


[00843] I-40C

- [00844]** The compound 1, 4-dibromobenzene (14.0 g, 59.8 mmol) was added to tetrahydrofuran (150 mL) at room temperature; the mixture was cooled to -78°C; and a 2.5 mol/L solution of butyllithium tetrahydrofuran (24.0 mL, 60.0 mmol) was added. The mixture was stirred for 40 min, and then a solution of N-cyclobutylidene-2-methylpropane-2-sulfinamide (**Compound I-40B**) (7.0 g, 40.4 mmol) in tetrahydrofuran (150 mL) was added dropwise under nitrogen protection. Then, the mixture was naturally warmed to room temperature and stirred for 2.5 hours, and then diluted with saturated aqueous ammonium chloride (200 mL), extracted with ethyl acetate (200 mL × 3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a yellow solid of N-(1-(4-bromophenyl)cyclobutyl)-2-methylpropane-2-sulfinamide (**Compound I-40C**) (1.8 g, yield 64%).

- 20 **[00845]** LC-MS, M/Z (ESI): 330.0 [M+H]⁺

[00846] Third step: methyl 4-(1-((tert-butylsulfinyl)amino)cyclobutyl)benzoate (**Compound I-40D**)

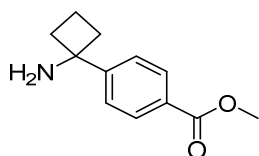


[00847] I-40D

[00848] The starting material N-(1-(4-bromophenyl)cyclobutyl)-2-methylpropane-2-

sulfonamide (**Compound I-40C**) (1.80 g, 5.47 mmol) was added to methanol (30 mL) at room temperature; triethylamine (2.76 g, 27.4 mmol) and 1, 1'-bis(diphenylphosphino)ferrocene palladium dichloride (450 mg, 0.55 mmol) were added, and carbon monoxide was charged. The mixture was heated to 85°C and stirred for 24 hours. The mixture was diluted with water (50 mL), extracted with ethyl acetate (50 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on thin-layer silica gel plate (dichloromethane: methanol (V/V) = 10: 1) to obtain a yellow liquid of methyl 4-(1-((tert-butylsulfinyl)amino)cyclobutyl)benzoate (**Compound I-40D**) (1.30 g, 77% yield).

10 **[00849]** Fourth step: methyl 4-(1-aminocyclobutyl)benzoate (**Compound I-40E**)

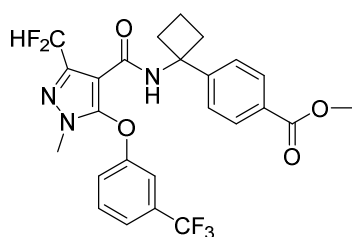


[00850] **I-40E**

[00851] The compound methyl 4-(1-((tert-butylsulfinyl)amino)cyclobutyl)benzoate (**Compound I-40D**) (600 mg, 1.94 mmol) was added to dichloromethane (10 mL), and a 4 mol/L solution of hydrochloric acid in dioxane (2.0 mL, 8.0 mmol) was added; the mixture was stirred at room temperature for 4 hours; a saturated solution of sodium bicarbonate (30 mL) was added, and the mixture was extracted with dichloromethane (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated and separated to obtain a colorless liquid crude product of methyl 4-(1-aminocyclobutyl)benzoate (**Compound I-40E**) (350 mg, yield 88%).

20 **[00852]** LC-MS, M/Z (ESI): 206.5 [M+H]⁺

[00853] Fifth step: methyl 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoate (**Compound I-40F**)

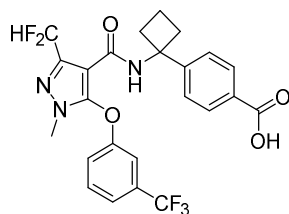


[00854] **I-40F**

[00855] The compound 3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl) phenoxy)-1H-pyrazole-4-carboxylic acid (100 mg, 0.30 mmol) was added to N, N-dimethylformamide (3 mL); methyl 4-(1-aminocyclobutyl)benzoate (**Compound I-40E**) (74 mg, 0.36 mmol), and O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (125 mg, 0.33 mmol) and N, N-diisopropylethylamine (85 mg, 0.66 mmol) were added at room temperature. The mixture was stirred for 16 hours at room temperature; and the mixture was diluted by addition of water (10 mL), extracted with ethyl acetate (10 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified on a thin-layer silica gel plate (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a white solid of methyl 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoate (**Compound I-40F**) (60 mg, yield 39%).

[00856] LC-MS, M/Z (ESI): 524.5 [M+H]⁺

[00857] Sixth step: 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoic acid (**Compound I-40**)



[00858] **I-40**

[00859] A starting material methyl 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoate (**Compound I-40F**) (60 mg, 0.12 mmol) was added to tetrahydrofuran (1 mL), methanol (1 mL) and water (1 mL) at room temperature; and lithium hydroxide (20 mg, 0.47 mmol) was added. The mixture was stirred at room temperature for 16 hours; the pH was adjusted to 2 with 1N hydrochloric acid; and the reaction mixture was concentrated to prepare, by the acidic preparation method A, a white solid of 4-(1-(3-(difluoromethyl)-1-methyl-5-(3-(trifluoromethyl)phenoxy)-1H-pyrazole-4-carboxamido)cyclobutyl)benzoic acid (**Compound I-40**) (16 mg, yield 27%).

[00860] ¹H NMR (400MHz, DMSO-d₆) δ12.7 (s, 1H), 8.52 (s, 1H), 7.74 (d, 2H), 7.68 (d, 1H), 7.62 (d, 1H), 7.50 (s, 1H), 7.34 (t, 1H), 7.25 (d, 2H), 7.18 (t, 1H), 3.78 (s, 3H), 2.33-2.28 (m,

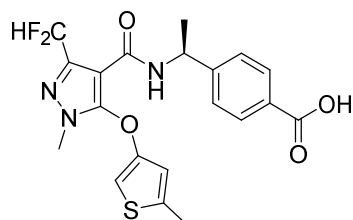
2H), 2.16-2.13 (m, 2H), 1.70-1.65 (m, 1H), 1.59-1.54 (m, 1H).

[00861] LC-MS, M/Z (ESI): 510.5 [M+H]⁺

[00862] Example 41: Preparation of Compound I-41

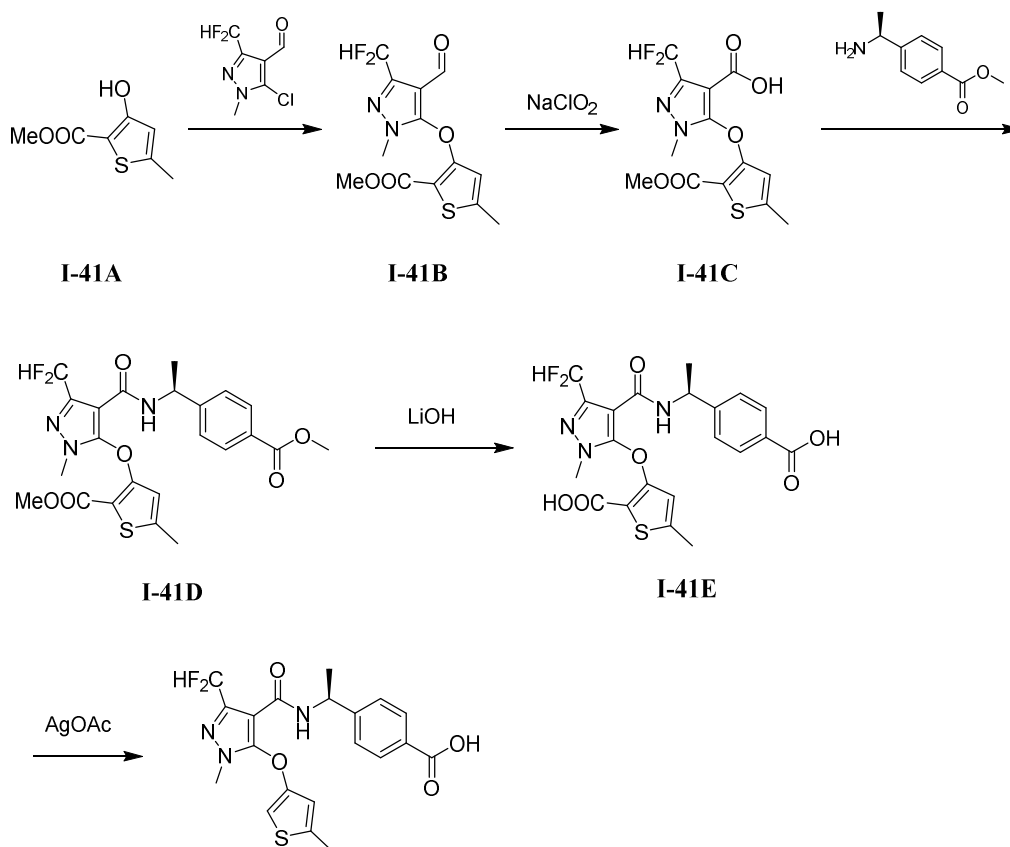
[00863] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-methylthiophen-3-yl)oxy)-1H-pyrazole-5

4-carboxamido)ethyl)benzoic acid (**Compound I-41**)



[00864] **I-41**

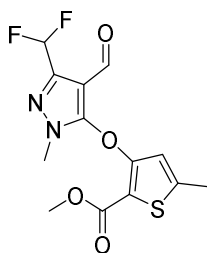
[00865] The synthesis scheme for **Compound I-41** is shown below:



[00866]

I-41

[00867] First step: methyl 3-((3-(difluoromethyl)-4-formyl-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylate (**Compound I-41B**)

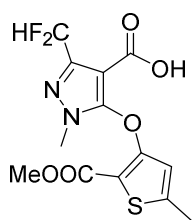


[00868] I-41B

[00869] The compound 5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carbaldehyde (120 mg, 0.62 mmol) was added to N, N-dimethylformamide (5 mL); methyl 3-hydroxy-5-methylthiophene-2-carboxylate (143 mg, 0.74 mmol), cesium carbonate (403 mg, 1.24 mmol), cuprous iodide (24.0 mg, 0.12 mmol) and 1, 10-phenanthroline (45.0 mg, 0.25 mmol) were added at room temperature and heated to 130°C in the microwave and stirred for 1 h; then the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (15 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a colorless liquid of methyl 3-((3-(difluoromethyl)-4-formyl-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylate (**Compound I-41B**) (32.0 mg, yield 16%).

[00870] LC-MS, M/Z (ESI): 331.4 (M+1).

[00871] Second step: 3-(difluoromethyl)-5-((2-(methoxycarbonyl)-5-methylthiophen-3-yl)oxy)-1-methyl-1H-pyrazole-4-carboxylic acid (**Compound I-41C**)



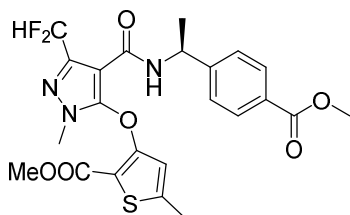
[00872] I-41C

[00873] The compound methyl 3-((3-(difluoromethyl)-4-formyl-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylate (**Compound I-41B**) (30.0 mg, 0.09 mmol) was added to tert-butanol (6 mL) and water (3 mL) at room temperature; 2-methyl-2-butene (18.9 mg, 0.27 mmol), sodium chlorite (24.0 mg, 0.27 mmol), and sodium dihydrogen phosphate (27.0 mg, 2.34 mmol) were added. The mixture was stirred for 16 hours at room temperature; and then the mixture was diluted with water (5 mL), extracted with ethyl acetate (10 mL ×3)

and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered, and concentrated to obtain a white solid of 3-(difluoromethyl)-5-((2-(methoxycarbonyl)-5-methylthiophen-3-yl)oxy)-1-methyl-1H-pyrazole-4-carboxylic acid (**Compound I-41C**) (32.0 mg, yield 100%).

5 **[00874]** LC-MS, M/Z (ESI): 347.4 (M+1)

[00875] Third step: methyl (S)-3-((3-(difluoromethyl)-4-((1-(4-(methoxycarbonyl)phenyl)ethyl)carbamoyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylate (**Compound I-41D**)

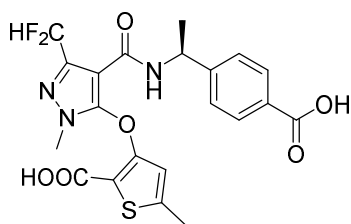


[00876] **I-41D**

10 **[00877]** The compound 3-(difluoromethyl)-5-(3-ethyl-4-fluorophenoxy)-1-methyl-1H-pyrazole-4-carboxylic acid (24.0 mg, 0.07 mmol) was added to dichloromethane (5 mL); (S)-methyl 4-(1-aminoethyl) benzoate (15.0 mg, 0.08 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-Tetramethyluronium hexafluorophosphate (40.0 mg, 0.10 mmol) and N, N-diisopropylethylamine (14.0 mg, 0.10 mmol) were added; and the mixture was stirred at room
15 temperature for 16 hours, diluted with water (20 mL), extracted with ethyl acetate (10 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered, concentrated, and the residue purified by silica gel column separation (petroleum ether: ethyl acetate (V/V) = 3: 1) to obtain a colorless liquid of methyl (S)-3-((3-(difluoromethyl)-4-((1-(4-(methoxycarbonyl)phenyl)ethyl)carbamoyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-
20 methylthiophene-2-carboxylate (**Compound I-41D**) (25.0 mg, yield 71.0%).

[00878] LC-MS, M/Z (ESI): 508.6 (M+1).

[00879] Fourth step: (S)-3-((4-((1-(4-carboxyphenyl)ethyl)carbamoyl)-3-(difluoromethyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylic acid (**Compound I-41E**)

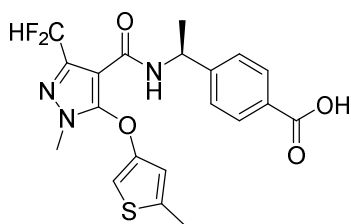


[00880] I-41E

[00881] The starting material methyl (S)-3-((3-(difluoromethyl)-4-((1-(4-methoxycarbonyl)phenyl)ethyl)carbamoyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylate (38.0 mg, 0.08 mmol) was added to tetrahydrofuran (5 mL) and water (3 mL) at room temperature, and lithium hydroxide (10.0 mg, 0.22 mmol) was added. The mixture was stirred at room temperature for 3 hours; the pH was adjusted to 6 with 1N hydrochloric acid; and the mixture was extracted with ethyl acetate (10 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated to obtain a white solid crude product of (S)-3-((4-((1-(4-carboxyphenyl)ethyl)carbamoyl)-3-(difluoromethyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylic acid (**Compound I-41E**) (33.0 mg, yield 92%).

[00882] LC-MS, M/Z (ESI): 480.3 (M+1)

[00883] Fifth step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-methylthiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-41**)



[00884] I-41

[00885] The starting material (S)-3-((4-((1-(4-carboxyphenyl)ethyl)carbamoyl)-3-(difluoromethyl)-1-methyl-1H-pyrazol-5-yl)oxy)-5-methylthiophene-2-carboxylic acid (30.0 mg, 0.06 mmol) was added to N-methylpyrrolidone (6 mL) at room temperature; silver acetate (10.2 mg, 0.06 mmol) was added; and the mixture was heated to 100°C under nitrogen protection and stirred for 0.5 hour. The reaction mixture was filtered to prepare, via the Acidic Preparation Method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-methylthiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-41**)

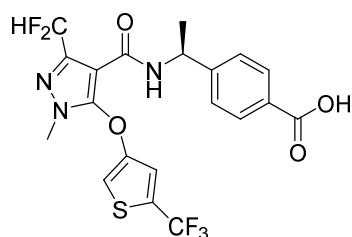
(6.7 mg, 25% yield).

[00886] ^1H NMR (400m Hz, CDCl_3) δ 12.8 (s, 1H), 7.99 (d, 2H), 7.20 (t, 1H), 7.20 (d, 2H), 6.51 (s, 1H), 6.39 (d, 1H), 6.20 (s, 1H), 5.22 (t, 1H), 3.75 (s, 3H), 2.45 (s, 3H), 1.43 (d, 3H)。

[00887] LC-MS, M/Z (ESI): 436.1 (M+1)

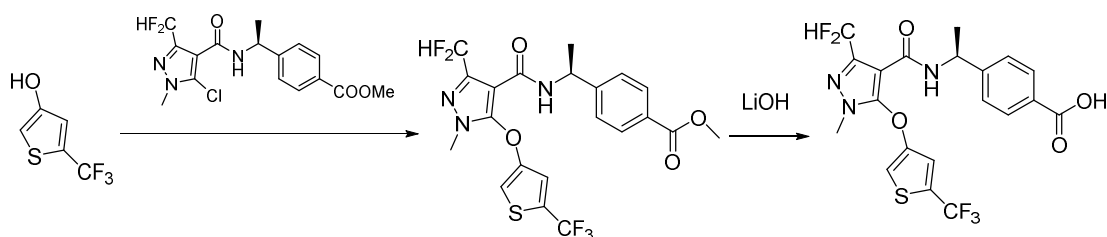
5 **[00888]** Example 42: Preparation of Compound I-42

[00889] (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-42**)



[00890] **I-42**

[00891] The synthesis scheme for **Compound I-42** is shown below:

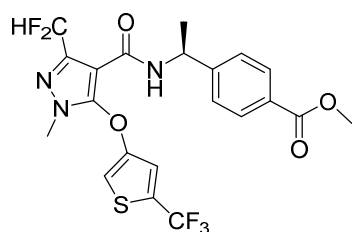


10 **[00892]** **I-42A**

I-42B

I-42

[00893] First step: methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-42B**)



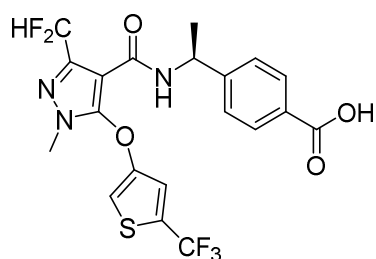
15 **[00894]** **I-42B**

[00895] The compound methyl (S)-4-(1-(5-chloro-3-(difluoromethyl)-1-methyl-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Intermediate A**) (1.0 g, 2.69 mmol) was added to dimethyl

sulfoxide (5 mL); 5-(trifluoromethyl) thiophen-3-ol (908 mg, 5.40 mmol), potassium carbonate (1.13 g, 8.18 mmol), cuprous iodide (206 mg, 1.08 mmol), and 1, 10-phenanthroline (388 mg, 2.16 mmol) were added at room temperature; and the mixture was heated to 120°C in the microwave and stirred for 4 hours. Then, the mixture was cooled to room temperature, diluted with water (20 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 3) to obtain a white solid of methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-42B**) (980 mg, yield 72%).

[00896] LC-MS, M/Z (ESI): 504.2 (M+1)

[00897] Second step: (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-42**)



[00898] **I-42**

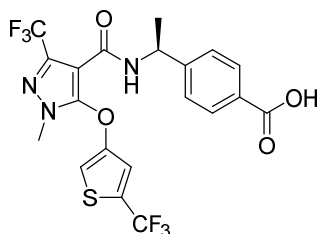
[00899] The starting material methyl (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-42B**) (980 mg, 1.89 mmol) was added to tetrahydrofuran (5 mL); water (3 mL) and lithium hydroxide (238 mg, 5.67 mmol) were added at room temperature; and the mixture was stirred for 3 hours at room temperature. Then, the reaction mixture was concentrated to prepare, via the acidic preparation method A, a white solid of (S)-4-(1-(3-(difluoromethyl)-1-methyl-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-42**) (345 mg, 36% yield).

[00900] ¹H NMR (400m Hz, DMSO-d₆) δ12.8 (s, 1H), 8.20 (d, 1H), 7.79 (d, 2H), 7.67 (s, 1H), 7.25 (d, 2H), 7.23 (d, 1H), 7.09 (t, 1H), 4.99-4.92 (m, 1H), 3.77 (s, 3H), 1.29 (d, 3H).

[00901] LC-MS, M/Z (ESI): 490.2 (M+1)

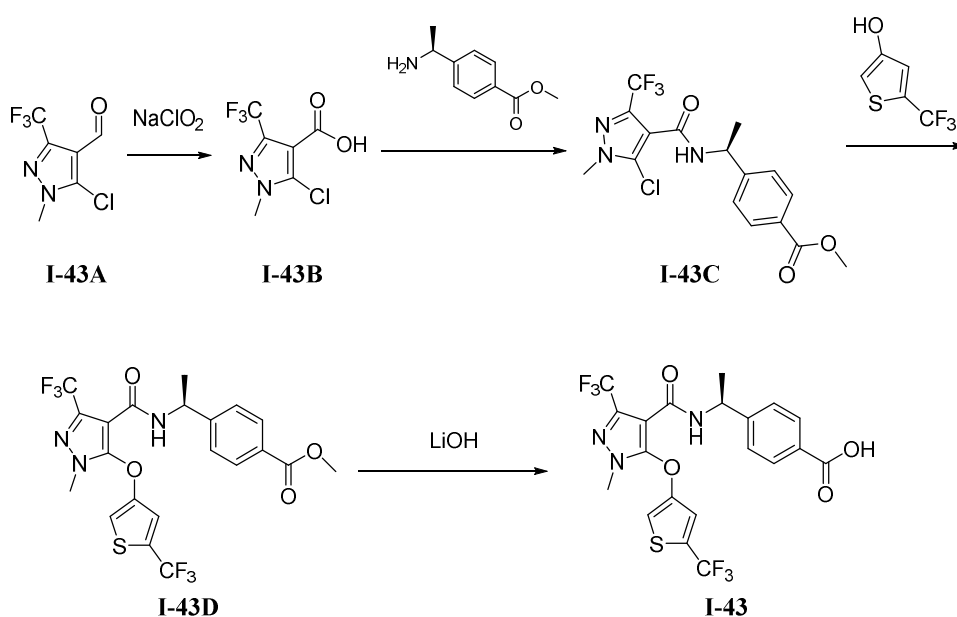
[00902] Example 43: Preparation of Compound I-43

[00903] (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-43**)



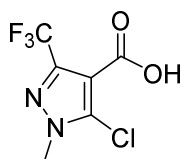
[00904] **I-43**

5 [00905] The synthesis scheme for **Compound I-43** is shown below:



[00906]

[00907] First step: 5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxylic acid (**Compound I-43B**)



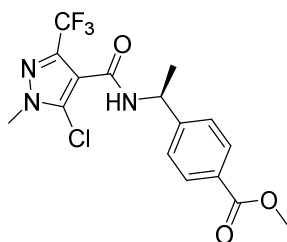
[00908] **I-43B**

10 [00909] The compound 5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carbaldehyde (700 mg, 3.30 mmol) was added to tert-butanol (20 mL) and water (5 mL) at room temperature; 2-methyl-2-butene (1.80 g, 25.7 mmol), sodium chlorite (1.48 g, 16.4 mmol) and sodium dihydrogen phosphate (3.10 g, 25.8 mmol) were added. The mixture was stirred for 14 hours at room temperature; and then the mixture was diluted with water (50 mL), extracted with ethyl

acetate (30 mL ×3) and separated. The organic phases were combined and dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid crude product of 5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxylic acid (**Compound I-43B**) (680 mg, yield 90%).

[00910] LC-MS, M/Z (ESI): 229.6 (M+1).

[00911] Second step: methyl (S)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-43C**)

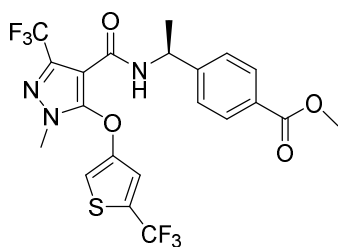


[00912] I-43C

[00913] The compound 5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxylic acid (**Compound I-43B**) (680 mg, 2.98 mmol) was added to N, N-dimethylformamide (20 mL); methyl (S)-4-(1-aminoethyl) benzoate (537 mg, 3.00 mmol), O-(7-azabenzotriazol-1-yl)-N, N, N, N-tetramethyluronium hexafluorophosphate (1.70 g, 4.47 mmol) and N, N-diisopropylethylamine (1.90 g, 14.7 mmol) were added and stirred at room temperature for 16 hours; the mixture was diluted with water (200 mL), extracted with ethyl acetate (30 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a colorless solid of methyl (S)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-43C**) (700 mg, yield 60%).

[00914] LC-MS, M/Z (ESI): 390.5 (M+1).

[00915] Third step: methyl (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-43D**)

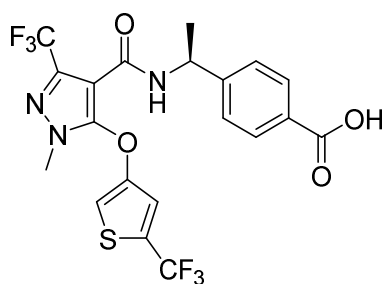


[00916] **I-43D**

[00917] The compound methyl (S)-4-(1-(5-chloro-1-methyl-3-(trifluoromethyl)-1H-pyrazole-4-carboxamido)ethyl)benzoate (500 mg, 1.3 mmol) was added to dimethyl sulfoxide (10 mL); 5-(trifluoromethyl) thiophen-3-ol (216 mg, 1.3 mmol), potassium carbonate (360 mg, 2.6 mmol), cuprous iodide (100 mg, 0.5 mmol) and 1, 10-phenanthroline (100 mg, 0.5 mmol) were added at room temperature; and the mixture was heated to 95°C in the microwave and stirred for 5 hours. Then, the mixture was cooled to room temperature, diluted with water (50 mL), extracted with ethyl acetate (20 mL ×3) and separated. The organic phases were combined, dried over anhydrous sodium sulfate, filtered and concentrated, and the residue was separated and purified by a silica gel column (petroleum ether: ethyl acetate (V/V) = 1: 1) to obtain a yellow solid of methyl (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-43D**) (200 mg, yield 30%).

[00918] LC-MS, M/Z (ESI): 522.1 (M+1).

[00919] Fourth step: (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-43**)



[00920] **I-43**

[00921] The starting material methyl (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoate (**Compound I-43D**) (100 mg, 0.19 mmol) was added to tetrahydrofuran (30 mL); water (10 mL) and lithium hydroxide (24 mg, 0.57 mmol) were added at room temperature; and the mixture was heated to

50°C and stirred for 2 hours. The reaction mixture was concentrated to prepare, by the acidic preparation method A, a yellow solid of (S)-4-(1-(1-methyl-3-(trifluoromethyl)-5-((5-(trifluoromethyl)thiophen-3-yl)oxy)-1H-pyrazole-4-carboxamido)ethyl)benzoic acid (**Compound I-43**) (54 mg, 56% yield).

5 **[00922]** ¹H NMR (400mHz, DMSO-d₆) δ12.8 (s, 1H), 8.64 (d, 1H), 7.80 (d, 2H), 7.68 (s, 1H), 7.31 (d, 1H), 7.24 (d, 2H), 4.92 (t, 1H), 3.80 (s, 3H), 1.26 (d, 3H).

[00923] LC-MS, M/Z (ESI): 508.1 (M+1)

[00924] Assay Examples for Biological Activity and Related Properties

[00925] Assay Example 1: assay for EP4 antagonistic effect

10 **[00926]** The control compound and the compounds prepared by Examples 1 to 43 were tested separately for antagonistic effect of EP4, and the assay was performed in a CHO stable cell line, which highly expresses the human EP4 receptor.

[00927] After trypsinization, cells were resuspended in a buffer (1× HBSS, 0.1% BSA, 20 mM HEPES, and 500 μM IBMX) and 8000 cells were seeded per well in 384-well plates in a seeding volume of 15μL. A working solution the compound with a 8X concentration was prepared with an experimental buffer, and then 2.5 μL of the working solution of the compound with 8X concentration was added to the 384-well plate, respectively, and incubated at 37°C for 30 min. An agonist PGE₂ working solution with 8X concentration of (4 nM) was prepared with the experimental buffer and 2.5 μL per well was added to the 384-well plate (the final PGE₂ concentration was 0.5 nM) and incubated at 37°C for 30 min. After the reaction was completed, the amount of cAMP in the cells was quantified according to the method in the instructions of the cAMP test kit (Perkin Elmer, Cat # TRF0263). The antagonistic effects (IC₅₀ value) were calculated for the test compounds.

[00928] Table 1 Antagonistic effects on EP4 by test compounds

Test compounds	IC ₅₀ (nM)
Control Compound	44
Compound I-1	3.7
Compound I-2	1.9
Compound I-3	2.8
Compound I-5	2.2
Compound I-6	8.2
Compound I-9	29
Compound I-28	5.7

Compound I-29	14
Compound I-30	12
Compound I-35	6.6
Compound I-42	8.6

[00929] Experimental results indicate that the compounds of the present application have a good antagonistic effect on EP4; compared with the control compound, the antagonistic effect of most of compounds is more than 5 times that of the control compound; and the compounds of the present disclosure exhibit a better antagonistic effect on the EP4 receptor.

5 **[00930]** Assay Example 2: Assay of calcium current inhibition on EP4 receptor

[00931] The control compound and the compounds prepared in Examples 1 to 43 were respectively tested for their inhibitory effect on EP4 calcium current, and the assay was performed on the 293 cells, which overexpress the human EP4 receptor.

[00932] The cells were rapidly thawed in a 37°C water bath, centrifuged, resuspended, and
10 counted. The cell suspensions were seeded at 20 µL/well in two 384-well plates (20,000 cells/well) and placed in an incubator (a 37°C, 5% CO₂) overnight. Preparation of a 2X Fluo-4 Direct™ (Invitrogen, Cat # F10471) loading buffer: 77 mg of probenecid was added to 1 mL of a FLIPR buffer, with a concentration of 250 mM. 10 mL of FLIPR buffer and 0.2 mL of probenecid (250 mM) were added to each tube of Fluo-4 Direct™ crystals (F10471).

15 **[00933]** One of the cell plates was taken out of the incubator and the medium was removed. 20 µL of an assay buffer and 2X Fluo-4 Direct™ non-wash loading buffer were added to the 384-well cell culture plate to a final volume of 40 µL, and then the culture plate was incubated in an incubator (a 37°C, 5% CO₂) for 50 minutes, and then at room temperature for 10 minutes, and then the plate was placed in the FLIPR. 10 µL of the buffer was transferred to the cell plate and
20 the fluorescence signal was read. The agonist PGE₂ was formulated as a 10 mM stock in a DMSO solvent and 10 concentration points of the 6X working solution were serially diluted using the buffer. 10 µL of the agonist PGE₂ was transferred to the cell plate, the fluorescence signal was read to calculate EC₈₀ value.

[00934] The agonist PGE₂ was prepared at 6X EC₈₀ concentration and the compound to be
25 tested was formulated as a 10 mM stock solution in the DMSO solvent and 10 concentration points of the 6X compound working solution was serially diluted using the buffer.

[00935] Another cell plate was taken to remove media and 20 µL of the assay buffer and 2X

Fluo-4 Direct™ non-wash loading buffer were added; the cell plate was incubated in a 37°C, 5% CO₂ incubator for 50 minutes and room temperature for 10 minutes and then placed in the FLIPR. 10µL of the compound working solution, DMSO, and the EP4 full antagonist were transferred to the cell plate and the fluorescence signal was read. 10 µL of the agonist PGE₂, with a concentration of 6X EC₈₀ was transferred to the cell plate, fluorescence signal was read to calculate the inhibition rate.

[00936] Inhibition (%) = 100-(test group-EP4 full antagonist group)/(DMSO group-EP4 full antagonist group) * 100

[00937] Compound's IC₅₀ value of EP4 calcium current inhibition was calculated based on the inhibition rate at different concentrations of the compound.

[00938] Table 2 Effects of test compounds on EP4 calcium current inhibition

Test compounds	IC ₅₀ (nM)
Control Compounds	21
Compound I-1	15
Compound I-2	7.4
Compound I-3	3
Compound I-4	4.0
Compound I-5	12
Compound I-6	8.0
Compound I-8	16.5
Compound I-10	14
Compound I-11	8.0
Compound I-12	19
Compound I-13	10
Compound I-15	18
Compound I-17	18
Compound I-26	8.5
Compound I-28	8
Compound I-30	15
Compound I-33	12
Compound I-35	6

Compound I-41	16
Compound I-42	15

[00939] The experimental results indicate that the compounds of the present application exhibit better inhibition effects on the calcium current of EP4, and they are superior to the control compound, especially the Compound I-4; and the inhibition effects of the compounds on the calcium current of EP4 are increased by more than 5 time, compared with the control compound.

5 The compounds of the present disclosure exhibit better inhibition effects on the calcium current of EP4.

[00940] Assay Example 3: Radioligand binding assay for EP4 receptor

[00941] The bindings, to the radioligand EP4, of a control compound and the compounds prepared by Examples 1 to 43 were tested using recombinant human EP4 receptor membrane protein (prepared from 293 cells overexpressing human EP4 receptor). The compounds to be tested and PGE₂ were formulated as 10 mM stocks in a DMSO solvent, and then 8 concentration points of 4×working solution were serially diluted using a buffer (50 mM HBSS, 0.1% BSA, 500 mM NaCl). 1 μL of the compound working solution, DMSO, and the PGE₂ working solution were added to the assay plate, respectively; 100 μL of a EP4 receptor membrane protein (20 μg /well) and 100 μL of a radioligand [³H]-PGE₂ (PerkinElmer, Cat: NET428250UC, Lot: 2469552) (final concentration of 1.5 nM) were added and incubated at sealed room temperature for 1 hour. Unifilter-96 GF/C filter plate (Perkin Elmer) was soaked with 0.5% BSA, 50 μL per well, for at least 30 min at room temperature. After binding was complete, the reaction mixture was harvested through a GF/C plate using a Perkin Elmer Filtermate Harvester, followed by washing the filter plate and drying the filter plate for 1 hour at 50°C. After drying, the bottom of the filter plate wells was sealed using a Perkin Elmer Unifilter-96 sealing tape, and 50 μL of MicroScint TM-20 cocktail (Perkin Elmer) was added to seal the top of the filter plate. The ³H counts captured on the filter were read using a Perkin Elmer MicroBeta2 Reader.

[00942] Data were analyzed using GraphPad Prism 5 and inhibition rates were calculated according to the following formula:

[00943] Inhibition (%) = 100-(test group-PGE₂ group)/(DMSO group-PGE₂ group) * 100

[00944] The IC₅₀ and Ki values of the compounds determined by the radioligand binding assay for EP4 receptor were calculated based on the inhibition rate of the different concentrations of

the compounds.

[00945] Table 3 IC₅₀ and Ki values for test compounds determined by radioligand binding assay for EP4 receptor

Test compounds	IC ₅₀ (nM)	Ki (nM)
Control Compounds	30	16
Compound I-1	16	8.9
Compound I-2	5.5	3.0
Compound I-3	9.0	4.9
Compound I-4	7.4	4.0
Compound I-5	5.3	2.9
Compound I-6	11	5.8
Compound I-10	4.0	2.2
Compound I-11	6.8	3.7
Compound I-15	12	6.6
Compound I-23	13	7.0
Compound I-30	12	6.6
Compound I-33	31	17
Compound I-35	28	15
Compound I-42	13	7.3

[00946] The experimental results indicate that, compared with the control compound, the compounds of the present disclosure have a better affinity with the EP4 receptor, which is superior to the control compound, especially Compound I-2 and Compound I-5; and their affinity with the EP4 receptor is increased by more than 5 times, compared with the control compound. The compounds of the present disclosure exhibit better affinity with the EP4 receptor.

[00947] Assay Example 4: Pharmacokinetic Assay

[00948] In this assay, the pharmacokinetic parameters of a control compound and the compounds prepared by Examples 1 to 43 in mouse, rat, and canine subjects were tested, respectively.

[00949] In a mouse pharmacokinetic assay, male ICR mice (20 g to 25 g) were selected and fasted overnight. Three mice were selected and intragastrically administered with 5 mg/kg of the corresponding compound. The blood of the mice was collected before the administration and at 15 min, 30 min, 1 h, 2 h, 4 h, 8 h, 24 h after the administration. Another 3 mice were intravenously administered with 1 mg/kg of the corresponding compound, and the blood thereof was collected before the administration and at 15 min, 30 min, 1 h, 2 h, 4 h, 8 h, 24 h after the

administration. The blood samples were centrifuged at 6800 g at 2°C to 8°C for 6 minutes, and plasma was collected and stored at -80°C. The plasma was taken at respective time point, mixed with 3-5 times the amount of acetonitrile containing internal standard, vortexed for 1 minute, centrifuged at 4°C for 10 minutes at 13000 rpm, supernatant was added 3 times the amount of water, and the appropriate mixture was taken for LC-MS/MS analysis. The primary pharmacokinetic parameters were analyzed with a WinNonlin 7.0 software, with non-compartmental model.

[00950] In a rat pharmacokinetic assay, male SD rats (180 g to 240 g) were selected and fasted overnight. Three rats were intragastrically administered with 5 mg/kg of the corresponding compound. Another 3 rats were intravenously administered with 1 mg/kg of the corresponding compound. The following operations were the same as the mouse pharmacokinetic assay.

[00951] In a canine pharmacokinetic assay, male Beagle dogs (8 kg to 10 Kg) were selected and fasted overnight. Three Beagle dogs were taken and intragastrically administered with 3 mg/kg of the corresponding compound. Another 3 Beagle dogs were intravenously administered with 1 mg/kg of the corresponding compound. The following operations were the same as the mouse pharmacokinetic assay.

[00952] Table 4-1 Pharmacokinetic assay results for intravenous administration in mouse model

Test compounds	Intravenous administration (1 mg/kg)			
	CL (L/h/kg)	V _z (L/kg)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compound	0.88	8.38	974	6.64
Compound I-5	0.76	5.62	1264	5.11

[00953] Table 4-2 Pharmacokinetic assay results for intragastric administration in mouse model

Test compounds	Oral gavage administration (5 mg/kg)			
	C _{max} (ng/mL)	T _{max} (hr)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compound	792	0.25	1766	30.93
Compound I-1	1114	0.33	2135	4.33
Compound I-2	1837	0.33	2139	15.68
Compound I-5	1450	0.33	2709	7
Compound I-6	1285	0.42	4436	7.22
Compound I-33	2528	0.33	8292	8.22
Compound I-35	1040	0.33	3060	7.12

Compound I-42	570	0.33	2444	23.25
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[00954] Table 5-1 Pharmacokinetic assay results for intravenous administration in rat model

Test compounds	Intravenous administration (1 mg/kg)			
	CL (L/h/kg)	V _z (L/kg)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compound	0.61	9.55	1425	10.9
Compound I-5	0.57	7.66	1668	9.3
Compound I-33	0.22	2.72	4668	9.02

[00955] Table 5-2 Pharmacokinetic assay results for intragastric administration in rat model

Test compounds	Oral gavage administration (5 mg/kg)			
	C _{max} (ng/mL)	T _{max} (hr)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compounds	1102	0.5	6849	12.67
Compound I-5	993	2.00	10785	9.87

[00956] Table 6-1 Pharmacokinetic assay results for intravenous administration in canine model

Test compounds	Intravenous administration (1 mg/kg)			
	CL (L/h/kg)	V _z (L/kg)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compound	0.24	3.21	3730	9.36
Compound I-5	0.21	2.84	4373	9.73

5 **[00957]** Table 6-2 Pharmacokinetic assay results for intragastric administration in canine model

Test compounds	Oral gavage administration (3 mg/kg)			
	C _{max} (ng/mL)	T _{max} (hr)	AUC _{0-t} (h*ng/mL)	T _{1/2} (h)
Control Compound	1357	1.0	6802	18.6
Compound I-5	4822	0.25	10341	19.9
Compound I-42	4853	0.25	7187	8.35

[00958] The experimental results indicate that compared to the control compound, the compounds of the present disclosure have lower clearance for intravenous administration and higher exposure for oral administration, in particular compound I-5; the exposure after oral administration in rats and canines is about 2 times higher than that of the control compound; and the compounds of the present disclosure exhibit superior pharmacokinetic properties and are good druggability.

[00959] Assay Example 5: Anti-tumor effect of test compounds in a CT-26 murine colon cancer tumor model, in combination with radiation therapy

[00960] In this assay, the anti-tumor effects of the control compound and the compounds prepared in Examples 1 to 43, in combination with radiation therapy, were tested in a CT-26 murine colon cancer tumor model.

[00961] After one week of adaptive feeding of mice, CT-26 cells in a log phase were resuspended in PBS, 5×10^5 CT-26 cells were inoculated subcutaneously at the right flank at 100 μ L per mouse, and tumor growth was observed regularly. When the tumor grew to an average volume of 60 to 80 mm³, the mice carrying the tumor were randomly divided into 5 groups based on the tumor volume size, each group containing 10 mice of 10. These group were a group of Control compound (150 mg/kg) in combination with radiotherapy (3Gry), a group of Compound I-2 (150 mg/kg) in combination with radiotherapy, a group of Compound I-5 (150 mg/kg) in combination with radiotherapy, a group of radiotherapy (3Gry) alone, and a group of vehicle control. The mice were intragastrically administered with the corresponding compound once daily, for 23 days in total; radiation therapy was performed for one-time on the first day of the administration. Tumor volumes and mouse body weights were measured twice a week, and tumor weights were weighed at the end of the experiment. Efficacy evaluations were performed based on relative tumor inhibition (TGI), safety evaluations were performed based on changes in animal body weight and death. Tumor volume and relative tumor inhibition were calculated as follows:

[00962] Tumor volume (TV) = $1/2 \times a \times b^2$, where a and b are the length and width of the tumor measurement, respectively.

[00963] Relative tumor inhibition rate TGI (%) = $(TW_c - TW_t / TW_c) \times 100\%$, where TW_c is the mean tumor weight of the vehicle control group and TW is the mean tumor weight of the treatment group.

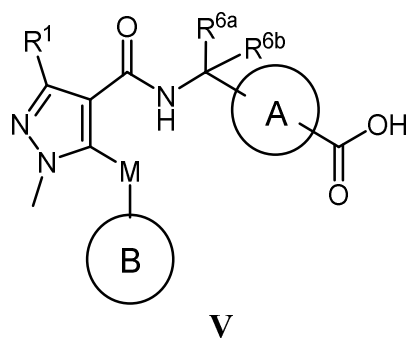
[00964] The experimental results are illustrated in FIG. 1. FIG. 1 indicates that the control compound, the test compounds 1-2 and 1-5, in combination with radiotherapy, all exhibited significant tumor inhibition effect at day 23 after the beginning of administration, and the relative tumor inhibition rates TGI (%) were 54%, 63%, and 79%, respectively, which are all statistically significantly different from the vehicle control group (p-mean smaller than 0.05); the Compound I-5 in combination with radiotherapy group was statistically significantly different from the group of radiotherapy alone (p smaller than 0.05), and better than the group

of the control compound in combination with radiotherapy. None of the groups in combination with radiotherapy neither had animal deaths nor exhibited apparent drug toxicity, exhibiting good tolerance during the treatment.

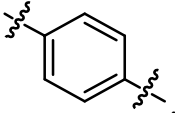
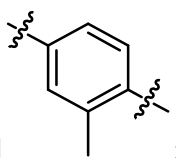
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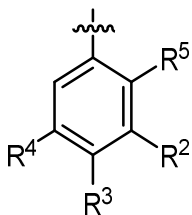
What is claimed is:

1. A compound, being a compound represented by Formula V, or being a tautomer, stereoisomer, hydrate, solvate or salt of the compound represented by Formula V:



, wherein

the ring A is selected from , and  ;

the ring B is  ;

R¹ is selected from -CH₃, -CHF₂, and -CF₃;

R² is selected from C₂-C₆ alkyl, C₃-C₆ cycloalkyl, phenyl, C₂-C₆ halogen-substituted alkyl, C₃-C₆ halogen-substituted cycloalkyl, C₂-C₆ hydroxy-substituted alkyl, C₂-C₆ cyano-substituted alkyl, and -X-R^{2a}, where X is selected from sulfur, and -CO-, and R^{2a} is selected from C₁-C₆ alkyl and C₁-C₆ halogen-substituted alkyl;

R³ is selected from hydrogen, halogen, C₁-C₂ alkyl, C₁-C₂ fluorine-substituted alkyl, and phenyl;

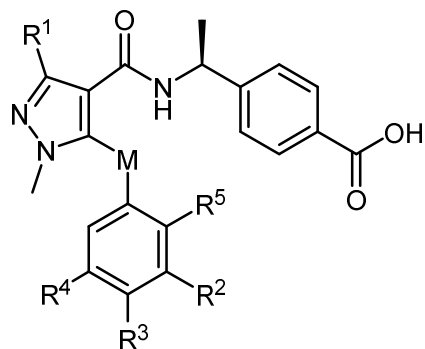
R⁴ is selected from hydrogen, halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₆ halogen-substituted alkyl, and C₁-C₆ halogen-substituted alkoxy;

R⁵ is selected from hydrogen and halogen;

one of R^{6a} and R^{6b} is hydrogen, and the other one of R^{6a} and R^{6b} is methyl; and

M is selected from oxygen, sulfur, and methylene.

2. The compound according to claim 1, being a compound represented by Formula II, or being a tautomer, stereoisomer, hydrate, solvate or salt of the compound represented by Formula II:



II, wherein

R¹ is selected from -CH₃, -CHF₂, and -CF₃, and preferably, R¹ is -CHF₂;

R² is selected from ethyl, propyl, isopropyl, n-butyl, isobutyl, tert-butyl, fluoro-substituted ethyl, fluoro-substituted propyl, fluoro-substituted isopropyl, fluoro-substituted butyl, fluoro-substituted isobutyl, hydroxyethyl, hydroxyisopropyl, cyanoethyl, phenyl, and -X-R^{2a}, where X is selected from sulfur, and -CO-, and R^{2a} is selected from methyl, ethyl, fluoro-substituted methyl, and fluoro-substituted ethyl;

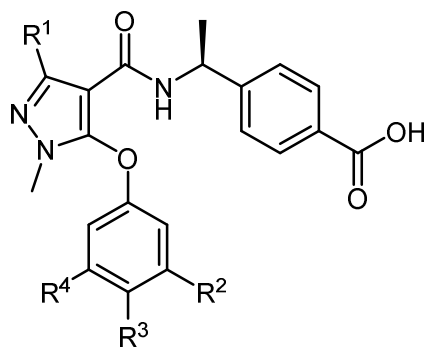
R³ is selected from hydrogen, fluorine, chlorine, methyl, ethyl, fluoro-substituted methyl, fluoro-substituted ethyl, and phenyl;

R⁴ is selected from hydrogen, fluorine, chlorine, methyl, ethyl, fluoro-substituted methyl, and fluoro-substituted ethyl;

R⁵ is selected from hydrogen, fluorine, and chlorine; and

M is selected from oxygen, sulfur, and methylene.

3. A compound, being a compound represented by Formula I, or being a tautomer, stereoisomer, hydrate, solvate or salt of the compound represented by Formula I:



I, wherein

R¹ is selected from -CH₃, -CHF₂, and -CF₃;

R² is selected from C₂-C₆ alkyl, C₃-C₆ cycloalkyl, C₂-C₆ halogen-substituted alkyl, and C₃-C₆ halogen-substituted cycloalkyl;

R³ is selected from hydrogen, halogen, C₁-C₂ alkyl, and C₁-C₂ fluorine-substituted alkyl;

R⁴ is selected from hydrogen, halogen, C₁-C₆ alkyl, C₁-C₆ alkoxy, C₁-C₆ halogen-substituted alkyl, and C₁-C₆ halogen-substituted alkoxy.

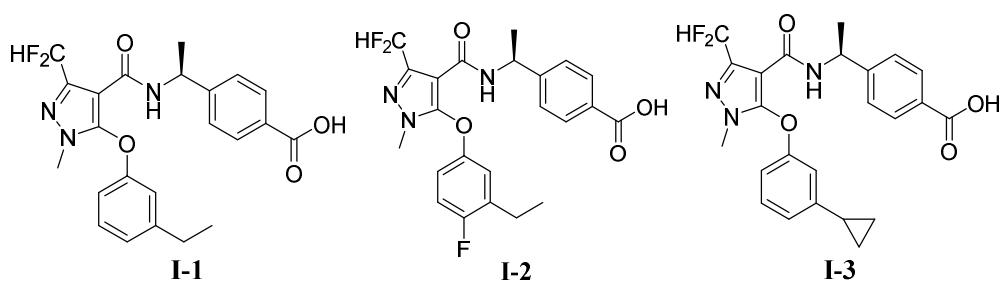
4. The compound according to claim 3, wherein

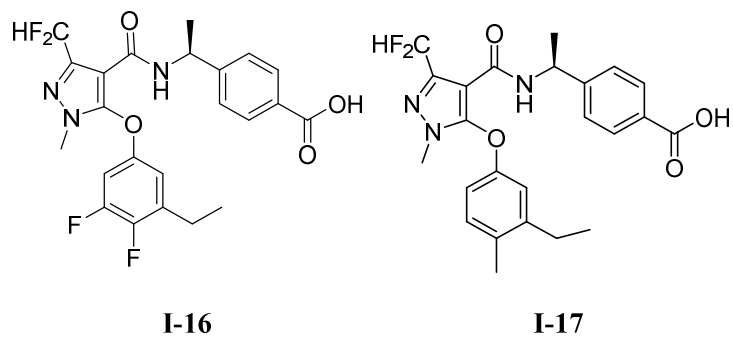
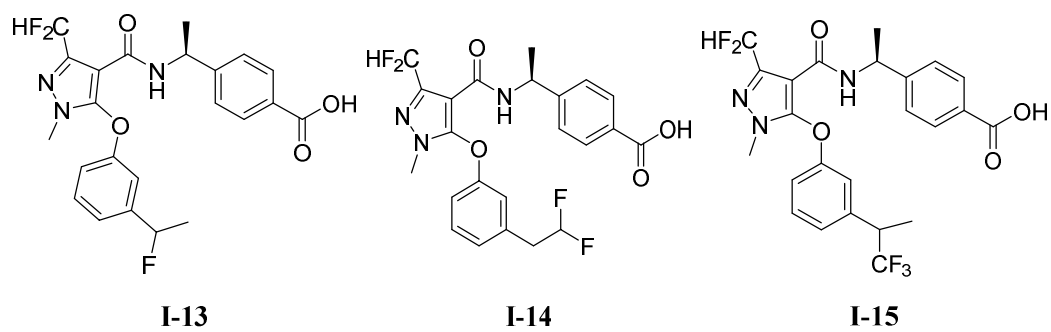
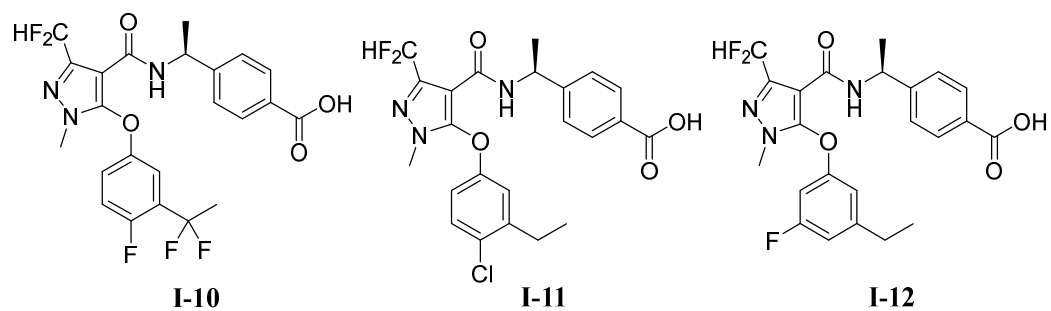
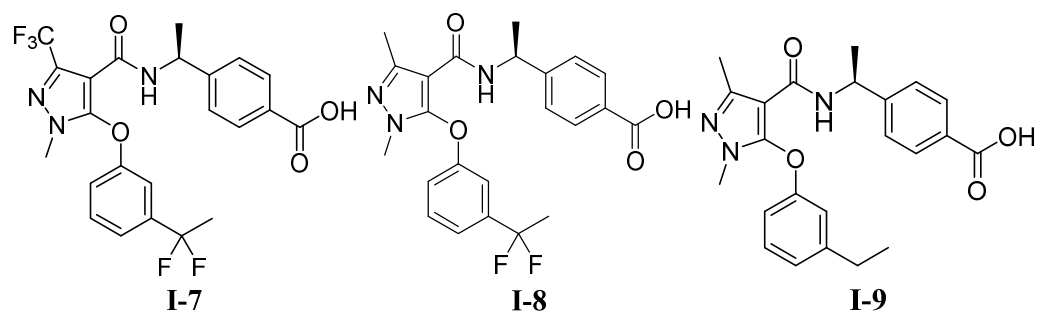
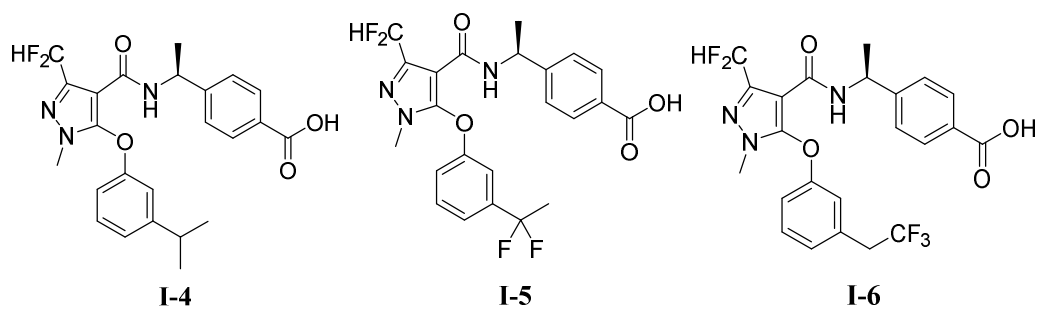
R² is selected from C₂-C₃ alkyl, C₃-C₆ cycloalkyl, C₂-C₃ fluorine-substituted alkyl, and C₃-C₆ fluorine-substituted cycloalkyl, and preferably, R² is selected from -CH₂CH₃, -CH(CH₃)₂, cyclopropyl, -CF₂CH₃, and -CH₂CF₃;

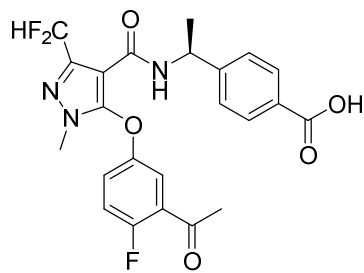
R³ is selected from hydrogen, fluorine, and chlorine;

R⁴ is selected from hydrogen, fluorine, chlorine, C₁-C₄ alkyl, C₁-C₄ alkoxy, C₁-C₄ fluorine- or chlorine-substituted alkyl, and C₁-C₄ fluorine- or chlorine-substituted alkoxy, and preferably, R⁴ is selected from hydrogen, fluorine, and chlorine.

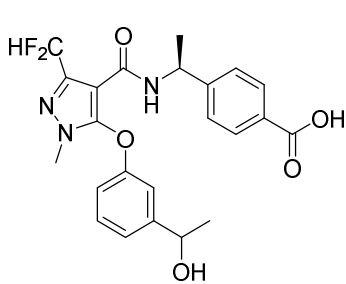
5. A compound, being any one of the following compounds, or being a tautomer, stereoisomer, hydrate, solvate or pharmaceutically acceptable salt of any one of the following compounds:



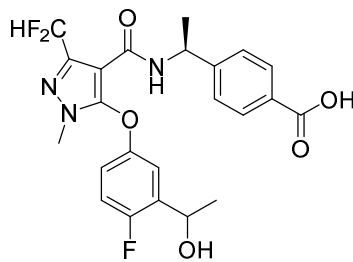




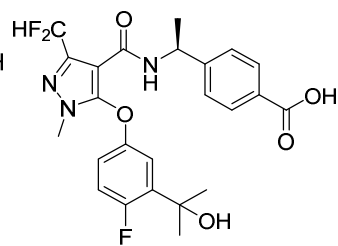
I-21



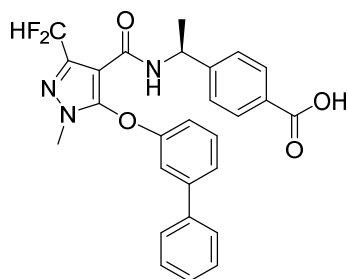
I-22



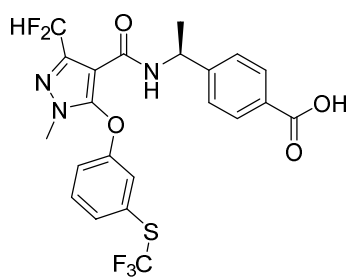
I-23



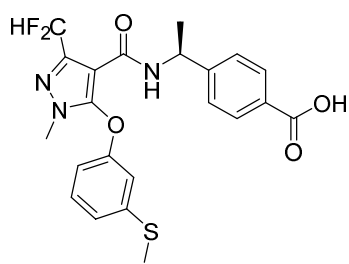
I-24



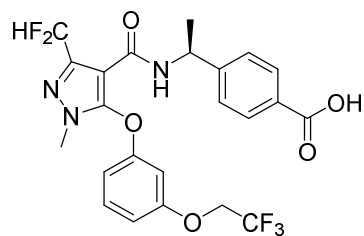
I-27



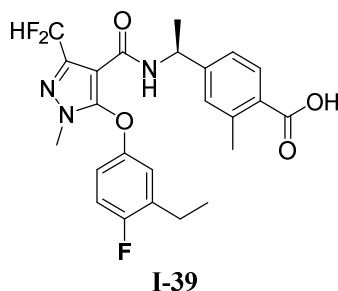
I-29



I-30



I-33



6. A pharmaceutical composition, comprising:

a pharmaceutically acceptable excipient; and

the compound according to any one of claims 1 to 5.

7. Use of the compound according to any one of claims 1 to 5 or the pharmaceutical composition according to claim 6 in the preparation of a medicament for the treatment or prevention of at least one disease selected from the group consisting of an inflammatory disease, a pain, a cancer, a metabolic disease, and a urinary system disease;

wherein the pain is selected from osteoarthritis pain and endometriosis-induced pain, and

wherein the urinary system disease is overactive bladder.

8. The use according to claim 7, wherein the inflammatory disease is selected from arthritis and rheumatoid arthritis.

9. The use according to claim 7, wherein the medicament is adapted to be administered in combination with a radiation therapy and/or an antibody therapy, the antibody therapy being selected from the group consisting of a CTLA4 antibody therapy, a PDL1 antibody therapy, a PD1 antibody therapy, and combinations thereof.

10. The use according to claim 7, wherein the cancer is selected from solid cancers, preferably from breast cancer, cervical cancer, colorectal cancer, endometrial cancer, glioblastoma, head and neck cancer, kidney cancer, liver cancer, lung cancer, medulloblastoma, ovarian cancer, pancreatic cancer, prostate cancer, skin cancer, and urethral cancer.

11. The use according to claim 7, wherein the metabolic disease is diabetes.

12. The compound according to any one of claims 1 to 5 or the pharmaceutical

composition according to claim 6 for use in therapy a disease selected from an inflammatory disease, a pain, a cancer, a metabolic disease, and a urinary system disease;

wherein the pain is selected from osteoarthritis pain and endometriosis-induced pain, and

wherein the urinary system disease is overactive bladder.

13. The compound or the pharmaceutical composition for use according to claim 12, wherein the inflammatory disease is selected from arthritis and rheumatoid arthritis.

14. The compound or the pharmaceutical composition for use according to claim 12, wherein the cancer is selected from solid cancers, preferably from breast cancer, cervical cancer, colorectal cancer, endometrial cancer, glioblastoma, head and neck cancer, kidney cancer, liver cancer, lung cancer, medulloblastoma, ovarian cancer, pancreatic cancer, prostate cancer, skin cancer, and urethral cancer.

15. The compound or the pharmaceutical composition for use according to claim 12, wherein the compound or the pharmaceutical composition is administered in combination with a radiation therapy and/or an antibody therapy, the antibody therapy being selected from the group consisting of a CTLA4 antibody therapy, a PDL1 antibody therapy, a PD1 antibody therapy, and combinations thereof.

16. The compound or the pharmaceutical composition for use according to claim 12, wherein the metabolic disease is diabetes.