

TITLE OF INVENTION

PYRAZOLOPYRIMIDINE DERIVATIVES AS KINASE INHIBITOR

TECHNICAL FIELD

The present invention relates to a pyrazolopyrimidine derivative having kinase inhibitory
5 activity, a process for preparing the same and use thereof.

BACKGROUND OF ART

Protein kinase is an enzyme that catalyzes phosphorylation of specific residues of other
proteins, and plays an important role in signal-transduction pathways that transduce extracellular
signals to the nucleus. Further, it is involved in various diseases in vivo. In the onset or
10 development of inflammatory disease, autoimmune disease, proliferative disease or
hyperproliferative disease, and/or immunity mediated disease, there is various evidence that T-
cells (or T-lymphocytes) and B-cells (or B-lymphocytes) play an important role.

Janus kinase (hereinafter referred to as "JAK") is a cytoplasmic protein tyrosine kinase
15 that plays pivotal roles in regulating cell function in the limbo-hematopoietic system.
Cytokines are known to play an important role in regulating inflammation, immunity and
normal cell function, and JAK activates STAT (Signal Transducer and Activators of
Transcription) proteins through tyrosine phosphorylation to provide rapid signaling pathways to
cytokines. JAK/STAT signaling is known to be associated with allergies, asthma, autoimmune
20 diseases (e.g., transplant rejection, rheumatoid arthritis, amyotrophic lateral sclerosis, multiple
sclerosis etc.), solid cancers, blood cancers (e.g., leukemia, lymphoma and so on).

The JAK family is classified into four members: JAK 1, JAK 2, JAK 3, and TYK 2.
Members of the JAK family pair with each other to mediate signals from a variety of cytokines.
25 It includes JAK2 and JAK1 associated with hematopoietic growth factor signaling, and a
combination of TYK2 and JAK2 is important for interferon signaling and contributes to host
tolerance. JAK2 can induce anemia, thrombocytopenia, leukopenia, especially when it is
involved in the hematopoietic growth factor signaling and causes excessive inhibition.

30 The expression of JAK1, JAK2, and TYK2 was found to be widely distributed,

whereas the expression of JAK3 was restricted to lymphocytes and is associated with signaling for the common gamma chains, members of IL-2, IL-4, IL-7, IL-9, IL-15 and IL-21 receptors, particularly the common gamma chain of the IL-2 family. As soon as the cytokine is bound, the receptor carries adjacent JAK3 nearby, which induces autophosphorylation of the β -chain C-terminus. As a result, it causes activation of the STAT protein, which is an important step in retransmitting the signal to the nucleus. JAK3 controls the signal pathways of various cytokines through this process. This makes JAK3 as an attractive target for immunosuppression.

B cells play an important role in the development of autoimmune and/or inflammatory diseases. Protein-based therapeutic agents that reduce B cells, for example Rituxan, are effective in autoantibody-induced inflammatory diseases such as rheumatoid arthritis. Thus, protein kinase inhibitors that play a role in B cell activation are useful therapeutic agents for the treatment of B cell-mediated diseases, for example, for the production of autoantibodies.

Signal transduction through B cell receptor (BCR) regulates various B cell responses, including proliferation and differentiation into mature antibody-producing cells. BCR is an important regulatory element of B cell activity, and abnormal signal transduction can cause the formation of pathogenic autoantibodies leading to a plurality of autoimmune and/or inflammatory diseases and the proliferation of deregulated B cell.

Bruton's tyrosine kinase (hereinafter, referred to as "BTK") is an important regulator of the development, activation, signaling and survival of B-cells. BTK is involved in signal transduction pathways initiated by binding various extracellular ligands to their cell surface receptors. Following ligation of the B cell antigen receptor (BCR), the activity of BTK by the coincident action of the protein tyrosine kinases Lyn and Syk is required for the induction of the phospholipase C- γ 2-mediated calcium mobilization. Therefore, inhibition of BTK can be a useful therapeutic approach in blocking the onset process of B-cell mediated diseases.

As mentioned above, Janus kinase and TEC-based kinases play an important role in the activation of T-cells and/or B-cells involved in the development of inflammatory diseases,

autoimmune diseases, proliferative diseases or hyperproliferative diseases, and immunity mediated diseases. Therefore, the development of substances that effectively inhibit these diseases can be useful as a related therapeutic agent. Specific examples of the diseases which can be treated and prevented include cancer, transplant rejection, multiple sclerosis, rheumatoid arthritis, psoriatic arthritis, psoriasis, asthma, allergic dermatitis, atopic dermatitis, eczema, type I diabetes, diabetic complication, ulcerative colitis, Crohn's disease, autoimmune thyroid disorder, systemic depilation, Sjogren's syndrome and the like.

JAK3 kinase inhibitor, tofacitinib (CP-690550) (Pfizer Inc.) is currently approved and marketed for the treatment of rheumatoid arthritis. In addition, a BTK kinase inhibitor, ibrutinib (PCI-32765) (Pharmacyclics) is in a clinical stage, but severe side effects such as skin rash and diarrhea have been reported in clinical cases. Thus, there is a need to develop a more stable and effective substance that inhibits JAK and/or BTK (see, Nat Rev Rheumatol. 2009 Jun 5(6) 317-24; Expert Opin Investig Drugs. 2014 Aug 23(8) 1067-77; Drug Discov Today 2014 Aug 19(8) 1200-4; WO2002/096909; WO2010-009342).

Therefore, the present inventors has found a new compound having an excellent inhibitory activity as a kinase inhibitor, thereby completing the present invention. Specifically, the compounds of the present invention show affinity for JAK and/or BTK. The compounds belonging to the present invention themselves have mainly a kinase-inhibitory activity, but do not exclude a possibility of exhibiting a pharmacological action as an efficacious agent by a special body environment or by products of metabolic process, after absorption into the body.

DETAILED DESCRIPTION OF THE INVENTION

TECHNICAL PROBLEM

It is an object of the present invention to provide a pyrazolopyrimidine derivative having kinase inhibitory activity, and a process for preparing the same.

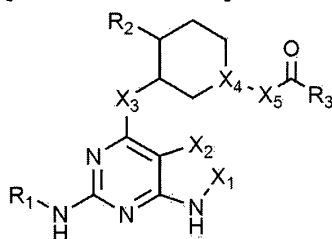
It is another object of the present invention to provide a pharmaceutical composition comprising the pyrazolopyrimidine derivative as an active ingredient.

TECHNICAL SOLUTION

In order to achieve the above objects, the present invention provides a compound

represented by the following Chemical Formula 1, or a pharmaceutically acceptable salt thereof:

[Chemical Formula 1]



in Chemical Formula 1,

5 R_1 is C_{6-10} aryl; or C_{2-10} heteroaryl containing one to three of N, O, or S;

wherein R_1 is unsubstituted, or substituted with a substituent selected from the group consisting of piperazinyl unsubstituted or substituted with C_{1-4} alkyl; benzyl unsubstituted or substituted with C_{1-4} alkoxy; one or two C_{1-4} alkyl unsubstituted or substituted with morpholino, $-N(C_{1-4} \text{ alkyl})_2$, C_{1-4} alkoxy, cyano, or $-CONH(C_{1-4} \text{ alkyl})$; C_{1-4} haloalkyl; C_{3-6} cycloalkyl; 10 morpholino; $-CO-(\text{morpholino})$; morpholino and halogen; $-N(C_{1-4} \text{ alkyl})_2$; $-NHCO(C_{2-4} \text{ alkenyl})$; $-NHCO(\text{pyrrolidinyl})$; C_{1-4} alkoxy unsubstituted or substituted with $-N(C_{1-4} \text{ alkyl})_2$; C_{6-10} aryloxy; pyrazolyl unsubstituted or substituted with one or two C_{1-4} alkyl; pyrrolidinyl; tetrahydropyranyl; and halogen,

R_2 is hydrogen, C_{1-4} alkyl, or halogen;

15 R_3 is C_{1-4} alkyl unsubstituted or substituted with cyano, or halogen; C_{2-6} alkenyl unsubstituted or substituted with one or two substituents independently selected from the group consisting of cyano, C_{3-6} cycloalkyl, and $-N(C_{1-4} \text{ alkyl})_2$; or C_{2-4} alkynyl unsubstituted or substituted with C_{3-6} cycloalkyl,

X_1 is CR_4 or N,

20 wherein R_4 is hydrogen, C_{1-4} alkyl, or halogen,

X_2 is CR_5 , or N,

wherein R_5 is hydrogen, C_{1-4} alkyl, C_{1-4} alkoxy, C_{1-4} haloalkyl, halogen, cyano, or C_{1-4} alkylthio,

X_3 is NR_6 , O, or S,

25 wherein R_6 is hydrogen or C_{1-4} alkyl,

X_4 is CH, or N, and

X₅ is a bond, or NH.

Preferably, R₁ is benzothiazolyl, isothiazolyl, isoxazolyl, phenyl, pyrazolyl, or pyridinyl.

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Preferably, R₁ is phenyl, wherein the R₁ is piperazinyl substituted with C₁₋₄ alkyl; morpholino; -CO-(morpholino); -N(C₁₋₄ alkyl)₂; -NHCO(C₂₋₄ alkenyl); -NHCO(pyrrolidinyl); C₁₋₄ alkoxy substituted with -N(C₁₋₄ alkyl)₂; phenoxy; pyrazolyl unsubstituted or substituted with one or two C₁₋₄ alkyl; or pyrrolidinyl. More preferably, R₃ is C₂₋₄ alkenyl unsubstituted or substituted with cyano, or C₂₋₄ alkenyl unsubstituted or substituted with cyano or -N(C₁₋₄ alkyl)₂.

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Preferably, R₁ is pyrazolyl, wherein the R₁ is benzyl substituted with C₁₋₄ alkoxy; one or two C₁₋₄ alkyl unsubstituted or substituted with morpholino, -N(C₁₋₄ alkyl)₂, C₁₋₄ alkoxy, cyano, or -CONH(C₁₋₄ alkyl); C₁₋₄ haloalkyl; C₃₋₆ cycloalkyl; or tetrahydropyranyl. More preferably, R₃ is C₂₋₄ alkenyl unsubstituted or substituted with cyano or -N(C₁₋₄ alkyl)₂; or C₂₋₄ alkynyl.

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Preferably, R₁ is unsubstituted benzothiazolyl, isothiazolyl substituted with C₁₋₄ alkyl, or unsubstituted isoxazolyl. More preferably, R₃ is C₂₋₄ alkenyl unsubstituted or substituted with cyano or -N(C₁₋₄ alkyl)₂; or C₂₋₄ alkynyl.

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Preferably, R₁ is

unsubstituted benzothiazolyl;

isothiazolyl substituted by C₁₋₄ alkyl;

isoxazolyl unsubstituted or substituted by C₁₋₄ alkyl;

pyrazolyl substituted by benzyl substituted by C₁₋₄ alkoxy, one or two C₁₋₄ alkyl unsubstituted or substituted by morpholino, -N(C₁₋₄ alkyl)₂, C₁₋₄ alkoxy, cyano, or -CONH(C₁₋₄ alkyl), C₁₋₄ haloalkyl, C₃₋₆ cycloalkyl, or tetrahydropyranyl;

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phenyl substituted by piperazinyl substituted by C₁₋₄ alkyl, morpholino, -CO-(morpholino), morpholino and halogen, -N(C₁₋₄ alkyl)₂, -NHCO(C₂₋₄ alkenyl), -NHCO(pyrrolidinyl), C₁₋₄ alkoxy substituted by -N(C₁₋₄ alkyl)₂, phenoxy, pyrazolyl

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unsubstituted or substituted by one or two C₁₋₄ alkyl, or pyrrolidinyl; or
unsubstituted pyridinyl.

Preferably, R₂ is hydrogen, methyl, or fluoro.

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Preferably, R₃ is -CH₂Cl, -CH₂CN, -CH=CH₂, -CH=CHCH₃, -C(CN)=CHCH(CH₃)₂,
-C(CN)=CH(cyclopentyl), -C(CN)=CH(cyclopropyl), -C(CN)=CHC(CH₃)₃,
C(CN)=CHCH(CH₃)₂, -C≡CCH₃, -C≡CCH₂CH₃, or -C≡C-(cyclopropyl).

10 Preferably, X₁ is CH, or N.

Preferably, X₂ is CR₅, and R₅ is hydrogen, methyl, fluoro, chloro, cyano, or methylthio.

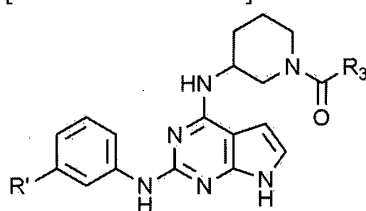
Preferably, X₃ is NH, N(CH₃), S, or O.

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Preferably, X₄ is N, and X₅ is a bond; or X₄ is CH, and X₅ is NH.

Preferably, the compound represented by Chemical Formula 1 is represented by the
following Chemical Formula 1-1:

20 [Chemical Formula 1-1]



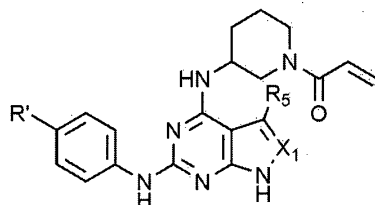
in Chemical Formula 1-1,

R' is -NHCO(C₂₋₄ alkenyl); -NHCO(pyrrolidinyl); or pyrazolyl unsubstituted or
substituted with two C₁₋₄ alkyl,

25 R₃ is C₁₋₄ alkyl unsubstituted or substituted with cyano; or C₂₋₄ alkenyl unsubstituted or
substituted with cyano or -N (C₁₋₄ alkyl).

Preferably, the compound represented by Chemical Formula 1 is represented by the following Chemical Formula 1-2:

[Chemical Formula 1-2]



5 in Chemical Formula 1-2,

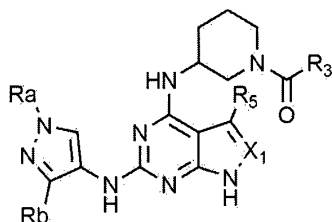
X₁ is CH, or N,

R' is piperazinyl unsubstituted or substituted with C₁₋₄ alkyl; morpholino; -CO-(molpolino); -N(C₁₋₄ alkyl)₂; C₁₋₄ alkoxy unsubstituted or substituted with -N(C₁₋₄ alkyl)₂; phenoxy; pyrazolyl unsubstituted or substituted with one or two C₁₋₄ alkyl; or pyrrolidinyl, and

10 R₅ is hydrogen or halogen.

Preferably, the compound represented by Chemical Formula 1 is represented by the following chemical formula 1-3:

[Chemical Formula 1-3]



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in Chemical Formula 1-3,

X_1 is CH, or N,

Ra is benzyl unsubstituted or substituted with C₁₋₄ alkoxy; C₁₋₄ alkyl unsubstituted or substituted with morpholino, -N(C₁₋₄ alkyl)₂, C₁₋₄ alkoxy, cyano, or -CONH(C₁₋₄ alkyl); C₁₋₄ haloalkyl; C₃₋₆ cycloalkyl; or tetrahydropyranyl,

Rb is hydrogen or C₁₋₄ alkyl,

R₃ is C₂₋₄ alkenyl unsubstituted or substituted with cyano or -N(C₁₋₄ alkyl)₂; or C₂₋₄ alkynyl, and

R₅ is hydrogen or halogen.

Representative examples of the compounds represented by Chemical Formula 1 are as follows:

- 1) (R)-1-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 2) (R)-N-(3-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide,
- 3) (S)-N-(3-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide,
- 4) (R)-1-(3-(2-(4-morpholinophenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 5) (R)-1-(3-(2-(4-(pyrrolidin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 6) (R)-1-(3-(2-(4-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 7) (R)-1-(3-(2-(4-(2-(diethylamino)ethoxy)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 8) (R)-1-(3-(2-(4-(morpholine-4-carbonyl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 9) (R)-1-(3-(2-(4-(dimethylamino)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 10) (R)-1-(3-(2-(3-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 11) (S)-1-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 12) (R)-1-(3-(2-(benzo[d]thiazol-6-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 13) (R)-1-(3-(2-(4-phenoxyphenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 14) (R)-1-(3-(2-(1-methyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-

- ylamino)piperidin-1-yl)prop-2-en-1-one,
- 15) (R)-1-(3-(2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 16) (R)-1-(3-(2-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 17) (R)-1-(3-(2-(1-(2-methoxyethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 18) (R)-1-(3-(2-(1-(2-morpholinoethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 19) (R)-1-(3-(2-(1-(2-(diethylamino)ethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 20) (R)-1-(3-(2-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 21) (R)-1-(3-(2-(1-ethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 22) (R)-1-(3-(2-(1,3-dimethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 23) (R)-1-(3-(2-(isoxazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 24) (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide,
- 25) (R)-1-(3-(2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)but-2-yn-1-one,
- 26) (R)-N-(3-(4-(1-(2-cyanoacetyl)piperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)acrylamide,
- 27) 1-((3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 28) 3-((3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile,
- 29) N-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-

4-ylamino)cyclohexyl)acrylamide,

- 30) (R)-1-(3-(6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 31) (R)-1-(3-(6-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 32) (R)-1-(3-(6-(1-methyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 33) (R)-1-(3-(6-(1-ethyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 34) (R)-1-(3-(6-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 35) (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide,
- 36) (R)-1-(3-(5-chloro-2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 37) (R)-1-(3-(5-chloro-2-(1-ethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 38) (R)-1-(3-(5-chloro-2-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 39) (R)-1-(3-(5-chloro-2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 40) (R)-1-(3-(5-chloro-2-(1-(2-morpholinoethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 41) (R)-1-(3-(5-chloro-2-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 42) (R)-1-(3-(3-chloro-6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 43) (R)-1-(3-(3-chloro-6-(4-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 44) (R)-1-(3-(3-chloro-6-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-1H-

- pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 45) (R)-1-(3-(3-chloro-6-(1-methyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 46) (R)-1-(3-(3-chloro-6-(1-ethyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 5 d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 47) (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide,
- 48) (R)-1-(3-(3-chloro-6-(1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 10 49) (R)-1-(3-(3-chloro-6-(1-cyclopropyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 50) (R)-1-(3-(3-chloro-6-(1-isopropyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 51) (R)-1-(3-(3-chloro-6-(1-propyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 15 d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 52) (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)acetonitrile,
- 53) (R)-1-(3-(6-(1-*tert*-butyl-1H-pyrazol-4-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 20 54) (R)-1-(3-(3-chloro-6-(1-(2-morpholinoethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 55) (R)-1-(3-(3-chloro-6-(1-isobutyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 56) (R)-1-(3-(3-chloro-6-(1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 25 pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 57) (R)-1-(3-(3-chloro-6-(1-(2-(diethylamino)ethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 58) (R)-1-(3-(3-chloro-6-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,
- 30 59) (R)-1-(3-(3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-

- ylamino)piperidin-1-yl)prop-2-en-1-one,
- 60) (R)-2-(4-((4-((1-acryloylpiperidin-3-yl)amino)-5-chloro-7H-pyrrolo[2,3-d]pyrimidin-2-yl)amino)-1H-pyrazol-1-yl)-N-methylacetamide,
- 61) (R)-1-(3-((5-chloro-2-((4-morpholinophenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 5 62) (R)-1-(3-((5-chloro-2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 63) (R)-1-(3-((5-chloro-2-((1-isobutyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 10 64) (R)-1-(3-((3-chloro-6-((1-(difluoromethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 65) (R)-1-(3-((6-((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 66) (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 15 67) (R)-1-(3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 68) (R)-1-(3-((5-chloro-2-((4-(morpholine-4-carbonyl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 20 69) (R)-1-(3-((5-chloro-2-((1-methyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 70) (R)-1-(3-((2-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-5-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 71) (R)-1-(3-((6-((1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 25 72) (R)-1-(3-((6-((1-(2-morpholinoethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 73) (R)-1-(3-((6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 30 74) (R)-1-(3-((6-((1-(2-methoxyethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-

- d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 75) (R)-1-(3-((2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 76) 1-((3S,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one,
- 5 d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one,
- 77) 1-((3S,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one,
- 78) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 10 79) (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile,
- 80) (R)-1-(3-((2-((4-(3,5-dimethyl-1H-pyrazol-1-yl)phenyl)amino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 81) (R)-1-(3-((2-(benzo[d]thiazol-6-ylamino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 15 d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 82) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 83) (R)-1-(3-((5-fluoro-2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 20 84) 1-((3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 85) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 86) 1-((3S,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one,
- 25 d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one,
- 87) (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile,
- 88) (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 30 89) (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-((1-cyclopropyl-1H-pyrazol-4-

- yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 90) (R)-1-(3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 91) (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
5 d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 92) 1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 93) 1-((3R,4R)-3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 10 94) 1-((3R,4R)-3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 95) (R)-1-(3-((5-chloro-2-((1-(2-(diethylamino)ethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 96) (R)-1-(3-((5-chloro-2-(isoxazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-
15 yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 97) (R)-1-(3-((5-chloro-2-((2-fluoro-4-morpholinophenyl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 98) (R)-1-(3-((5-chloro-2-((3-fluoro-4-morpholinophenyl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 20 99) (R)-1-(3-((5-chloro-2-((5-methylisoxazol-4-yl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 100) (R)-1-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one,
- 101) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-
25 yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one,
- 102) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-
yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one,
- 103) (R)-1-(3-((2-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one,
- 30 104) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-

- yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 105) (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 106) 1-((3R,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
5 d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 107) 1-((3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 108) (R)-1-(3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-
pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 109) (R)-1-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-
10 d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 110) (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-
pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 111) (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
15 d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 112) (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-3-chloro-1H-
pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 113) (R)-1-(3-((3-chloro-6-((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 114) (R)-1-(3-((3-chloro-6-((1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-yl)amino)-1H-
20 pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 115) (R)-1-(3-((3-chloro-6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 116) (R)-1-(3-((3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-
25 yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 117) (R)-1-(3-((3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-
yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 118) (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one,
- 119) (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-
30

- d]pyrimidin-4-yl)(methylamino)piperidin-1-yl)prop-2-en-1-one,
- 120) 1-((3R,4R)-3-((3-chloro-6-((1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 121) 1-((3R,4R)-3-((3-chloro-6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 5 122) 1-((3R,4R)-3-((3-chloro-6-((1-(2-(diethylamino)ethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 123) 1-((3R,4R)-3-((3-chloro-6-((1-(2-morpholinoethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 10 124) 1-((3R,4R)-3-((3-chloro-6-((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 125) 1-((3R,4R)-3-((3-chloro-6-((1-(3-methoxybenzyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 126) 2-(4-((4-((3R,4R)-1-acryloyl-4-methylpiperidin-3-yl)amino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-yl)amino)-1H-pyrazol-1-yl)acetonitrile,
- 15 127) 1-((3R,4R)-3-((3-chloro-6-((1-(difluoromethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 128) 1-((3R,4R)-3-((3-chloro-6-((1-methyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 20 129) (R)-1-(3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 130) (R)-1-(3-((6-(isoxazol-4-ylamino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 131) (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methylamino)piperidin-1-yl)prop-2-en-1-one,
- 25 132) 1-((3R,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 133) 1-((3R,4R)-3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 30 one,

- 134) 2-(4-((4-(((3R,4R)-1-acryloyl-4-methylpiperidin-3-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-6-yl)amino)-1H-pyrazol-1-yl)acetonitrile,
- 135) 1-((3R,4R)-3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 5 136) (R)-1-(3-((5-chloro-2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 137) (R)-1-(3-((5-chloro-2-(pyridin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 138) 3-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropanenitrile,
- 10 d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropanenitrile,
- 139) (R)-3-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile,
- 140) (R,E)-1-(3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)but-2-en-1-one,
- 15 141) 1-((R)-3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-cyclopropylprop-2-yn-1-ol,
- 142) 1-((R)-3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)pent-2-yn-1-ol,
- 143) 1-((R)-3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)but-2-yn-1-ol,
- 20 144) (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one,
- 145) 1-((3R,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 25 146) (R)-1-(3-((6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 147) (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one,
- 148) (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 30

- 149) (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 150) (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 5 151) (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one,
- 152) (R)-1-(3-((2-((4-morpholinophenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one,
- 153) (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one,
- 10 154) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 155) (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 15 156) (R,E)-4-((1-(2-cyano-4-methylpent-2-enoyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 157) (R,E)-4-((1-(2-cyano-3-cyclopropylacryloyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile,
- 158) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 20 159) 1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one,
- 160) (E)-2-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 25 161) (E)-2-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 162) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 30

- 163) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4,4-dimethylpent-2-enenitrile,
- 164) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 5 165) (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopentylacrylonitrile,
- 166) (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 167) (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 10 168) (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 169) (E)-2-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carbonyl)-3-
- 15 cyclopropylacrylonitrile,
- 170) (E)-2-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 171) (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile,
- 20 172) (R,E)-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 173) (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)acrylonitrile,
- 25 174) (R,E)-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 175) (R,E)-3-cyclopropyl-2-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile,
- 176) (R,E)-3-cyclopropyl-2-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)acrylonitrile,
- 30

- 177) (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 178) (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 5 179) (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 180) (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-4-methylpent-2-enenitrile,
- 181) (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile,
- 10 182) (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile,
- 183) 2-chloro-1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)ethan-1-one,
- 15 184) (R)-1-(3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 185) (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 186) (R)-1-(3-((6-((1-isobutyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 20 187) (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 188) (R)-1-(3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 25 189) (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 190) (R)-1-(3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 191) (R)-1-(3-((6-((1-methyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one,
- 30

192) (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one, and

193) 1-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one.

5

In addition, the compounds of the present invention may exist in the form of salts, especially pharmaceutically acceptable salts. As salts, salts commonly used in the art, such as acid addition salts formed by pharmaceutically acceptable free acids can be used without limitation. The term "pharmaceutically acceptable salt" as used herein refers to any organic or
10 inorganic addition salt of the compound represented by Chemical Formula 1, whose concentration is relatively non-toxic and harmless to a patient and activates effectively and whose side effects do not degrade the beneficial efficacy of the above compounds.

Pharmaceutically acceptable salts can be obtained by conventional methods using
15 inorganic or organic acids. For example, the pharmaceutically acceptable salt can be prepared by dissolving the compound represented by Chemical Formula 1 in a water-miscible organic solvent. e.g., acetone, methanol, ethanol or acetonitrile, followed by adding an organic acid or an inorganic acid, and filtering and drying the precipitated crystals. Alternatively, it may be prepared by removing a solvent or an excessive amount of acid from the acid-added reaction
20 mixture under reduced pressure, followed by drying the residue, or by adding a different organic solvent and then filtering the precipitated salt. At this time, the preferred salts may include salts derived from hydrochloric acid, hydrobromic acid, sulfuric acid, phosphoric acid, nitric acid, acetic acid, glycolic acid, lactic acid, pyruvic acid, malonic acid, succinic acid, glutaric acid, fumaric acid, malic acid, mandelic acid, tartaric acid, citric acid, ascorbic acid, palmitic acid,
25 maleic acid, hydroxymaleic acid, benzoic acid, hydroxybenzoic acid, phenylacetic acid, cinnamic acid, salicylic acid, methanesulfonic acid, benzenesulfonic acid or toluenesulfonic acid, and the like.

A pharmaceutically unacceptable salt or solvate of the compound of Chemical
30 Formula 1 may be used as an intermediate when preparing the compound of Chemical Formula

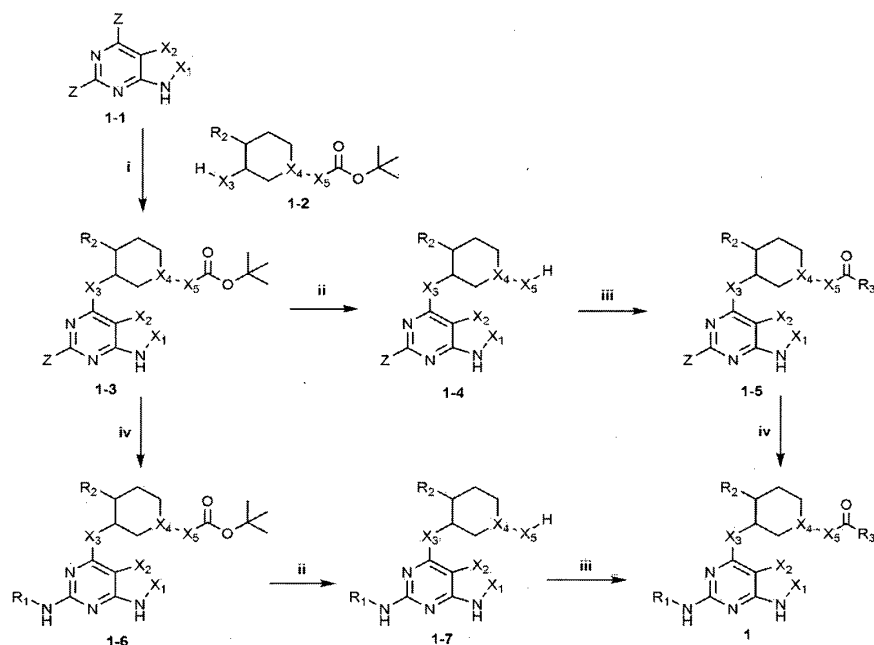
1, or the pharmaceutically acceptable salt or the solvate thereof.

The compound of Chemical Formula 1 according to the present invention includes not only pharmaceutically acceptable salts thereof, but all solvates and hydrates that can be prepared therefrom, and includes all possible stereoisomers as well. The solvate, the hydrate and the stereoisomer of the compound of Chemical Formula 1 may be prepared and used from the compound of Chemical Formula 1 using common methods.

In addition, the compound of Chemical Formula 1 according to the present invention may be prepared either in a crystalline form or in a non-crystalline form, and when the compound of Chemical Formula 1 is prepared in a crystalline form, it may be optionally hydrated or solvated. In the present invention, the compound of Chemical Formula 1 may not only include a stoichiometric hydrate, but include a compound containing various amounts of water. The solvate of the compound of Chemical Formula 1 according to the present invention includes both stoichiometric solvates and non-stoichiometric solvates.

Furthermore, as an example, the present invention can produce the compound represented by Chemical Formula 1 through Reaction Scheme 1 below.

[Reaction Scheme 1]



(in Reaction Scheme 1, X₁ to X₃, and R₁ to R₃ are as previously defined, and when X₄ is N, X₅ is a bond, and when X₄ is CH, X₅ is NH, and Z is halogen. Preferably, Z is chloro)

5 Step i is a step of preparing a compound represented by Chemical Formula 1-3 by reacting a compound represented by Chemical Formula 1-1 with a compound represented by Chemical Formula 1-2. The reaction is preferably carried out in the presence of N,N-diisopropylethylamine at room temperature to high temperature, and the solvent is preferably ethanol.

10

 Step ii is a step of preparing a compound represented by Chemical Formula 1-4 by eliminating a protecting group from a compound represented by Chemical Formula 1-3. The reaction is preferably carried out under a hydrochloric acid condition (preferably, 6 N hydrochloric acid condition), and the solvent is preferably methanol.

15

 Step iii is a step of preparing a compound represented by Chemical Formula 1-5 by reacting a compound represented by Chemical Formula 1-4 with R₃-acyl chloride. The reaction is preferably carried out at -20 °C to 0 °C under conditions of triethylamine or sodium hydrogencarbonate. Further, the solvent is preferably dichloromethane or a mixed solution of

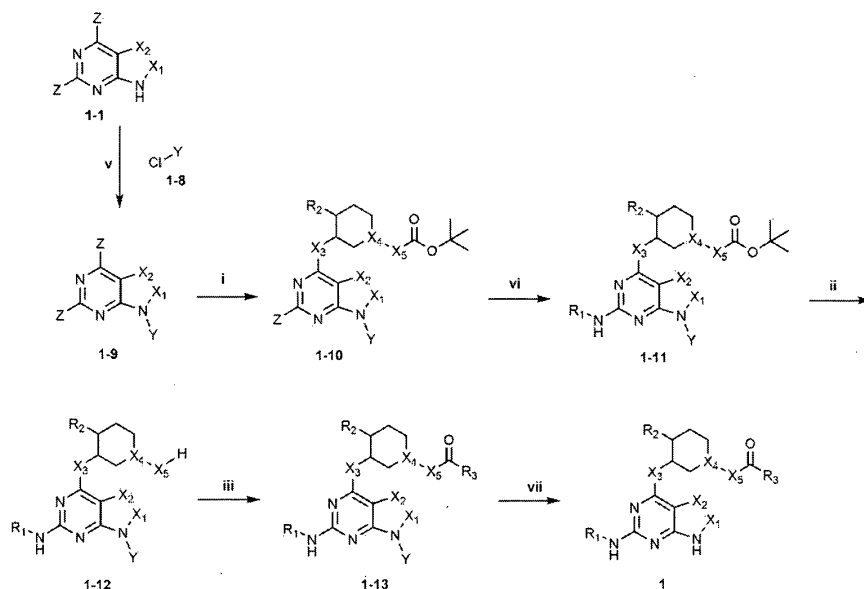
tetrahydrofuran and water. Alternatively, step iii may be carried out by reacting an R₃-carboxylic acid instead of an R₃-acyl chloride. In this case, the reaction is preferably carried out under the conditions of N,N-diisopropylethylamine, 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate at room temperature, and the
5 solvent is preferably tetrahydrofuran or N,N-dimethylformamide.

Step iv is a step of preparing a compound represented by Chemical Formula 1 by reacting a compound represented by Chemical Formula 1-5 with R₁-NH₂. The reaction is preferably carried out at a high temperature under conditions of trifluoroacetic acid, and the
10 solvent is preferably 2-butanol.

In addition, as shown in the above Reaction Scheme 1, starting from the compound represented by Chemical Formula 1-3, the compound represented by Chemical Formula 1-6, the compound represented by Chemical Formula 1-7 and the compound represented by
15 Chemical Formula 1 can be prepared in this order, and each of Steps iv, ii and iii is the same as described above, except for the reactants.

Furthermore, as an example, the present invention can produce the compound represented by Chemical Formula 1 through Reaction Scheme 2 below.

20 [Reaction Scheme 2]



(in Reaction Scheme 2, X₁ to X₃, and R₁ to R₃ are as previously defined, and when X₄ is N, X₅ is a bond, and when X₄ is CH, X₅ is NH. Y is 4-methylbenzyl sulfonyl or 2-(trimethylsilyl)ethoxymethyl, Z is halogen, and preferably Z is chloro)

5

Step v is a step of preparing a compound represented by Chemical Formula 1-9 by reacting a compound represented by Chemical Formula 1-1 with a compound represented by Chemical Formula 1-8. The reaction is preferably carried out at a temperature of 0 °C or less under a basic condition, and the solvent is preferably acetone or N,N-dimethylformamide.

10

Step i is a step for preparing a compound represented by Chemical Formula 1-10 from the compound represented by Chemical Formula 1-9, which is the same as Step i of Reaction Scheme 1 except for the reactants.

15

Step vi is a step of preparing a compound represented by Chemical Formula 1-11 by reacting a compound represented by Chemical Formula 1-10 with R₁-NH₂. The reaction is preferably carried out at a temperature of 100 °C to 120 °C under conditions of a ligand, a palladium catalyst and a base, and the solvent is preferably 1,4-dioxane.

20

Step ii is a step for preparing a compound represented by Chemical Formula 1-12 from

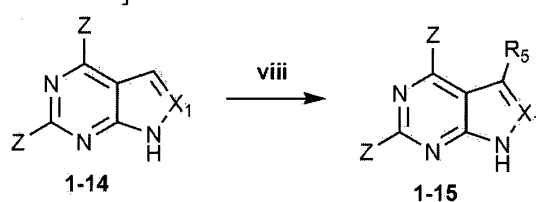
the compound represented by Chemical Formula 1-11, which is the same as Step ii of Reaction Scheme 1 except for the reactants.

Step iii is a step for preparing a compound represented by Chemical Formula 1-13
5 from the compound represented by Chemical Formula 1-12, which is the same as Step iii of
Reaction Scheme 1 except for the reactants.

Step vii is a step for preparing a compound represented by Chemical Formula 1 eliminating Y from a compound represented by Chemical Formula 1-13. When Y is 4-methylbenzylsulfonyl, the reaction is preferably carried out at a temperature of 40 °C to 60 °C under a basic condition, and the solvent is preferably methanol. Further, when Y is 2-(trimethylsilyl)ethoxymethyl, the reaction is preferably carried out at room temperature under trifluoroacetic acid conditions. The solvent is preferably dichloromethane.

15 In addition, the present invention provides a process for preparing a compound represented by the following chemical formula 1-15 as shown in the following Reaction Scheme 3 when X_2 is CR_5 and R_5 is halogen. The compound represented by Chemical Formula 1-15 can be used as the compound represented by Chemical Formula 1-1 in Reaction Schemes 1 and 2.

20 [Reaction Scheme 3]



(in Reaction Scheme 3, X_1 and R_5 are as previously defined, and Z is halogen. Preferably, Z is chloro)

25 Step viii is a step for preparing a compound represented by Chemical Formula 1-15 from the compound represented by Chemical Formula 1-14. The reaction is preferably carried out at room temperature to 60 °C in the presence of N-halosuccinimide, and the solvent is

preferably N,N-dimethylformamide.

Further, the present invention provides a pharmaceutical composition for preventing or treating diseases which are associated with kinase inhibitory actions, comprising the compound
5 represented by Chemical Formula 1, or a pharmaceutically acceptable salt, hydrate, solvate or isomer thereof as an active ingredient.

In this case, the diseases which are associated with kinase inhibitory actions includes inflammatory diseases, autoimmune diseases, proliferative diseases or hyperproliferative
10 diseases, and immunity mediated diseases.

As used herein, the term "prevention" refers to any act to delay or inhibit occurrence, spread or recurrence of the above-mentioned diseases by administration of the composition of the present invention, and "treatment" refers to any act to improve or change the symptoms of
15 the above diseases for the better by administration of the composition of the present invention.

The pharmaceutical composition according to the present invention can be formulated in types for oral or parenteral administrations according to a standard pharmaceutical practice. These formulations may contain additives such as pharmaceutically acceptable carrier, adjuvant
20 or diluent in addition to the active ingredient.

Suitable carriers include, for example, physiological saline, polyethylene glycol, ethanol, vegetable oil, and isopropyl myristate and the like. Diluents include, for example, lactose, dextrose, sucrose, mannitol, sorbitol, cellulose and/or glycine and the like, but are not
25 limited thereto. Further, the compounds of the present invention can be dissolved in oils, propylene glycol or other solvents commonly used in the preparation of injection solutions. Furthermore, the compounds of the present invention can be formulated in ointments or creams for topical application.

30 Pharmaceutical dosage forms of the compounds of the present invention may include

using the compounds in the form of pharmaceutically acceptable salts or solvates thereof, and using the compounds alone or as a combination and/or a suitable mixture together with other pharmaceutically active compounds.

5 The compounds of the present invention can be formulated into injection solutions by dissolving, suspending or emulsifying the compounds in a water-soluble solvent such as normal saline, 5% dextrose or a non-aqueous solvent such as synthetic fatty acid glyceride, higher fatty acid ester or propylene glycol. Formulations of the present invention may include conventional additives such as solubilizers, isotonic agents, suspending agents, emulsifying agents, stabilizers
10 and preservatives.

A preferred dose of the compound of the present invention may be varied according to the condition and weight of a patient, the severity of a disease, the type of a drug, and the route and duration of administration, but it may be suitably selected by those skilled in the art. In order
15 to achieve the desirable effects, however, the compound of the present invention may be administrated daily at a dose of 0.0001 to 100 mg/kg (body weight), and preferably 0.001 to 100 mg/kg (body weight). The administration may be performed once a day or in divided doses each day through an oral or parenteral route. Depending on the method of administration, the composition may contain the compound of the present invention in an amount of 0.001 to 99%
20 by weight, preferably 0.01 to 60% by weight.

The pharmaceutical composition according to the present invention may be administered to mammals such as a rat, a mouse, a domestic animal, a human, through various routes. The administration may be carried out through all possible methods, for example, oral,
25 rectal, intravenous, intramuscular, subcutaneous, intra-endometrial, intracerebroventricular injection.

ADVANTAGEOUS EFFECTS

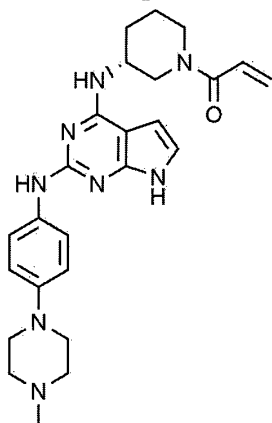
The compound represented by Chemical Formula 1 according to the present invention or a pharmaceutically acceptable salt, hydrate, solvate or isomer thereof can be usefully used for
30 the prevention or treatment of diseases which are associated with kinase inhibitory actions.

DETAILED DESCRIPTION OF THE EMBODIMENTS

Below, the present invention will be described in more detail by way of examples. However, these examples are provided for illustrative purposes only, and should not be construed as limiting the scope of the present invention to these examples.

5

Preparation Example 1: Preparation of 1(R)-1-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of *tert*-butyl (R)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (500.0 mg, 2.7 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695.0 μ L, 4.0 mmol) and *tert*-butyl-(R)-3-aminopiperidine-1-carboxylate (639.3 mg, 4.0 mmol) was added thereto. After stirring the reaction mixture at 110 °C for 12 hours, the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (900.0 mg, yield: 98.0%).

¹H NMR (500MHz, CDCl₃) δ 7.06(s, 1H), 6.38(s, 1H), 4.24-4.20(m, 1H), 3.84-3.82(m, 1H), 3.80-3.30(m, 3H), 2.01-1.90(m, 1H), 1.80-1.75(m, 1H), 1.65-1.55(m, 2H), 1.43(s, 9H)

20

Step 2: Preparation of (R)-2-chloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride

To *tert*-butyl (R)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-

carboxylate (474.0 mg, 1.4 mmol), 6 N hydrochloric acid solution (5.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 7.11(d, 1H), 6.63(d, 1H), 4.52-4.49(m, 1H), 3.66-3.63(m, 1H), 3.37-3.34(m, 1H), 3.02-2.90(m, 2H), 2.19-2.16(m, 1H), 2.12-2.09(m, 1H), 1.90-1.77(m, 2H)

Step 3: Preparation of (R)-1-(3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-2-chloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride(500.0 mg, 1.7 mmol) was dissolved in dichloromethane (10.0 mL), triethylamine (725.3 μL, 5.2 mmol) was added at 0 °C and then the mixture was stirred for 30 minutes. Acryloyl chloride (155.0 μL, 1.9 mmol) was added to the reaction mixture, following by stirring at 0 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (250.0 mg, yield: 66.0%).

¹H NMR (500MHz, CD₃OD) δ 7.02(s, 1H), 6.92-6.86(m, 1H), 6.57(s, 1H), 5.73-5.68(m, 1H), 4.60-4.28(m, 2H), 4.06-4.02(m, 1H), 3.25-2.96(m, 2H), 2.14-2.12(m, 1H), 1.92-1.88(m, 1H), 1.81-1.78(m, 1H), 1.67-1.63(m, 1H)

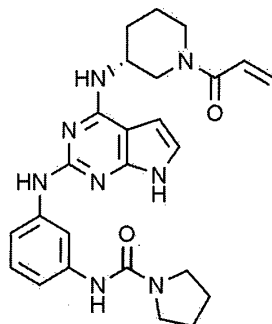
Step 4: Preparation of (R)-1-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

(R)-1-(3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-one (27.0 mg, 0.09 mmol) and 4-(4-methylpiperazin-1-yl)aniline (11.5 mg, 0.06 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (6.9 μL, 0.07 mmol) was added thereto and the reaction mixture was reacted at 110 °C for 12 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to obtain a title compound (2.3 mg, yield: 6.4%)

¹H NMR (500MHz, CD₃OD) δ 7.55-7.52(m, 2H), 6.92-90(m, 2H), 6.84-6.50(m, 2H,

6.41(s, 1H), 6.28-6.03(m, 1H), 5.80-5.46(m, 1H), 4.20-4.06(m, 3H), 3.16-3.13(m, 5H), 2.67-2.63(m, 5H), 2.38(s, 3H), 2.15-2.02(m, 1H), 1.91-1.89(m, 1H), 1.72-1.59(m, 2H)

Example 2: Preparation of (R)-N-(3-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide,



Step 1: Preparation of 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (3.0 g, 16.0 mmol) was dissolved in acetone (20.0 mL), 4-methylbenzenesulfonyl chloride (4.6 g, 23.9 mmol) was added thereto. After cooling to 0 °C, 2 M sodium hydroxide solution (12.0 mL) was slowly added dropwise and then stirred at room temperature for 2 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (2.9 g, yield: 80.0%).

¹H NMR (500MHz, CD₃OD) δ 8.12(d, 2H), 7.76(d, 1H), 7.37(d, 2H), 6.68(d, 1H), 2.43(s, 3H)

Step 2: Preparation of *tert*-butyl-(R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 2,4-dichloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine (500.0 mg, 1.5 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (382.0 μL, 2.2 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (322.0 mg, 1.6 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (681.0 mg, yield: 92.0%).

¹H NMR (500MHz, CDCl₃) δ 8.10(d, 1H), 7.39(d, 1H), 7.31-7.23(m, 4H), 4.17-

4.13(m, 1H), 3.70-3.60(m, 1H), 3.45-3.35(m, 3H), 2.40(s, 3H), 1.95-1.85(m, 1H), 1.70-1.65(m, 1H), 1.60-1.55(m, 2H), 1.40-1.37(m, 9H)

Step 3: Preparation of N-(3-nitrophenyl)pyrrolidine-1-carboxamide

5 After 1-isocyanato-3-nitrobenzene (1.0 g, 6.1 mmol) was dissolved in tetrahydrofuran (10.0 mL), pyrrolidine (0.6 mL, 7.3 mmol) was added thereto. The reaction mixture was allowed to react for 1 hour and then the solvent was concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (1.3 g, yield: 89.0%).

¹H NMR (500MHz, CD₃OD) δ 8.47(s, 1H), 7.85-7.81(m, 2H), 7.49(t, 1H), 3.49-
10 3.47(m, 4H), 1.98(s, 4H)

Step 4: Preparation of N-(3-aminophenyl)pyrrolidine-1-carboxamide

After N-(3-nitrophenyl)pyrrolidine-1-carboxamide(1.3 g, 5.5 mmol) was dissolved in methanol (30.0 mL), excess Raney nickel was added thereto and replaced with hydrogen gas. The
15 reaction mixture was stirred for 1 hour, filtered through celite, and the filtrate was concentrated to obtain a title compound (1.1 g, yield: 100.0%).

¹H NMR (500MHz, CD₃OD) δ 6.88-6.85(m, 1H), 6.74(s, 1H), 6.61(d, 1H), 6.31(d, 1H), 3.18(s, 4H), 1.83(s, 4H)

20 **Step 5: Preparation of *tert*-butyl-(*R*)-3-((2-((3-(pyrrolidin-1-carboxamido)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate**

After *tert*-butyl-(*R*)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-carboxylate (200.0 mg, 0.4 mmol) was dissolved in anhydrous 1,4-
25 dioxane (2.0 mL), N-(3-aminophenyl)pyrrolidine-1-carboxamide (73.8 mg, 0.4 mmol), palladium acetate (4.0 mg, 0.02 mmol), 4,5-*bis*-9,9-dimethylxanthene (20.8 mg, 0.04 mmol) and cesium carbonate (234.1 mg, 0.7 mmol) were added thereto. The reaction mixture was reacted at 120 °C for 30 minutes using microwave. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure.
30 The residue was isolated by column chromatography to obtain a title compound (190.0 mg,

yield: 78.2%).

¹H NMR (500MHz, CDCl₃) δ 8.57(s, 1H), 7.98(d, 1H), 7.59-7.55(m, 1H), 7.22-7.18(m, 2H), 6.91-6.89(m, 2H), 6.51(s, 1H), 6.30(s, 1H), 3.72-3.69(m, 1H), 3.47-3.39(m, 7H), 2.33(m, 3H), 1.95-1.90(m, 6H), 1.69-1.64(m, 2H), 1.59-1.56(m, 1H), 1.44-1.36(m, 9H)

5

Step 6: Preparation of (R)-N-(3-((4-(piperidin-3-ylamino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-2-yl)amino)phenyl) pyrrolidine-1-carboxamide hydrochloride

To *tert*-butyl-(R)-3-((2-((3-(pyrrolidin-1-carboxamido)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-carboxylate (190.0 mg, 0.3 mmol), 6 N hydrochloric acid solution (5.0 mL, excessive amount) was added. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 8.38(s, 1H), 8.02(d, 2H), 7.47-7.44(m, 2H), 7.38-7.36(m, 1H), 7.32-7.28(m, 1H), 6.98-6.86(m, 3H), 3.69-3.66(m, 1H), 3.48-3.39(m, 5H), 2.98-2.94(m, 1H), 2.82-2.80(m, 1H), 2.43(s, 3H), 2.16-2.14(m, 1H), 2.06-1.98(m, 7H), 1.79-1.75(m, 1H)

15

Step 7: Preparation of (R)-N-(3-((4-(1-acryloylpiperidin-3-yl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-2-yl)amino)phenyl)pyrrolidine-1-carboxamide

After (R)-N-(3-((4-(piperidin-3-ylamino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-2-yl)amino)phenyl) pyrrolidin-1-carboxamide hydrochloride (160.8 mg, 0.3 mmol) was dissolved in dichloromethane (10.0 mL), triethylamine (110.0 μL, 0.8 mmol) was added thereto at 0 °C and stirred for 30 minutes. Acryloyl chloride (60.6 μL, 0.4 mmol) was added to the reaction mixture, followed by stirring at 0 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (250.0 mg, yield: 66.9%).

25

¹H NMR (500MHz, CD₃OD) δ 8.13(s, 1H), 7.96(d, 1H), 7.31-6.99(m, 5H), 6.85-6.45(m, 2H), 6.30-5.94(m, 1H), 6.80-5.32(m, 1H), 4.30-4.07(m, 2H), 3.45(s, 4H), 3.20-3.16(m, 2H), 3.10-2.70(m, 1H), 2.33(s, 3H), 2.15-2.10(m, 1H), 1.94(s, 4H), 1.89-1.87(m, 1H), 1.75-1.57(m, 2H)

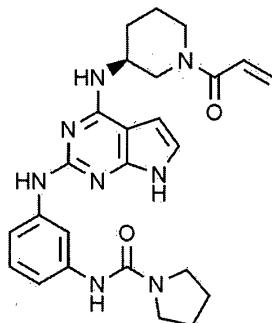
30

Step 8: Preparation of (R)-N-(3-((4-(1-acryloylpiperidin-3-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide

(R)-N-(3-((4-(1-acryloylpiperidin-3-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide (25.0 mg, 0.04 mmol) was dissolved in methanol (1.0 mL). Potassium hydroxide (4.5 mg, 0.08 mmol) was added to the reaction mixture and stirred at 50 °C for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (2.0 mg, yield: 11.1%).

¹H NMR (500MHz, CD₃OD) δ 8.09(s, 1H), 7.18-7.12(m, 3H), 7.10-6.43(m, 2H), 6.43(s, 1H), 6.31-6.00(m, 1H), 5.81-5.41(m, 1H), 4.21-4.16(m, 1H), 3.52-3.41(m, 4H), 3.43-3.17(m, 2H), 3.19-3.00(m, 1H), 2.89-2.76(m, 1H), 2.11-2.07(m, 1H), 1.93-1.92(m, 4H), 1.82-1.80(m, 1H), 1.79-1.59(m, 2H)

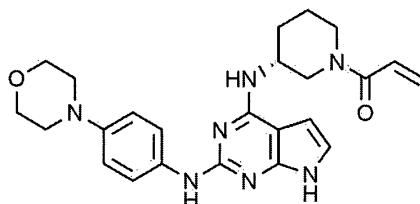
Example 3: Preparation of (S)-N-(3-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)pyrrolidine-1-carboxamide,



A title compound (10 mg, yield: 25.3%) was prepared in the same manner as in Example 2, except that *tert*-butyl-(*R*)-3-aminopiperidine-1-carboxylate was used instead of *tert*-butyl-(*S*)-3-aminopiperidine-1-carboxylate in Example 2.

¹H NMR (500MHz, CD₃OD) δ 8.09(s, 1H), 7.18-7.12(m, 3H), 7.10-6.43(m, 2H), 6.43(s, 1H), 6.31-6.00(m, 1H), 5.81-5.41(m, 1H), 4.21-4.16(m, 1H), 3.52-3.41(m, 4H), 3.43-3.17(m, 2H), 3.19-3.00(m, 1H), 2.89-2.76(m, 1H), 2.11-2.07(m, 1H), 1.93-1.92(m, 4H), 1.82-1.80(m, 1H), 1.79-1.59(m, 2H)

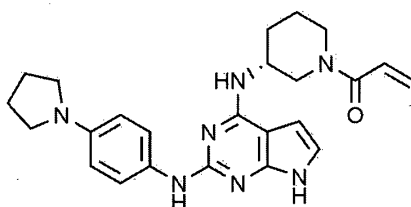
Example 4: Preparation of (R)-1-(3-(2-(4-morpholinophenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one,



5 A title compound (9.6 mg, yield: 20.0%) was prepared in the same manner as in Example 1, except that 4-morpholinoaniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.55-7.50(m, 2H), 6.90-6.88(m, 2H), 6.74-6.63(m, 2H), 6.54(s, 1H), 6.41-6.02(m, 1H), 5.80-5.45(m, 1H), 4.21-4.05(m, 2H), 3.83-3.81(m, 4H),
10 3.05-3.03(m, 4H), 2.79-2.73(m, 3H), 2.15-2.10(m, 1H), 1.94-1.90(m, 1H), 1.75-1.59(m, 2H)

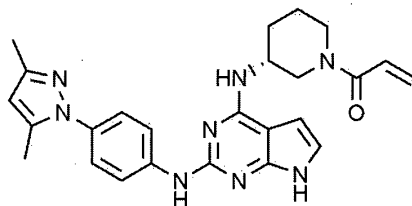
Example 5: Preparation of (R)-1-(3-(2-(4-(pyrrolidin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



15 A title compound (5.6 mg, yield: 11.9%) was prepared in the same manner as in Example 1, except that 4-(pyrrolidin-1-yl) aniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.45-7.36(m, 2H), 6.85-6.50(m, 4H), 6.47(s, 1H),
20 6.39-6.05(m, 1H), 5.85-5.54(m, 1H), 4.21-4.08(m, 2H), 3.26-3.05(m, 6H), 2.20-2.13(m, 1H), 2.00-1.98(m, 4H), 1.92-1.90(m, 1H), 1.76-1.58(m, 3H)

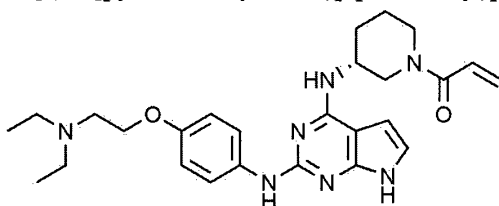
Example 6: Preparation of (R)-1-(3-(2-(4-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (17.3 mg, yield: 34.0%) was prepared in the same manner as in Example 1, except that 4-(3,5-dimethyl-1H-pyrazol-1-yl)aniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

5 ^1H NMR (500MHz, CD_3OD) δ 7.86-7.84(m, 2H), 7.24-7.23(m, 2H), 7.04-6.68(m, 2H), 6.44(s, 1H), 6.24-6.01(m, 2H), 5.76-5.50(m, 1H), 4.35-4.11(m, 2H), 3.50-2.74(m, 3H), 2.20(s, 6H), 2.00-1.92(m, 1H), 1.87-1.60(m, 3H)

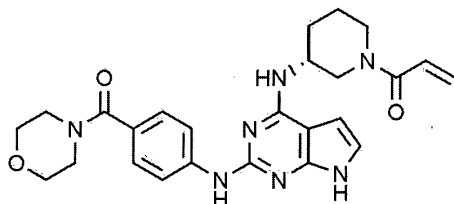
Example 7: Preparation of (R)-1-(3-(2-(4-(2-(diethylamino)ethoxy)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (10.0 mg, yield: 19.0%) was prepared in the same manner as in Example 1, except that 4-(2-(diethylamino)ethoxy)aniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

15 ^1H NMR (500MHz, CD_3OD) δ 7.54-7.50(m, 2H), 6.85-6.82(m, 2H), 6.74-6.50(m, 2H), 6.40(s, 1H), 6.30-6.06(m, 1H), 5.80-5.46(m, 1H), 4.21-4.06(m, 4H), 3.20-3.10(m, 1H), 2.92-2.90(m, 3H), 2.72-2.66(m, 5H), 2.20-2.15(m, 1H), 1.91-1.88(m, 1H), 1.85-1.59(m, 2H), 1.12-1.05(m, 6H)

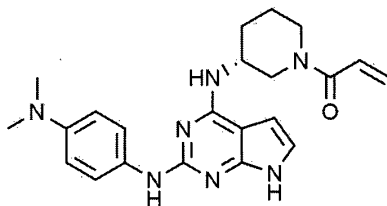
20 **Example 8: Preparation of (R)-1-(3-(2-(4-(morpholine-4-carbonyl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (6.9 mg, yield: 13.3%) was prepared in the same manner as in Example 1, except that (4-aminophenyl)(morpholino)methanone was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.84-7.79(m, 2H), 7.34-7.33(m, 2H), 6.82-6.80(m, 1H), 6.64-6.55(m, 1H), 6.44(s, 1H), 6.27-6.03(m, 1H), 5.80-5.46(m, 1H), 4.40-4.16(m, 2H), 3.68-3.57(m, 9H), 3.16-3.11(m, 1H), 2.90-2.73(m, 1H), 2.20-2.17(m, 1H), 2.15-1.93(m, 1H), 1.85-1.61(m, 2H)

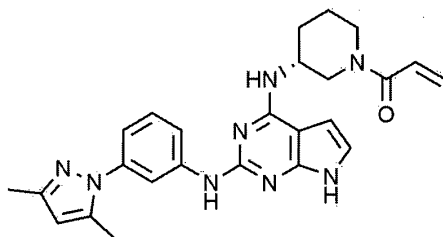
Example 9: Preparation of (R)-1-(3-(2-(4-(dimethylamino)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (8.1 mg, yield: 18.4%) was prepared in the same manner as in Example 1, except that N,N-dimethylbenzene-1,4-diamine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.48-7.43(m, 2H), 6.83-6.47(m, 4H), 6.40(s, 1H), 6.27-6.02(m, 1H), 5.78-5.46(m, 1H), 4.21-4.10(m, 2H), 3.26-3.07(m, 2H), 2.84(s, 6H), 2.80-2.74(m, 1H), 2.14-2.13(m, 1H), 1.94-1.90(m, 1H), 1.62-1.55(m, 2H)

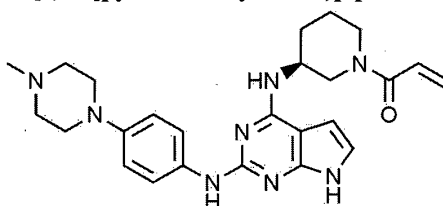
Example 10: Preparation of (R)-1-(3-(2-(3-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (9.8 mg, yield: 19.6%) was prepared in the same manner as in Example 1, except that 3-(3,5-dimethyl-1H-pyrazol-1-yl) aniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

5 ^1H NMR (500MHz, CD_3OD) δ 8.19-8.17(m, 1H), 7.60-7.50(m, 1H), 7.35-7.31(m, 1H), 6.90-6.88(m, 1H), 6.64-6.55(m, 2H), 6.44-6.42(m, 1H), 6.23-6.03(m, 2H), 5.85-5.55(m, 1H), 4.36-4.00(m, 2H), 3.67-3.40(m, 1H), 3.16-3.13(m, 1H), 2.77-2.74(m, 1H), 2.28(s, 3H), 2.24(s, 3H), 2.18-2.05(m, 1H), 1.86-1.48(m, 3H)

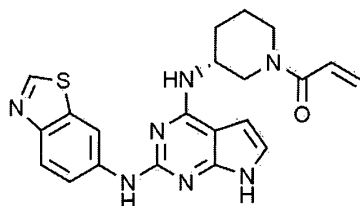
10 **Example 11: Preparation of (S)-1-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (9.8 mg, yield: 19.6%) was prepared in the same manner as in Example 1, except that *tert*-butyl-(*R*)-3-aminopiperidine-1-carboxylate was used instead of *tert*-butyl (S)-3-aminopiperidine-1-carboxylate in Example 1.

15 ^1H NMR (500MHz, CD_3OD) δ 7.58-7.51(m, 2H), 6.92-6.86 (m, 2H), 6.50-6.41(m, 2H), 6.28-6.03 (m, 1H), 5.80-5.46(m, 1H), 4.20-4.06(m, 2H), 3.72-3.66 (m, 2H), 3.12(m, 4H), 2.65 (m, 4H), 2.36(s, 3H), 2.15-2.02(m, 1H), 1.91-1.89(m, 2H), 1.72-1.59(m, 2H)

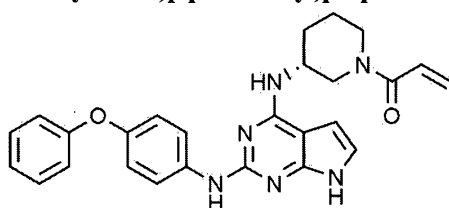
20 **Example 12: Preparation of (R)-1-(3-(2-(benzo[d]thiazol-6-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (7.5 mg, yield: 29.8%) was prepared in the same manner as in Example 1, except that benzo[d]thiazol-6-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

5 ^1H NMR (500MHz, CD_3OD) δ 8.97(s, 1H), 8.80(d, 1H), 7.87(d, 1H), 7.62-7.56(m, 1H), 6.81-6.43(m, 3H), 6.27-5.93(m, 1H), 5.80-5.33(m, 1H), 4.28-4.23(m, 2H), 3.26-2.74(m, 3H), 2.20-2.14(m, 1H), 1.94-1.93(m, 1H), 1.68-1.63(m, 2H)

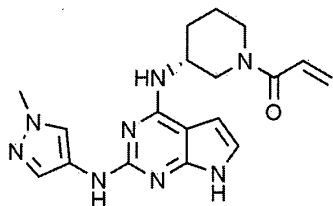
Example 13: Preparation of (R)-1-(3-(2-(4-phenoxyphenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (6.0 mg, yield: 12.2%) was prepared in the same manner as in Example 1, except that 4-phenoxyaniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

15 ^1H NMR (500MHz, CD_3OD) δ 7.37-7.35(m, 2H), 7.29-7.28(m, 2H), 7.12-7.10(m, 2H), 7.03-6.98(m, 3H), 6.71-6.49(m, 2H), 6.37-6.10(m, 1H), 5.70-5.58(m, 1H), 4.22-4.19(m, 1H), 3.28-3.25(m, 1H), 2.89-2.78(m, 1H), 2.59-2.54(m, 2H), 2.07-2.05(m, 1H), 1.85-1.81(m, 1H), 1.79-1.53(m, 2H)

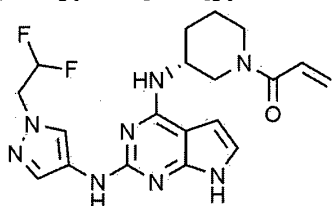
Example 14: Preparation of (R)-1-(3-(2-(1-methyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (10.0 mg, yield: 28.6%) was prepared in the same manner as in Example 1, except that 1-methyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

5 ^1H NMR (500MHz, CD_3OD) δ 7.89(d, 1H), 7.59(d, 1H), 6.84-6.52(m, 2H), 6.40(s, 1H), 6.25-6.07(m, 1H), 5.78-5.50(m, 1H), 4.22-4.16(m, 2H), 3.81(s, 3H), 3.27-2.80(m, 2H), 2.17-2.14(m, 1H), 1.93-1.91(m, 1H), 1.76-1.61(m, 2H)

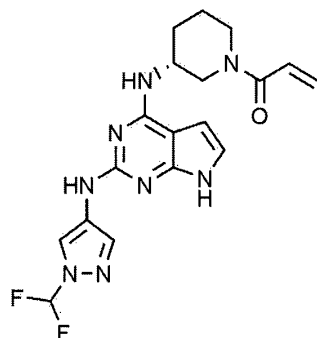
Example 15: Preparation of (R)-1-(3-(2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (14.0 mg, yield: 30.9%) was prepared in the same manner as in Example 1, except that 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

15 ^1H NMR (500MHz, CD_3OD) δ 8.12-7.99(m, 1H), 7.61-7.52(m, 1H), 6.83-6.52(m, 2H), 6.40(s, 1H), 6.26-6.05(m, 2H), 5.79-5.50(m, 1H), 4.48-4.43(m, 2H), 4.24-4.21(m, 1H), 4.11-4.08(m, 2H), 3.16-3.11(m, 1H), 2.16-2.13(m, 1H), 1.95-1.89(m, 1H), 1.73-1.61(m, 2H)

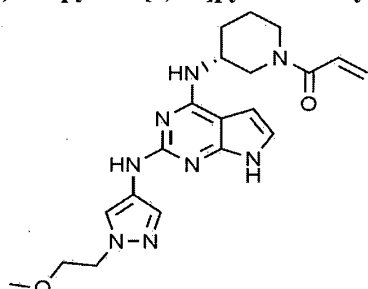
Example 16: Preparation of (R)-1-(3-(2-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.0 mg, yield: 24.2%) was prepared in the same manner as in Example 1, except that 1-(difluoromethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 8.39(d, 1H), 7.76(d, 1H), 7.36-7.34(m, 2H), 7.22-7.02(m, 1H), 6.77-6.57(m, 2H), 6.41(s, 1H), 6.25-6.06 (m, 1H), 5.78-5.52(m, 1H), 4.12-4.08(m, 2H), 3.16-3.14(m, 1H), 2.72-2.65(m, 1H), 2.19-2.17(m, 1H), 1.96-1.92(m, 1H), 1.82-1.73(m, 1H), 1.64-1.61(m, 2H)

Example 17: Preparation of (R)-1-(3-(2-(1-(2-methoxyethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

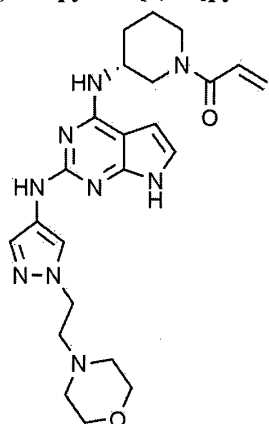


A title compound (6.5 mg, yield: 27.1%) was prepared in the same manner as in Example 1, except that 1-(2-methoxyethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 8.00(d, 1H), 7.55(d, 1H), 6.74-6.52(m, 2H), 6.41(s, 1H), 6.26-6.06(m, 1H), 5.78-5.51(m, 1H), 4.23-4.19(m, 3H), 4.15-4.10(m, 1H), 3.71-3.67 (m, 4H), 3.35 (s, 3H), 3.16-3.14(m, 1H), 2.18-2.17(m, 1H), 1.96-1.93(m, 1H), 1.65-1.63(m, 2H)

Example 18: Preparation of (R)-1-(3-(2-(1-(2-morpholinoethyl)-1H-pyrazol-4-

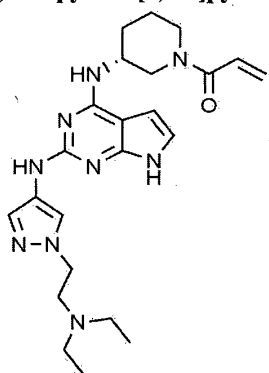
ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (17.5 mg, yield: 34.3%) was prepared in the same manner as in Example 1, except that 1-(2-morpholinoethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.98(d, 1H), 7.55(d, 1H), 6.73-6.56(m, 2H), 6.41(s, 1H), 6.40-6.07(m, 1H), 5.78-5.51(m, 1H), 4.23-4.18(m, 4H), 3.65-3.64(m, 4H), 3.30-3.16 (m, 1H), 2.78-2.71(m, 3H), 2.46 (s, 4H), 2.18-2.13(m, 1H), 1.94-1.92(m, 1H), 1.62-1.60(m, 2H)

Example 19: Preparation of (R)-1-(3-(2-(1-(2-(diethylamino)ethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

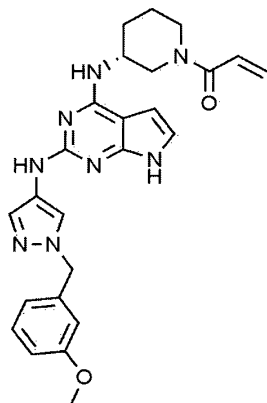


A title compound (4.5 mg, yield: 10.0%) was prepared in the same manner as in Example 1, except that 1-(2-(diethylamino)ethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.99(d, 1H), 7.56(d, 1H), 6.74-6.57(m, 2H), 6.42(s, 1H), 6.41-6.08(m, 1H), 5.78-5.52(m, 1H), 4.25-4.19(m, 4H), 3.31-3.12(m, 1H), 3.02-2.92(m,

2H), 2.66-2.60(m, 6H), 2.20-2.17(m, 1H), 2.00-1.92(m, 1H), 1.80-1.72(m, 2H), 1.71-1.59(m, 2H), 1.34-1.29(m, 4H)

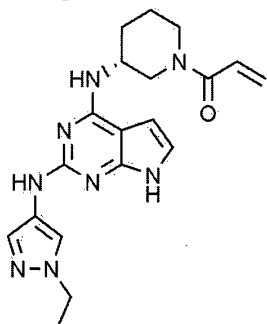
Example 20: Preparation of (R)-1-(3-(2-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (9.0 mg, yield: 17.6%) was prepared in the same manner as in Example 1, except that 1-(3-methoxybenzyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 8.09(d, 1H), 7.57(d, 1H), 7.23-7.22 (m, 1H), 6.85-6.78(m, 3H), 6.76-6.70(m, 2H), 6.40(s, 1H), 6.21-6.18(m, 1H), 5.73-5.71(m, 1H), 5.24-5.19 (m, 2H), 4.13-4.11 (m, 1H), 3.74 (s, 3H), 3.83-3.75(m, 1H), 3.38-3.01(m, 1H), 2.07-2.01(m, 1H), 1.92-1.87(m, 1H), 1.86-1.71(m, 1H), 1.69-1.49(m, 2H)

Example 21: Preparation of (R)-1-(3-(2-(1-ethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

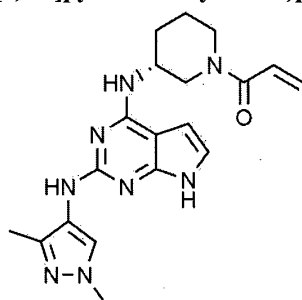


A title compound (15.0 mg, yield: 36.6%) was prepared in the same manner as in

Example 1, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.96(d, 1H), 7.52(d, 1H), 6.73-6.52(m, 2H), 6.39(s, 1H), 6.26-6.06(m, 1H), 5.78-5.51(m, 1H), 4.23-4.21(m, 3H), 4.20-4.07(m, 3H), 3.28-3.08(m, 1H), 2.16-2.14(m, 1H), 1.92-1.90(m, 1H), 1.61-1.59(m, 1H), 1.39-1.23(m, 2H)

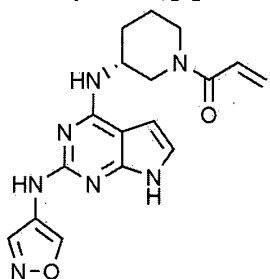
Example 22: Preparation of (R)-1-(3-(2-(1,3-dimethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (3.5 mg, yield: 8.5%) was prepared in the same manner as in Example 1, except that 1,3-dimethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.83(d, 1H), 6.73-6.56(m, 2H), 6.55(s, 1H), 6.40-6.39(m, 1H), 5.35-5.33(m, 1H), 4.18-4.10(m, 3H), 3.76(s, 3H), 3.22-3.16(m, 1H), 2.22(s, 3H), 2.19-2.17(m, 1H), 2.16-2.14(m, 1H), 1.95-1.90(m, 1H), 1.78-1.70(m, 1H), 1.65-1.60(m, 2H)

Example 23: Preparation of (R)-1-(3-(2-(isoxazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



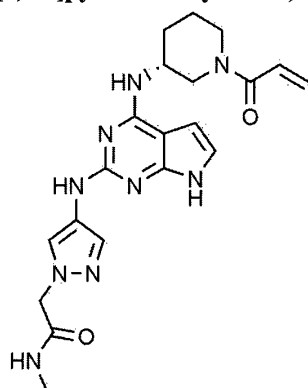
A title compound (4.5 mg, yield: 11.8%) was prepared in the same manner as in Example 1, except that isoxazole-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline

in Example 1.

¹H NMR (500MHz, CD₃OD) δ 9.04(d, 1H), 8.49(d, 1H), 6.78-6.57(m, 2H), 6.43-6.12(m, 1H), 5.56-5.34(m, 1H), 4.18-4.12(m, 3H), 3.25-3.16(m, 1H), 2.20-2.16(m, 1H), 1.93-1.91(m, 1H), 1.79-1.74(m, 1H), 1.71-1.60(m, 2H)

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Example 24: Preparation of (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide

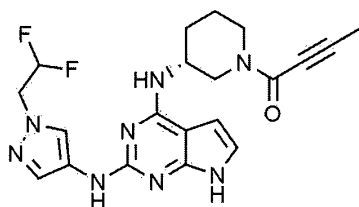


A title compound (11.9 mg, yield: 40.2%) was prepared in the same manner as in Example 1, except that 2-(4-amino-1H-pyrazol-1-yl)-N-methylacetamide was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.96-7.90(m, 1H), 7.57-7.50(m, 1H), 6.86-6.50(m, 1H), 6.27-6.09(m, 1H), 5.80-5.56(m, 1H), 4.42-4.29(m, 2H), 4.14-4.09(m, 2H), 3.97-3.41(m, 3H), 2.15-2.09(m, 1H), 1.93-1.86(m, 2H), 1.64-1.60(m, 1H), 1.45-1.40(m, 3H)

15

Example 25: Preparation of (R)-1-(3-(2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)but-2-yn-1-one



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Step 1: Preparation of tert-butyl-(R)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-

yl)amino)piperidine-1-carboxylate

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (500.0 mg, 2.7 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695.0 μ L, 4.0 Mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (639.3 mg, 4.0 eq) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and the organic layer was isolated, treated with magnesium sulfate, filtered, and concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (900.0 mg, yield: 98%).

¹H NMR (500MHz, CD₃OD) δ 7.06(s, 1H), 6.38(s, 1H), 4.24-4.20(m, 1H), 3.84-3.82(m, 1H), 3.80-3.30(m, 3H), 2.01-1.90(m, 1H), 1.80-1.75(m, 1H), 1.65-1.55(m, 2H), 1.43(s, 9H)

Step 2: Preparation of (R)-2-chloro-N-(piperidin-3-yl) -7H-pyrrolo[2,3-d]pyrimidine-4-amine

To *tert*-butyl-(R)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidine-4-yl)amino)piperidine-1-carboxylate (474.0 mg, 1.4 mmol), 6 N hydrochloric acid solution (5.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30 minutes, the reaction mixture was neutralized with saturated sodium bicarbonate solution. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure to obtain a title compound without isolation.

¹H NMR (500MHz, CD₃OD) δ 7.11(d, 1H), 6.63(d, 1H), 4.52- 4.49(m, 1H), 3.66-3.63(m, 1H), 3.37-3.34(m, 1H), 3.02-2.90(m, 2H), 2.19-2.16(m, 1H), 2.12-2.09(m, 1H), 1.90-1.77(m, 2H)

Step 3: Preparation of (R)-1-(3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidine-4-yl)amino)piperidin-1-yl)but-2-yn-1-one

After 2-butenic acid (84.1 mg, 0.2 mmol) was dissolved in tetrahydrofuran (3.0 mL), N,N-diisopropylethylamine (69.2 mg, 0.4 mmol) and 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazole[4,5-b]pyridinium 3-oxide hexafluorophosphate (90.6 mg, 0.2 mmol) was added thereto and stirred for 30 minutes. (R)-2-chloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine-3-amine (50.0 mg, 0.2 mmol) was added to this reaction mixture, followed by

stirring at room temperature for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (49.0 mg, yield: 77.8%).

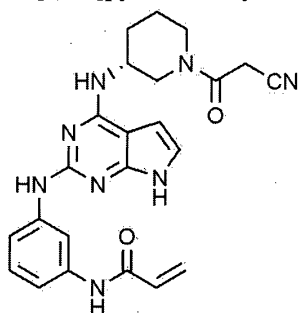
¹H NMR (500MHz, CD₃OD) δ 7.03-7.01(m, 1H), 6.61-6.55(m, 1H), 4.45-3.83(m, 3H), 3.65-3.58(m, 1H), 2.94-2.80(m, 1H), 2.15-2.09(m, 1H), 1.92-1.90(m, 2H), 1.85-1.79(m, 3H), 1.67-1.58(m, 1H)

Step 4: Preparation of (R)-1-(3-(2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)but-2-yn-1-one

(R)-1-(3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidine-4-yl)amino)piperidin-1-yl)but-2-yn-1-one (25.0 mg, 0.08 mmol) and 2,2-difluoroethyl-1H-pyrazol-4-amine (7.2 mg, 0.05 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (4.8 μL, 0.06 mmol) was added to the reaction mixture and reacted at 110 °C for 5 hours, and then the solvent was concentrated. This reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol. The residue was isolated by column chromatography to obtain a title compound (5.1 mg, yield: 24.3%).

¹H NMR (500MHz, CD₃OD) δ 8.08(s, 1H), 7.65-7.50(m, 1H), 6.76-6.73(m, 1H), 6.44-6.38(m, 1H), 6.30-6.06(m, 1H), 5.60-4.29(m, 6H), 3.29-3.15(m, 1H), 2.15-2.11(m, 1H), 2.04-2.00(m, 1H), 1.94-1.91(m, 1H), 1.85-1.65(m, 4H)

Example 26: Preparation of (R)-N-(3-(4-(1-(2-cyanoacetyl)piperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)acrylamide



Step 1: Preparation of *tert*-butyl-(R)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (500.0 mg, 2.7 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695.0 μ L, 4.0 mmol) and *tert*-butyl-(*R*)-3-aminopiperidine-1-carboxylate (639.3 mg, 4.0 eq) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (900.0 mg, yield: 98.0%).

¹H NMR (500MHz, CD₃OD) δ 7.06(s, 1H), 6.38(s, 1H), 4.24-4.20(m, 1H), 3.84-3.82(m, 1H), 3.80-3.30(m, 3H), 2.01-1.90(m, 1H), 1.80-1.75(m, 1H), 1.65-1.55(m, 2H), 1.43(s, 9H)

10

Step 2: Preparation of (R)-2-chloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine

To *tert*-butyl-(*R*)-3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (474.0 mg, 1.4 mmol), 6 N hydrochloric acid solution (5.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30 minutes, the reaction mixture was neutralized with saturated sodium bicarbonate solution. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure to obtain a title compound without isolation.

¹H NMR (500MHz, CD₃OD) δ 7.11(d, 1H), 6.63(d, 1H), 4.52- 4.49(m, 1H), 3.66-3.63(m, 1H), 3.37-3.34(m, 1H), 3.02-2.90(m, 2H), 2.19-2.16(m, 1H), 2.12-2.09(m, 1H), 1.90-1.77(m, 2H)

20

Step 3: Preparation of (R)-1-(3-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After cyanoacetic acid (40.6 mg, 0.4 mmol) was dissolved in tetrahydrofuran (3.0 mL), N,N-diisopropylethylamine (140 μ L, 0.8 mmol) and 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazole[4,5-b]-pyridinium 3-oxide hexafluorophosphate (181.3 mg, 0.4 mmol) were added thereto and stirred for 30 minutes. (R)-2-chloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine (100.0 mg, 0.3 mmol) was added to this reaction mixture and stirred at room temperature for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then

25

30

concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (49.0 mg, yield: 77.8%).

¹H NMR (500MHz, CD₃OD) δ 6.99(d, 1H), 6.45(d, 1H), 4.20-4.00(m, 2H), 3.84-3.62(m, 1H), 3.48-3.41(m, 1H), 2.97(s, 2H), 2.10(s, 1H), 1.85-1.82(m, 1H), 1.72-1.70(m, 1H)

5

Step 4: Preparation of N-(3-nitrophenyl)acrylamide

3-Nitroaniline (300.0 mg, 2.2 mmol) was dissolved in tetrahydrofuran (3.0 mL) to which triethylamine (450.0 μL, 3.3 mmol) was added, and the mixture was stirred for 10 minutes. After cooling to 0 °C, acryloyl chloride (260.0 μL, 3.3 mmol) was added, followed by stirring for 2 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (365.0 mg, yield: 87.0%).

¹H NMR (500MHz, CD₃OD) δ 8.70(s, 1H), 7.98-7.93(m, 2H), 7.59-7.55(m, 1H), 6.45-6.42(m, 2H), 5.83(s, 1H)

15

Step 5: Preparation of N-(3-aminophenyl)acrylamide

N-(3-nitrophenyl)acrylamide (365.0 mg, 1.9 mmol) was dissolved in ethanol (3.0 mL). Tin chloride (857.0 mg, 3.8 mmol) was added thereto and the mixture was stirred at 100 °C for 2 hours and then neutralized with 1 N sodium hydroxide solution. The reaction mixture was extracted with dichloromethane and concentrated under reduced pressure to obtain a title compound (300.3 mg, yield: 98.0%).

¹H NMR (500MHz, CD₃OD) δ 7.47(s, 1H), 7.26-7.23(m, 1H), 7.01(t, 1H), 6.74(d, 1H), 6.45-6.42(m, 1H), 6.30-6.21(m, 1H), 5.74-5.72(m, 1H)

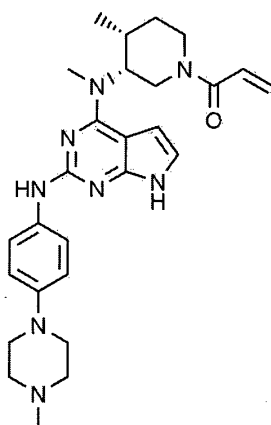
Step 6: Preparation of (R)-N-(3-(4-(1-(2-cyanoacetyl)piperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-2-ylamino)phenyl)acrylamide

(R)-1-(3-((2-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (25.0 mg, 0.08 mmol) and N-(3-aminophenyl)acrylamide (7.2 mg, 0.05 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (4.8 μL, 0.06 mmol) was added to the reaction mixture and reacted at 110 °C for 5 hours, and then the solvent was concentrated. This

reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol. The residue was isolated by column chromatography to obtain a title compound (5.1 mg, yield: 24.3%).

¹H NMR (500MHz, CD₃OD) δ 7.22-7.21(m, 1H), 7.19-7.18(m, 1H), 7.15-7.14(m, 1H), 6.97-7.95(m, 1H), 6.79(s, 1H), 6.47-6.42(m, 2H), 6.37-6.31(m, 1H), 5.76-5.73(m, 1H), 5.49(s, 1H), 4.59-4.52(m, 1H), 4.40-4.34(m, 1H), 3.95-3.87(m, 1H), 3.44(s, 2H), 2.81-2.74(m, 1H), 2.17-2.15(m, 1H), 2.01-1.96(m, 1H), 1.85-1.83(m, 1H), 1.73-1.60(m, 2H)

Example 27: Preparation of 1-((3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 2,4-dichloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine

2,4-Dichloro-7H-pyrrolo[2,3-d]pyrimidine (3.0 g, 15.9 mmol) and 4-toluenesulfonyl chloride (3.7 g, 23.9 mmol) were dissolved in acetone (20.0 mL). After cooling to 0 °C, 2 M sodium hydroxide solution (12.0 ml) was slowly added dropwise thereto, stirred for 2 hours, and then filtered through acetone to obtain a title compound (2.9 g, yield: 80.0%).

¹H NMR (500MHz, CD₃OD) δ 8.12(d, 2H), 7.76(d, 1H), 7.37(d, 2H), 6.68(d, 1H), 2.43(s, 3H)

Step 2: Preparation of tert-butyl (3R, 4R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidine-1-carboxylate

2,4-dichloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine (1.5 g, 5.1 mmol) and (3R,4R)-1-

benzyl-N,4-dimethylpiperidin-3-amine (965.0 mg, 5.1 mmol) were dissolved in ethanol (20.0 mL). N,N-Diisopropylethylamine (4.2 g, 30.6 mmol) was added dropwise thereto and reacted at 110 °C for 12 hours, then concentrated under reduced pressure, and the residue was isolated with column chromatography to obtain a title compound (1.3 g, yield: 85.0%).

5 ¹H NMR (500MHz, CD₃OD) δ 8.18(d, 2H), 7.45(d, 1H), 7.37-7.22(m, 7H), 6.62(s, 1H), 3.76-3.51(m, 2H), 3.49-3.31(m, 3H), 2.89-2.45(m, 1H), 2.38(s, 3H), 2.35-2.13(m, 1H), 1.70-1.56(m, 3H), 0.88(s, 3H)

10 **Step 3: Preparation of *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate**

After *tert*-butyl (3R,4R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidine-1-carboxylate (100 mg, 0.3 mmol), palladium acetate (3.04 mg, 0.02 mmol), 4,5-bis(diphenylphosphino)-9,9-dimethylxanthene (7.87 mg, 0.01 mmol), 4-(4-methylpiperazin-1-yl)aniline (53.26 mg, 0.2 mmol) and cesium carbonate (184 mg, 0.6 mmol) were dissolved in dioxane (1.0 mL), and the mixture was reacted at 120 °C using a microwave apparatus for 30 minutes. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (75.0 mg, yield: 65.0%).

20 ¹H NMR (500MHz, CD₃OD) δ 7.93(d, 2H), 7.64(d, 1H), 7.31-7.22(m, 4H), 7.20-7.16(m, 4H), 6.98(d, 2H), 6.12(d, 1H), 3.50(s, 2H), 3.32-3.31(m, 1H), 3.15(s, 4H), 2.72(t, 1H), 2.68(s, 3H), 2.69(s, 4H), 2.65(d, 1H), 2.51(s, 1H), 2.43(s, 1H), 2.35(s, 3H), 2.27(s, 3H), 2.20(s, 1H), 1.85(s, 1H), 1.56(s, 1H), 0.89(d, 3H)

25 **Step 4: Preparation of *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate**

After *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (100 mg, 0.2 mmol) was dissolved in methanol (3.0 mL), potassium hydroxide (16 mg, 0.4

mmol) was added thereto. After reacting at 70 °C for 12 hours, the methanol was removed, and water and diethyl ether were added. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (80.0 mg, yield: 88.0%).

5 ¹H NMR (500MHz, CD₃OD) δ 7.58(d, 2H), 7.32-7.29(m, 4H), 7.24-7.22(m, 1H), 6.94(d, 2H), 6.77(d, 1H), 6.56(d, 1H), 3.50(s, 2H), 3.33-3.30(m, 1H), 3.15(s, 4H), 2.72(t, 1H), 2.70(s, 4H), 2.67(d, 1H), 2.52(s, 1H), 2.45(s, 1H), 2.37(s, 3H), 2.29(s, 3H), 2.23(s, 1H), 1.87(s, 1H), 1.57(s, 1H), 0.88(d, 3H)

10 **Step 5: Preparation of N4-methyl-N2-(4-(4-methylpiperazin-1-yl)phenyl)-N4-((3R,4R)-4-methylpiperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine**

After *tert*-butyl (3R, 4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (70.0 mg, 0.1 mmol) was dissolved in methanol, an excess of 10 wt. % palladium carbon was added and
15 replaced with hydrogen gas. After stirring for 5 hours, the reaction mixture was filtered through celite. The filtrate was concentrated to obtain a title compound (65.0 mg, yield: 98.0%).

¹H NMR (500MHz, CD₃OD) δ 7.71(d, 2H), 6.94(d, 2H), 6.93(d, 1H), 6.47(d, 1H), 3.37-3.33(m, 2H), 3.16(s, 4H), 2.74(t, 1H), 2.72(s, 4H), 2.66(d, 1H), 2.54(s, 1H), 2.42(s, 1H), 2.36(s, 3H), 2.30(s, 3H), 2.24(s, 1H), 1.88(s, 1H), 0.99(d, 3H)

20

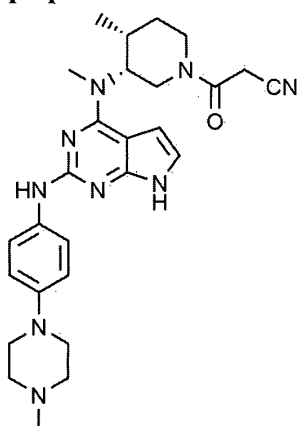
Step 6: Preparation of 1-((3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

N4-methyl-N2-(4-(4-methylpiperazin-1-yl)phenyl)-N4-((3R, 4R)-4-methylpiperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine (40 mg, 0.1 mmol) was dissolved in
25 dichloromethane (1.0 mL) to which triethylamine (40.0 μL, 0.3 mmol) was added, and the mixture was stirred for 10 minutes. After cooling to 0 °C, acryloyl chloride (8.0 μL, 0.1 mmol) was added, followed by stirring for 2 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (11.0 mg, yield: 23.0%).

30 ¹H NMR (500MHz, CD₃OD) δ 7.16(d, 2H), 7.09(s, 1H), 6.98(d, 2H), 6.83-6.39(m,

2H), 6.345-6.19(m, 1H), 5.77-5.65(m, 1H), 4.76(s, 1H), 3.81-3.77(m, 2H), 3.68-3.63(m, 2H), 3.36(s, 3H), 3.22(s, 4H), 2.64(s, 4H), 2.37(s, 3H), 2.24(s, 1H), 1.64(s, 1H), 1.59(s, 1H), 0.99-0.97(m, 3H)

5 **Example 28: Preparation of 3-((3R,4R)-4-methyl-3-(methyl(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile**



Step 1: Preparation of 2,4-dichloro-7-tosyl-7H-pyrrolo [2,3-d]pyrimidine

10 2,4-Dichloro-7H-pyrrolo[2,3-d]pyrimidine (3.0 g, 15.9 mmol) and 4-toluenesulfonyl chloride (3.7 g, 23.9 mmol) were dissolved in acetone (20.0 mL). After cooling to 0 °C, 2 M sodium hydroxide solution (12.0 mL) was slowly added dropwise, followed by stirring for 2 hours, and then washed and filtered with acetone to obtain a title compound (2.9 g, yield: 80.0%).

¹H NMR (500MHz, CD₃OD) δ 8.12(d, 2H), 7.76(d, 1H), 7.37(d, 2H), 6.68(d, 1H),
15 2.43(s, 3H)

Step 2: Preparation of *tert*-butyl (3R, 4R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidine-1-carboxylate

20 2,4-dichloro-7-tosyl-7H-pyrrolo [2,3-d]pyrimidine (1.5 g, 5.1 mmol) and (3R, 4R)-1-benzyl-N,4-dimethylpiperidine-3-amine (965.0 mg, 5.1 mmol) were dissolved in ethanol(20.0 mL). N,N-Diisopropylethylamine (4.2 g, 30.6 mmol) was added dropwise and the mixture was reacted at 110 °C for 12 hours and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (1.3 g, yield: 85.0%).

¹H NMR (500MHz, CD₃OD) δ 8.18(d, 2H), 7.45(d, 1H), 7.37-7.22(m, 7H), 6.62(s, 1H), 3.76-3.51(m, 2H), 3.49-3.31(m, 3H), 2.89-2.45(m, 1H), 2.38(s, 3H), 2.35-2.13(m, 1H), 1.70-1.56(m, 3H), 0.88(s, 3H)

5 **Step 3: Preparation of *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate**

After *tert*-butyl (3R,4R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidine-1-carboxylate (100 mg, 0.3 mmol), palladium acetate
10 (3.04 mg, 0.02 mmol), 4,5-bis(diphenylphosphino)-9,9-dimethylxanthene (7.87 mg, 0.01 mmol), 4-(4-methylpiperazin-1-yl)aniline (53.26 mg, 0.2 mmol) and cesium carbonate (184 mg, 0.6 mmol) were dissolved in dioxane (1.0 mL), the mixture was reacted at 120 °C using a microwave apparatus for 30 minutes. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by
15 column chromatography to obtain a title compound (75.0 mg, yield: 65%).

¹H NMR (500MHz, CD₃OD) δ 7.93(d, 2H), 7.64(d, 1H), 7.31-7.22(m, 4H), 7.20-7.16(m, 4H), 6.98(d, 2H), 6.12(d, 1H), 3.50(s, 2H), 3.32-3.31(m, 1H), 3.15(s, 4H), 2.72(t, 1H), 2.68(s, 3H), 2.69(s, 4H), 2.65(d, 1H), 2.51(s, 1H), 2.43(s, 1H), 2.35(s, 3H), 2.27(s, 3H), 2.20(s, 1H), 1.85(s, 1H), 1.56(s, 1H), 0.89(d, 3H)

20

Step 4: Preparation of *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate
25 (100.0 mg, 0.2 mmol) was dissolved in methanol (3.0 mL), potassium hydroxide (16.0 mg, 0.4 mmol) was added thereto. After reacting at 70 °C for 12 hours, methanol was removed, and water and diethyl ether were added. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column
30 chromatography to obtain 80.0 mg (yield: 88.0%) of the title compound (80.0 mg, yield: 88.0%).

¹H NMR (500MHz, CD₃OD) δ 7.58(d, 2H), 7.32-7.29(m, 4H), 7.24-7.22(m, 1H), 6.94(d, 2H), 6.77(d, 1H), 6.56(d, 1H), 3.50(s, 2H), 3.33-3.30(m, 1H), 3.15(s, 4H), 2.72(t, 1H), 2.70(s, 4H), 2.67(d, 1H), 2.52(s, 1H), 2.45(s, 1H), 2.37(s, 3H), 2.29(s, 3H), 2.23(s, 1H), 1.87(s, 1H), 1.57(s, 1H), 0.88(d, 3H)

Step 5: Preparation of N4-methyl-N2-(4-(4-methylpiperazin-1-yl)phenyl)-N4-((3R,4R)-4-methylpiperidin-[2,3-d]pyrimidine-2,4-diamine

After *tert*-butyl (3R,4R)-4-methyl-3-(methyl(2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (70.0 mg, 0.1 mmol) was dissolved in methanol, an excess of 10 wt. % palladium carbon was added and replaced with hydrogen gas. The reaction mixture was stirred for 5 hours, filtered through celite, and the filtrate was concentrated to obtain a title compound (65.0 mg, yield: 98.0%).

¹H NMR (500MHz, CD₃OD) δ 7.71(d, 2H), 6.94(d, 2H), 6.93(d, 1H), 6.47(d, 1H), 3.37-3.33(m, 2H), 3.16(s, 4H), 2.74(t, 1H), 2.72(s, 4H), 2.66(d, 1H), 2.54(s, 1H), 2.42(s, 1H), 2.36(s, 3H), 2.30(s, 3H), 2.24(s, 1H), 1.88(s, 1H), 0.99(d, 3H)

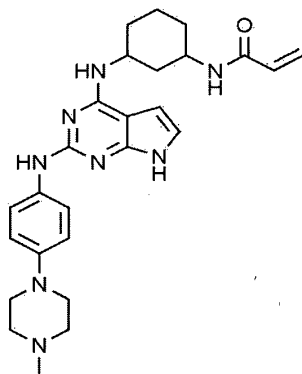
Step 6: Preparation of 3-((3R,4R)-4-methyl-3-(methyl 2-(4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-7-yl)-3-oxopropanenitrile

After N4-methyl-N2-(4-(4-methylpiperazin-1-yl)phenyl)-N4-((3R,4R)-4-methylpiperidin-[2,3-d]pyrimidine-2,4-diamine (50.0 mg, 0.1 mmol) and 2-cyanoacetic acid (20.1 mg, 0.1 mmol) were dissolved in tetrahydrofuran, 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxhexafluorophosphate (89.9 mg, 0.1 mmol) and N,N-diisopropylethylamine (30.6 μL, 0.2 mmol) were added thereto. The mixture was stirred for 20 hours and then water and diethyl ether were added. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (9.0 mg, yield: 20.0%).

¹H NMR (500MHz, CD₃OD) δ 7.49(d, 2H), 6.93(d, 2H), 6.91(d, 1H), 6.49(d, 1H), 4.09-4.01(dd, 1H), 3.89-3.82(m, 1H), 3.72-3.60(m, 1H), 3.58-3.52(m, 1H), 3.34(s, 3H), 3.16(s, 4H),

2.64(s, 4H), 2.46(s, 1H), 2.35(s, 3H), 2.19(s, 2H), 1.89-1.75(m, 1H), 1.69-1.55(m, 2H), 1.13-1.09(m, 3H)

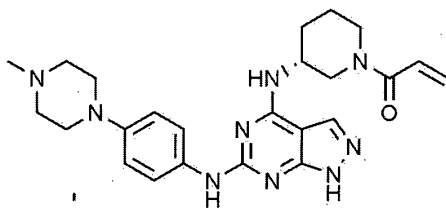
Example 29: Preparation of N-(3-(2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)cyclohexyl)acrylamide



A title compound (4.0 mg, yield: 17.4%) was prepared in the same manner as in Example 1, except that *tert*-butyl (3-aminocyclohexyl)carbonate was used instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.55(d, 2H), 6.94(d, 2H), 6.92(d, 1H), 6.84-6.69(m, 1H), 6.47-6.30(m, 1H), 6.28-6.23(m, 1H), 5.65-5.63(m, 1H), 4.45(s, 1H), 3.16(s, 4H), 3.09-2.98(m, 3H), 2.64-2.61(m, 4H), 2.33(s, 3H), 2.08-2.01(m, 1H), 1.92-1.91(m, 1H), 1.75-1.62(m, 2H)

Example 30: Preparation of (R)-1-(3-(6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of *tert*-butyl (R)-3-((6-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine (1.0 g, 5.3 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695 μL, 7.9 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (1.3 g, 6.35 mmol) were added thereto. The reaction mixture

was stirred at 110 °C for 12 hours, and the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (1.2 g, yield: 62.9%).

¹H NMR (500MHz, CD₃OD) δ 8.11(s, 1H), 4.26-4.07(m, 1H), 3.98-3.90(m, 1H),
5 3.70-3.65(m, 1H), 3.16-3.10(m, 2H), 2.10-2.05(m, 1H), 1.87-1.75(m, 1H), 1.71-1.58(m, 2H),
1.45-1.34(m, 9H)

Step 2: Preparation of (R)-6-chloro-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine hydrochloride

10 To *tert*-butyl (R)-3-((6-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (500.0 mg, 1.4 mmol), 6 N hydrochloric acid solution (5.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 8.17(s, 1H), 4.55-4.51(m, 1H), 3.68-3.65(m, 2H),
15 3.16-2.92(m, 2H), 2.21-2.10(m, 2H), 1.98-1.83(m, 2H)

Step 3: Preparation of (R)-1-(3-((6-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-6-chloro-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine
20 hydrochloride (421.0 mg, 1.5 mmol) was dissolved in dichloromethane (10.0 mL), triethylamine (608.8 μL, 4.4 mmol) was added at 0 °C, and the mixture was stirred for 30 minutes. Acryloyl chloride (124.2 μL, 1.5 mmol) was added to the reaction mixture and was stirred at 0 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column
25 chromatography to obtain a title compound (130.0 mg, yield: 29.3%).

¹H NMR (500MHz, CD₃OD) δ 8.10(s, 1H), 6.88-6.77(m, 1H), 6.19-6.16(m, 1H),
5.76-5.64(m, 1H), 4.47-4.24(m, 2H), 4.10(d, 1H), 3.98(d, 1H), 3.21-3.06(m, 1H), 2.17(s, 1H),
1.94-1.91(m, 1H), 1.83-1.79(m, 1H), 1.67-1.64(m, 1H)

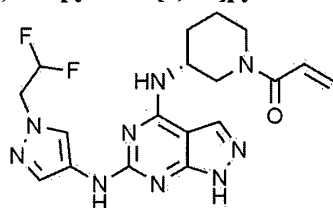
30 **Step 4: Preparation of (R)-1-(3-(6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-**

pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

(R)-1-(3-((6-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (30.0 mg, 0.10 mmol) and 4-(4-methylpiperazin-1-yl)aniline (12.5 mg, 0.07 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (6.0 μ L, 0.08 mmol) was added to the reaction mixture, followed by reacting at 110 $^{\circ}$ C for 12 hours, and then the solvent was concentrated. This reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol. The residue was isolated by column chromatography to obtain a title compound (6.6 mg, yield: 24.4%).

^{1} H NMR (500MHz, CD_3OD) δ 7.90-7.88(m, 1H), 7.59-7.53(m, 2H), 6.96-6.91(m, 2H), 6.90-6.49(m, 1H), 6.31-6.05(m, 1H), 5.84-5.48(m, 1H), 4.43-4.20(m, 1H), 4.07-4.05(m, 1H), 3.18-3.13(m, 4H), 2.75-2.73(m, 2H), 2.62-2.54(m, 5H), 2.32(s, 3H), 2.17-2.10(m, 1H), 1.99-1.88(m, 1H), 1.80-1.67(m, 2H)

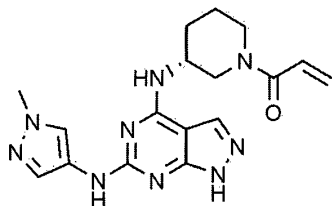
Example 31: Preparation of (R)-1-(3-(6-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (9.3 mg, yield: 28.2%) was prepared in the same manner as in Example 30, except that 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

^{1} H NMR (500MHz, CD_3OD) δ 8.12-8.5(m, 1H), 7.91(s, 1H), 7.65-7.55(m, 1H), 6.89-6.50(m, 1H), 6.27-6.07(m, 2H), 5.80-5.55(m, 1H), 4.58-4.45(m, 2H), 4.28-4.20(m, 1H), 4.11-4.03(m, 1H), 3.20-3.16(m, 2H), 2.75-2.69(m, 1H), 2.20-2.15(m, 1H), 1.94-1.80(m, 1H), 1.78-1.62(m, 2H)

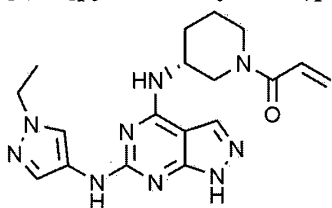
Example 32: Preparation of (R)-1-(3-(6-(1-methyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (5.2 mg, yield: 12.8%) was prepared in the same manner as in Example 30, except that 1-methyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

5 ^1H NMR (500MHz, CD_3OD) δ 7.95(d, 2H), 7.55(s, 1H), 6.86-6.56(m, 1H), 6.26-6.07(m, 1H), 5.79-5.53(m, 1H), 4.26-4.20(m, 1H), 4.09(d, 2H), 3.82(s, 3H), 3.22-3.16(m, 1H), 2.23-2.17(m, 1H), 1.96-1.92(m, 1H), 1.80-1.71(m, 1H), 1.67-1.60(m, 2H)

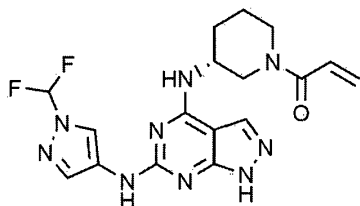
Example 33: Preparation of (R)-1-(3-(6-(1-ethyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.5 mg, yield: 17.9%) was prepared in the same manner as in Example 30, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

15 ^1H NMR (500MHz, CD_3OD) δ 7.99(d, 1H), 7.91(s, 1H), 7.57(d, 1H), 6.86-6.57(m, 1H), 6.26-6.07(m, 1H), 5.79-5.54(m, 1H), 4.27-4.21(m, 1H), 4.14-4.07(m, 2H), 3.45-3.36(m, 1H), 3.22-3.16(m, 2H), 2.20-2.17(m, 1H), 1.97-1.94(m, 1H), 1.80-1.72(m, 1H), 1.68-1.61(m, 2H), 1.46-1.41(d, 3H)

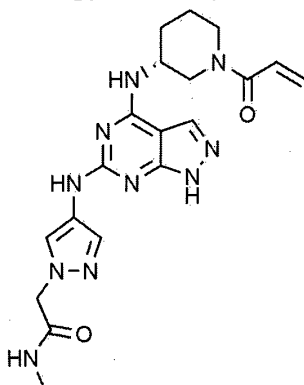
20 **Example 34: Preparation of (R)-1-(3-(6-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (6.5 mg, yield: 14.7%) was prepared in the same manner as in Example 30, except that 1-(difluoromethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

5 ^1H NMR (500MHz, CD_3OD) δ 8.42(s, 1H), 7.93(d, 1H), 7.79(d, 1H), 7.50-7.26(m, 1H), 6.85-6.61(m, 1H), 6.26-6.07(m, 1H), 5.79-5.55(m, 1H), 4.29-4.21(m, 1H), 4.09-4.04(m, 2H), 3.50-3.44(m, 1H), 2.21-2.17(m, 1H), 1.94-1.80(m, 1H), 1.71-1.66(m, 3H)

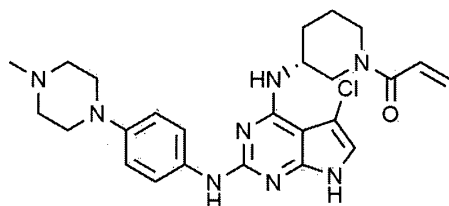
Example 35: Preparation of (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide



A title compound (12.4 mg, yield: 26.6%) was prepared in the same manner as in Example 30, except that 2-(4-amino-1H-pyrazol-1-yl)-N-methylacetamide was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

15 ^1H NMR (500MHz, CD_3OD) δ 8.10(d, 1H), 7.91(s, 1H), 7.65-7.59(m, 1H), 6.84-6.60(m, 1H), 6.26-6.07(m, 1H), 5.78-5.55(m, 1H), 4.79-4.77(m, 2H), 4.28-4.21(m, 1H), 4.07-4.05(m, 1H), 3.49-3.13(m, 2H), 2.76(d, 3H), 2.17-2.13(m, 1H), 1.94-1.82(m, 1H), 1.80-1.61(m, 3H)

Example 36: Preparation of (R)-1-(3-(5-chloro-2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 2,4,5-trichloro-7H-pyrrolo[2,3-d]pyrimidine

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (3.0 g, 16.0 mmol) and N-chlorosuccinimide (2.6 g, 19.2 mmol) were dissolved in N,N-dimethylformamide (20.0 mL), the mixture was stirred at 0 °C for 8 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (2.6 g, yield: 72.5%).

¹H NMR (500MHz, DMSO-d₆) δ 13.07(s, 1H), 7.94(s, 1H)

Step 2: Preparation of (R)-tert-butyl 3-((2,5-dichloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 2,4,5-trichloro-7H-pyrrolo[2,3-d]pyrimidine (2.6 g, 11.6 mmol) was dissolved in ethanol (25.0 mL), N,N-diisopropylethylamine (3.0 mL, 17.4 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (2.8 g, 13.9 mmol) were added thereto. After stirring the reaction mixture at 110 °C for 12 hours, the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (3.5 g, yield: 77.4%).

¹H NMR (500MHz, DMSO-d₆) δ 6.96(s, 1H), 6.09(bs, 1H), 4.14(bs, 1H), 3.60-3.30(m, 4H), 1.98-1.15(m, 13H)

Step 3: Preparation of (R)-2,5-dichloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride

To (R)-*tert*-butyl 3-((2,5-dichloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (3.5 g, 9.0 mmol), 6 N hydrochloric acid solution (15.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, DMSO-d₆) δ 12.2(bs, 1H), 9.53-9.51(m, 1H), 9.10-9.08(m, 1H),

7.34(s, 1H), 6.63-6.61(m, 1H), 4.48-4.47(m, 1H), 3.29-3.27(m, 1H), 3.17-3.14(m, 1H), 3.08-3.02(m, 1H), 2.71-2.69(m, 1H), 1.95-1.94(m, 1H), 1.85-1.84(m, 1H), 1.79-1.72(m, 2H)

Step 4: Preparation of (R)-1-(3-((2,5-dichloro-7H-pyrrolo [2,3-d] pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-2,5-dichloro-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride (3.4 g, 10.5 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water, sodium bicarbonate (2.6 g, 31.4 mmol) was added at 0 °C and the mixture was stirred for 30 minutes. Acryloyl chloride (1.3 mL, 10.5 mmol) was added to the reaction mixture and was stirred at 0 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (2.7 g, yield: 87.1%).

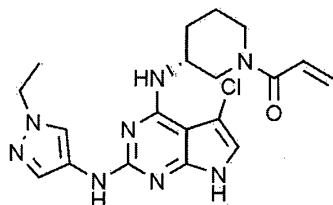
¹H NMR (500MHz, DMSO-d₆) δ 12.1(bs, 1H), 7.34-7.32(m, 1H), 6.81-6.79(m, 1H), 6.42-6.40(m, 1H), 5.68-5.59(m, 1H), 4.16-3.33(m, 4H), 3.10-3.02(m, 1H), 1.94-1.83(m, 2H), 1.66-1.50(m, 2H)

Step 5: Preparation of (R)-1-(3-(5-chloro-2-(4-(4-methylpiperazin-1-yl)phenylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

(R)-1-(3-((2,5-dichloro-7H-pyrrolo [2,3-d] pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (200.0 mg, 0.6 mmol) and 4-(4-methylpiperazin-1-yl)aniline (75.0 mg, 0.4 mmol) were dissolved in 2-butanol (15.0 mL). Trifluoroacetic acid (36.0 μL, 0.5 mmol) was added to the reaction mixture, followed by reacting at 110 °C for 12 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to obtain a title compound (4 mg, yield: 9.6%).

¹H NMR (500MHz, CD₃OD) δ 7.56-7.51(m, 2H), 6.95-6.91(m, 2H), 6.86-6.49(m, 2H), 6.26-6.04(m, 1H), 5.79-5.43(m, 1H), 4.39-4.25(m, 1H), 3.93-3.80(m, 1H), 3.70-3.63(m, 2H), 3.50-3.31(m, 1H), 3.13-3.12(m, 4H), 2.67-2.66(m, 4H), 2.37(s, 3H), 2.08-2.02(m, 1H), 1.91-1.84(m, 2H), 1.66-1.59(m, 1H)

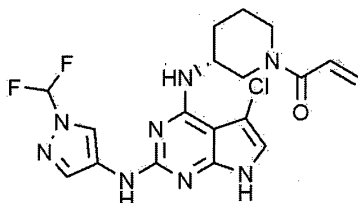
Example 37: Preparation of (R)-1-(3-(5-chloro-2-(1-ethyl-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (33.1 mg, yield: 23.5%) was prepared in the same manner as in Example 36, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.94-7.91(m, 1H), 7.54-7.50(m, 1H), 6.85-6.52(m, 2H), 6.25-6.06(m, 1H), 5.78-5.48(m, 1H), 4.39-4.28(m, 1H), 4.10(q, J = 7.0 Hz, 2H), 3.91-3.82(m, 1H), 3.71-3.60(m, 2H), 3.44-3.31(m, 1H), 2.09-2.01(m, 1H), 1.88-1.84(m, 2H), 1.66-1.59(m, 1H), 1.43(t, J = 7.0 Hz, 3H)

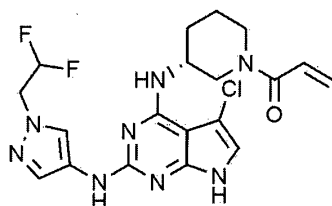
Example 38: Preparation of (R)-1-(3-(5-chloro-2-(1-(difluoromethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (6.6 mg, yield: 26.7%) was prepared in the same manner as in Example 36, except that 1-(difluoromethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 8.38-8.36(m, 1H), 7.76-7.73(m, 1H), 7.50-7.21(m, 1H), 6.85-6.50(m, 2H), 6.24-6.06(m, 1H), 5.78-5.51(m, 1H), 4.36-4.27(m, 1H), 3.86-3.62(m, 3H), 3.50-3.45(m, 1H), 2.10-2.08(m, 1H), 1.89-1.87(m, 2H), 1.70-1.60(m, 1H)

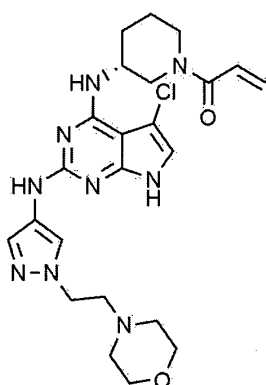
Example 39: Preparation of (R)-1-(3-(5-chloro-2-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (54.7 mg, yield: 32.6%) was prepared in the same manner as in Example 36, except that 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

5 ^1H NMR (500MHz, CD_3OD) δ 8.08-8.00(m, 1H), 7.62-7.55(m, 1H), 6.85-6.52(m, 2H), 6.25-5.99(m, 2H), 5.78-5.49(m, 1H), 4.47-4.44(m, 2H), 4.34-4.27(m, 1H), 3.88-3.86(m, 1H), 3.65-3.62(m, 2H), 3.44-3.40(m, 1H), 2.10-2.07(m, 1H), 1.85-1.84(m, 2H), 1.70-1.59(m, 1H)

Example 40: Preparation of (R)-1-(3-(5-chloro-2-(1-(2-morpholinoethyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

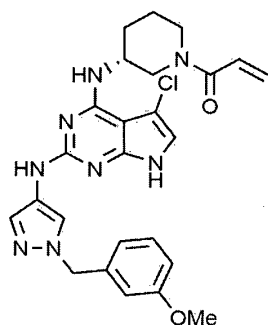


A title compound (5.2 mg, yield: 18.4%) was prepared in the same manner as in Example 36, except that 1-(2-morpholinoethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

15 ^1H NMR (500MHz, CD_3OD) δ 7.97-7.94(m, 1H), 7.57-7.55(m, 1H), 6.85-6.55(m, 1H), 6.25-6.05(m, 1H), 5.78-5.48(m, 1H), 4.34-4.21(m, 4H), 3.95-3.80(m, 1H), 3.75-3.60(m, 6H), 2.79(t, J=6.5Hz, 3H), 2.60-2.40(m, 4H), 2.09-2.06(m, 1H), 1.67-1.59(m, 1H), 2.03-2.00(m, 2H)

Example 41: Preparation of (R)-1-(3-(5-chloro-2-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

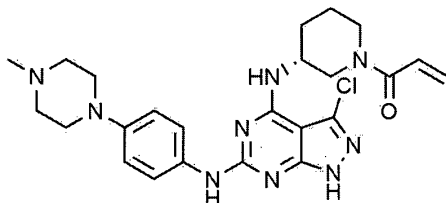
one



A title compound (4.4 mg, yield: 15.4%) was prepared in the same manner as in Example 36, except that 1-(3-methoxybenzyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 8.03-7.92(m, 1H), 7.57-7.53(m, 1H), 7.23-7.21(m, 1H), 6.86-6.45(m, 6H), 6.24-6.00(m, 1H), 5.80-5.45(m, 1H), 5.27-5.25(m, 2H), 4.30-4.18(m, 1H), 3.75(s, 3H), 3.70-3.40(m, 3H), 1.84-1.79(m, 2H), 1.63-1.59(m, 1H)

Example 42: Preparation of (R)-1-(3-(3-chloro-6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 3,4,6-trichloro-1H-pyrazolo [3,4-d] pyrimidine

4,6-Dichloro-1H-pyrazolo[3,4-d]pyrimidine (5.0 g, 26.5 mmol) and N-chlorosuccinimide (5.3 g, 39.7 mmol) were dissolved in N,N-dimethylformamide (50.0 mL). After stirring at room temperature for 24 hours, the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (3.3 g, yield: 56.0%).

Step 2: Preparation of tert-butyl (R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine (3.3 g, 14.8 mmol) was dissolved

in ethanol (50 mL), N,N-diisopropylethylamine (3.9 mL, 22.2 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (3.1 g, 15.6 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 3 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was
5 isolated by column chromatography to obtain a title compound (4.3 g, yield: 75.8%).

¹H NMR (500MHz, CD₃OD) δ 4.30-4.25(m, 1H), 3.76-3.74(m, 1H), 3.52-3.51(m, 2H), 3.45-3.40(m, 1H), 2.00-1.98-(m, 1H), 1.92-1.85(m, 1H), 1.80-1.75(m, 1H), 1.65-1.60(m, 1H), 1.45-1.34(m, 9H)

10 **Step 3: Preparation of (R)-3,6-dichloro-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine**

To *tert*-butyl (R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (4.3 g, 11.2 mmol), 6 N hydrochloric acid solution (30.0 mL, excessive amount) dissolved in methanol was added. After stirring at room temperature for 30
15 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 4.40-4.36(m, 1H), 3.44-3.40(m, 1H), 2.99-2.96(m, 1H), 2.74-2.66(m, 2H), 2.10-2.04(m, 1H), 1.89-1.81(m, 1H), 1.76-1.68(m, 1H)

20 **Step 4: Preparation of (R)-1-(3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one**

After (R)-3,6-dichloro-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine (4.0 g, 12.4 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran and distilled water, sodium bicarbonate (3.1 g, 3.7 mmol) was added at -20 °C and then stirred for 30 minutes.
25 Acryloyl chloride (4.1 mL, 13.0 mmol) was slowly added dropwise to the reaction mixture and then stirred at -20 °C for 30 minutes. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (4.0 g, yield: 95.2%).

¹H NMR (500MHz, CD₃OD) δ 6.86-6.78(m, 1H), 6.22-6.17(m, 1H), 5.78-5.68(m, 1H), 4.55-4.03(m, 3H), 3.71-3.66(m, 1H), 3.58-3.50(m, 1H), 2.12-2.00(m, 1H), 1.93-1.84(m,
30

2H), 1.73-1.68(m, 1H)

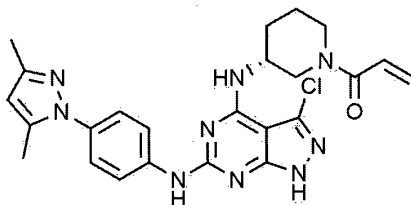
Step 5: Preparation of (R)-1-(3-(3-chloro-6-(4-(4-methylpiperazin-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one

5 (R)-1-(3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (30.0 mg, 0.09 mmol) and 4-(4-methylpiperazin-1-yl)aniline (11.2 mg, 0.06 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (5.6 μ L, 0.07 mmol) was added to the reactant, followed by reacting at 110 °C for 12 hours, and then the solvent was concentrated. This reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol.
10 The residue was isolated by column chromatography to obtain a title compound (15.0 mg, yield: 51.7%).

¹H NMR (500MHz, CD₃OD) δ 7.57-7.52(m, 2H), 6.92-6.90(m, 2H), 6.83-6.50(m, 1H), 6.28-6.07(m, 1H), 5.80-5.51(m, 1H), 4.40-3.88(m, 3H), 3.66-3.39(m, 2H), 3.13(s, 4H), 2.63(s, 4H), 2.35(s, 3H), 2.08-2.00(m, 1H), 1.91-1.84(m, 2H), 1.63-1.60(m, 1H)

15

Example 43: Preparation of (R)-1-(3-(3-chloro-6-(4-(3,5-dimethyl-1H-pyrazol-1-yl)phenylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



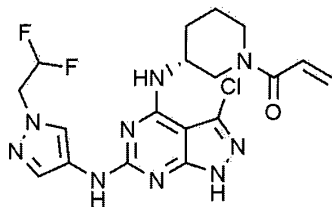
A title compound (6.1 mg, yield: 21.0%) was prepared in the same manner as in
20 Example 42, except that 4-(3,5-dimethyl-1H-pyrazol-1-yl)aniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.91-7.87(m, 2H), 7.30-7.28(m, 2H), 6.89-6.62(m, 1H), 6.27-6.10(m, 1H), 6.03(s, 1H), 5.79-5.58(m, 1H), 4.43-4.34(m, 1H), 4.01-3.87(m, 2H), 3.70-3.42(m, 2H), 2.20(s, 6H), 2.15-2.11(m, 1H), 1.95-1.85(m, 2H), 1.70-1.64(m, 1H)

25

Example 44: Preparation of (R)-1-(3-(3-chloro-6-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-

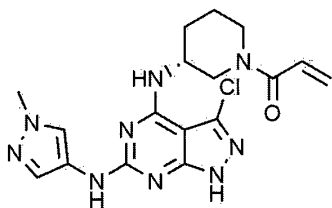
one



A title compound (10.1 mg, yield: 37.4%) was prepared in the same manner as in Example 42, except that 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.08-8.04(m, 1H), 7.65-7.61(m, 1H), 6.86-6.60(m, 1H), 6.27-6.01(m, 2H), 5.80-5.48(m, 1H), 4.58-4.30(m, 4H), 3.95-3.82(m, 1H), 3.65-3.38(m, 2H), 2.09-2.05(m, 1H), 1.92-1.85(m, 2H), 1.73-1.65(m, 1H)

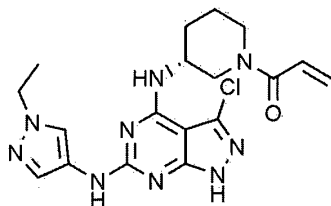
Example 45: Preparation of (R)-1-(3-(3-chloro-6-(1-methyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (10.1 mg, yield: 35.7%) was prepared in the same manner as in Example 42, except that 1-methyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.92(s, 1H), 7.55(s, 1H), 6.86-6.56(m, 1H), 6.26-6.10(m, 1H), 5.79-5.57(m, 1H), 4.43-4.29(m, 2H), 3.98-3.56(m, 5H), 3.46-3.41(m, 1H), 2.15-2.10(m, 1H), 1.91-1.86(m, 2H), 1.70-1.65(m, 1H)

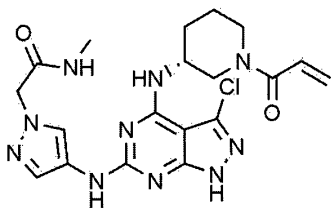
Example 46: Preparation of (R)-1-(3-(3-chloro-6-(1-ethyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (11.2 mg, yield: 38.6%) was prepared in the same manner as in Example 42, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

5 ^1H NMR (500MHz, CD_3OD) δ 7.96(s, 1H), 7.57(s, 1H), 6.86-6.56(m, 1H), 6.27-7.09(m, 1H), 5.80-5.56(m, 1H), 4.42-4.29(m, 2H), 4.14-4.09(m, 2H), 3.97-3.86(m, 1H), 3.66-3.41(m, 2H), 2.10-2.05(m, 1H), 1.93-1.86(m, 2H), 1.70-1.60(m, 1H), 1.43(s, 3H)

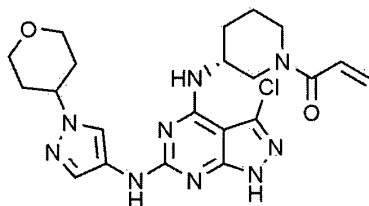
Example 47: Preparation of (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)-N-methylacetamide



A title compound (11.9 mg, yield: 37.2%) was prepared in the same manner as in Example 42, except that 2-(4-amino-1H-pyrazol-1-yl)-N-methyl acetamide was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

15 ^1H NMR (500MHz, CD_3OD) δ 8.06-8.02(m, 1H), 7.64-7.60(m, 1H), 6.85-6.60(m, 1H), 6.26-6.09(m, 1H), 5.79-5.50(m, 1H), 4.78(s, 2H), 4.42-4.32(m, 2H), 3.96-3.88(m, 1H), 3.81-3.41(m, 2H), 2.74(s, 3H), 2.25-2.10(m, 1H), 1.92-1.84(m, 2H), 1.70-1.65(m, 1H)

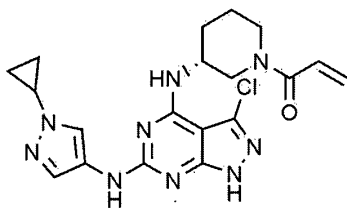
20 **Example 48: Preparation of (R)-1-(3-(3-chloro-6-(1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (79.3 mg, yield: 80.9%) was prepared in the same manner as in Example 42, except that 1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

5 ^1H NMR (500MHz, CD_3OD) δ 8.03(s, 1H), 7.59(s, 1H), 6.86-6.62(m, 1H), 6.27-6.09(m, 1H), 5.87-5.58(m, 1H), 4.38-4.32(m, 3H), 4.11-3.86(m, 4H), 3.59-3.44(m, 3H), 2.15-2.00(m, 5H), 1.92-1.87(m, 2H), 1.66-1.64(m, 1H)

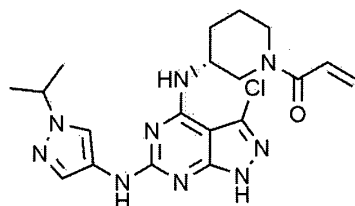
Example 49: Preparation of (R)-1-(3-(3-chloro-6-(1-cyclopropyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (20.2 mg, yield: 39.4%) was prepared in the same manner as in Example 42, except that 1-cyclo-1H-propyl-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

15 ^1H NMR (500MHz, CD_3OD) δ 8.00(s, 1H), 7.54(s, 1H), 6.86-6.61(m, 1H), 6.26-6.10(m, 1H), 5.79-5.58(m, 1H), 4.37-4.30(m, 1H), 3.97-3.85(m, 1H), 3.57(s, 1H), 3.45(s, 1H), 2.11(s, 1H), 1.87-1.67(m, 2H), 1.66(s, 1H), 1.32-1.28(m, 2H), 1.05(d, 4H)

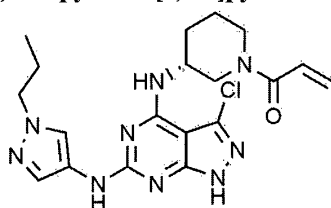
Example 50: Preparation of (R)-1-(3-(3-chloro-6-(1-isopropyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (5 mg, yield: 17.0%) was prepared in the same manner as in Example 42, except that 1-isopropyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

5 ^1H NMR (500MHz, CD_3OD) δ 7.97(s, 1H), 7.58(s, 1H), 6.86-6.67(m, 1H), 6.26-6.09(m, 1H), 5.80-5.57(m, 1H), 4.47-4.31(m, 2H), 3.97-3.85(m, 2H), 3.59-3.45(m, 1H), 2.10-2.00(m, 1H), 1.91-1.87(m, 2H), 1.73-1.66(m, 1H), 1.48(s, 6H)

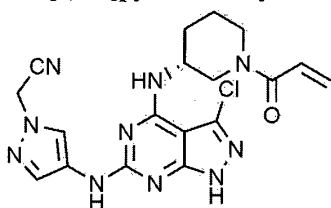
Example 51: Preparation of (R)-1-(3-(3-chloro-6-(1-propyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (47.7 mg, yield: 48.7%) was prepared in the same manner as in Example 42, except that 1-propyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

15 ^1H NMR (500MHz, CD_3OD) δ 7.96(s, 1H), 7.57(s, 1H), 6.87-6.63(m, 1H), 6.27-6.10(m, 1H), 5.80-5.57(m, 1H), 4.43-4.30(m, 2H), 4.11-3.87(m, 3H), 3.66-3.44(m, 2H), 2.15-2.00(s, 1H), 1.90-1.84(m, 5H), 1.70-1.65(m, 1H), 0.90-0.87(m, 3H)

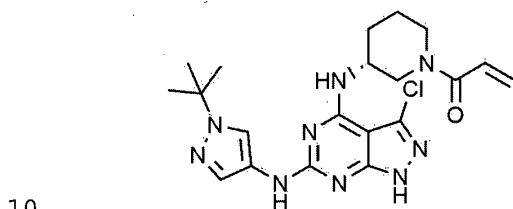
Example 52: Preparation of (R)-2-(4-(4-(1-acryloylpiperidin-3-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-ylamino)-1H-pyrazol-1-yl)acetonitrile



A title compound (52.8 mg, yield: 53.9%) was prepared in the same manner as in Example 42, except that 2-(4-amino-1H-pyrazol-1-yl)acetonitrile was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.16-8.10(m, 1H), 7.68-7.63(m, 1H), 6.85-6.65(m, 1H),
 5 6.28-6.10(m, 1H), 5.81-5.58(m, 1H), 5.30-5.26(m, 2H), 4.58-4.39(m, 2H), 3.96-3.82(m, 1H),
 3.68-3.44(m, 2H), 2.15-2.05(m, 1H), 1.89-1.85(m, 2H), 1.75-1.69(m, 1H)

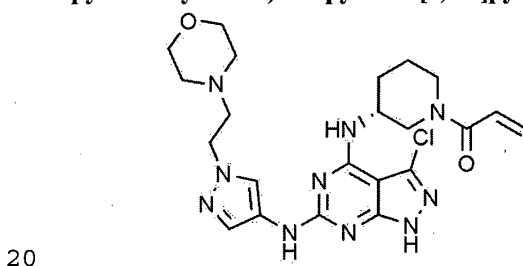
Example 53: Preparation of (R)-1-(3-(6-(1-*tert*-butyl-1H-pyrazol-4-ylamino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (27.3 mg, yield: 51.2%) was prepared in the same manner as in Example 42, except that 1-(*tert*-butyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.05(s, 1H), 7.61(s, 1H), 6.85-6.61(m, 1H), 6.26-6.09(m,
 15 1H), 5.79-5.58(m, 1H), 4.37-4.31(m, 1H), 3.97-3.84(m, 1H), 3.62-3.47(m, 1H), 2.09-2.07(m, 1H),
 1.97-1.84(m, 2H), 1.56(s, 9H), 1.34-1.28(m, 3H)

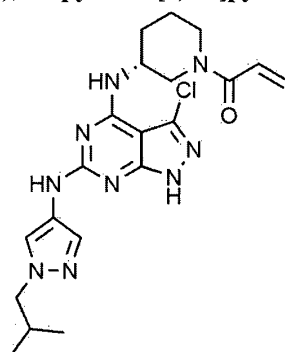
Example 54: Preparation of (R)-1-(3-(3-chloro-6-(1-(2-morpholinoethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (42.0 mg, yield: 69.0%) was prepared in the same manner as in Example 42, except that 1-(2-morpholinoethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.98(s, 1H), 7.61(s, 1H), 6.85-6.62(m, 1H), 6.27-6.10(m, 1H), 5.80-5.56(m, 1H), 4.40-4.35(m, 2H), 4.28-4.22(m, 2H), 3.92-3.85(m, 1H), 3.66(s, 4H), 3.51-3.42(m, 2H), 2.79(t, 2H), 2.48(s, 4H), 2.13-2.07(m, 1H), 1.91-1.86(m, 2H), 1.65-1.23(m, 2H)

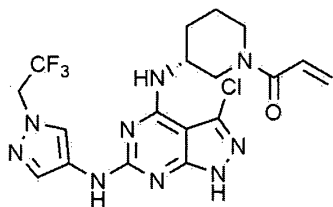
5 **Example 55: Preparation of (R)-1-(3-(3-chloro-6-(1-isobutyl-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (12.6 mg, yield: 23.6%) was prepared in the same manner as in Example 42, except that 1-isobutyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.95(s, 1H), 7.58(s, 1H), 6.87-6.62(m, 1H), 6.28-6.11(m, 1H), 5.81-5.57(m, 1H), 4.44-4.29(m, 2H), 3.67-3.94(m, 1H), 3.88(d, 2H), 3.65-3.41(m, 2H), 2.15-2.11(m, 2H), 1.93-1.87(m, 2H), 1.65-1.63(m, 1H), 0.91-0.90(d, 6H)

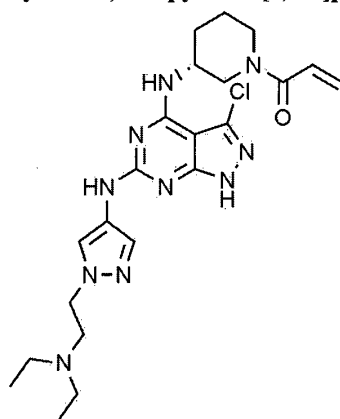
15 **Example 56: Preparation of (R)-1-(3-(3-chloro-6-(1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (55.8 mg, yield: 56.9%) was prepared in the same manner as in Example 42, except that 1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.16-8.10(m, 1H), 7.68-7.63(m, 1H), 6.87-6.65(m, 1H), 6.28-6.10(m, 1H), 5.81-5.59(m, 1H), 4.93-4.91(m, 2H), 4.78-4.29(m, 2H), 3.95-3.54(m, 2H), 3.44-3.38(m, 1H), 2.09-2.05(m, 1H), 1.89-1.75(m, 2H), 1.70-1.65(m, 1H)

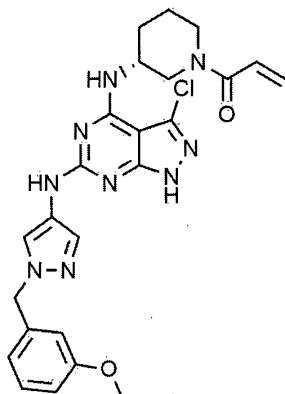
5 **Example 57: Preparation of (R)-1-(3-(3-chloro-6-(1-(2-(diethylamino)ethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (27.8 mg, yield: 47.6%) was prepared in the same manner as in Example 42, except that 1-(2-(diethylamino)ethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.03(s, 1H), 7.64(s, 1H), 6.86-6.12(m, 1H), 6.26-6.10(m, 1H), 5.80-5.56(m, 1H), 4.43-4.30(m, 3H), 3.97-3.86(m, 2H), 3.42(d, 4H), 3.21-3.17(m, 2H), 2.86-2.83(m, 2H), 2.10(s, 1H), 2.03-2.01(m, 2H), 1.67(s, 1H), 1.61-1.15(m, 6H)

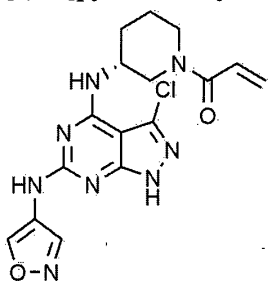
15 **Example 58: Preparation of (R)-1-(3-(3-chloro-6-(1-(3-methoxybenzyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one**



A title compound (23.4 mg, yield: 38.4%) was prepared in the same manner as in Example 42, except that 1-(3-methoxybenzyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.04(s, 1H), 7.61(s, 1H), 7.25(t, 1H), 6.85-6.59(m, 4H), 6.24-6.08(m, 1H), 5.77-5.56(m, 1H), 4.33-4.23(m, 3H), 3.75(s, 3H), 3.67-3.44(m, 2H), 2.03(s, 1H), 1.88-1.80(m, 2H), 1.63-1.60(m, 1H)

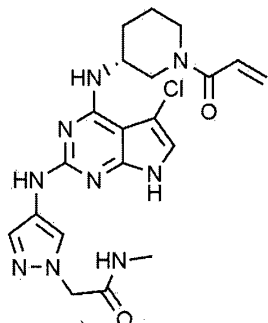
Example 59: Preparation of (R)-1-(3-(3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (23.1 mg, yield: 49.6%) was prepared in the same manner as in Example 42, except that isoxazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 9.04(s, 1H), 8.49(s, 1H), 6.85-6.62(m, 1H), 6.27-6.11(m, 1H), 5.80-5.58(m, 1H), 4.37-4.29(m, 2H), 3.97-3.95(m, 1H), 3.68-3.48(m, 2H), 2.10(s, 1H), 1.95-1.84(m, 2H), 1.66-1.65(m, 1H)

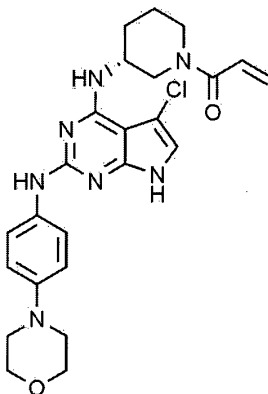
Example 60: Preparation of (R)-2-(4-((1-(acryloylpiperidin-3-yl)amino)-5-

chloro-7H-pyrrolo[2,3-d]pyrimidin-2-yl)amino)-1H-pyrazol-1-yl)-N-methylacetamide

A title compound (5.9 mg, yield: 22.8%) was prepared in the same manner as in Example 36, except that 2-(4-amino-1H-pyrazol-1-yl)-N-methylacetamide was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.98-8.05(m, 1H), 7.57-7.62(m, 1H), 6.52-6.86(m, 2H), 6.07-6.24(m, 1H), 5.50-5.77(m, 1H), 4.76-4.78(m, 2H), 4.25-4.50(m, 2H), 3.83-3.90(m, 1H), 3.78-3.80(m, 1H), 3.59-3.68(m, 1H), 3.39-3.48(m, 1H), 2.75(s, 3H), 1.95-2.01(m, 1H), 1.85-1.87(m, 1H), 1.58-1.68(m, 1H)

10

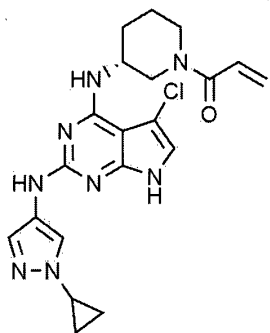
Example 61: Preparation of (R)-1-(3-((5-chloro-2-((4-morpholinophenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

A title compound (7.4 mg, yield: 27.2%) was prepared in the same manner as in Example 36, except that 4-morpholinoaniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.51-7.55(m, 2H), 6.89-6.90(m, 2H), 6.46-6.85(m, 2H), 6.03-6.25(m, 1H), 5.45-5.80(m, 1H), 3.96-4.18(m, 1H), 3.88-3.94(m, 1H), 3.80-3.82(m, 4H),

3.70-3.78(m, 1H), 3.53-3.66(m, 1H), 3.41-3.50(m, 1H), 3.35-3.40(m, 1H), 3.00-3.05(m, 4H), 1.97-2.02(m, 1H), 1.83-1.85(m, 1H), 1.59-1.65(m, 1H)

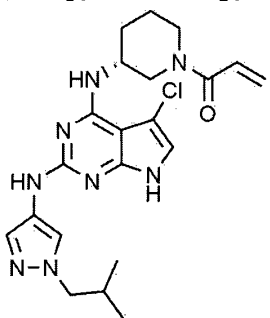
Example 62: Preparation of (R)-1-(3-((5-chloro-2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.2 mg, yield: 29.8%) was prepared in the same manner as in Example 36, except that 1-cyclo-1H-propyl-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.97-7.93(m, 1H), 7.49-7.45(m, 1H), 6.84-6.52(m, 2H), 6.24-6.06(m, 1H), 5.77-5.50(m, 1H), 4.31-4.27(m, 1H), 3.93-3.37(m, 5H), 2.08-2.05(m, 1H), 1.80-1.90(m, 1H), 1.55-1.70(m, 1H), 1.03-0.98(m, 4H)

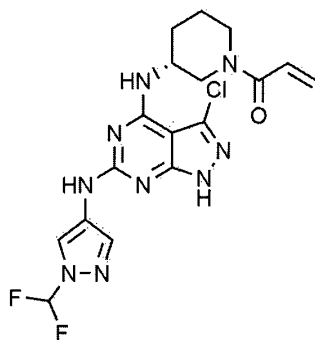
Example 63: Preparation of (R)-1-(3-((5-chloro-2-((1-isobutyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (4.9 mg, yield: 19.6%) was prepared in the same manner as in Example 36, except that 1-isobutyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.93-7.88(m, 1H), 7.55-7.51(m, 1H), 6.89-6.53(m, 2H), 6.26-6.07(m, 1H), 5.79-5.51(m, 1H), 4.38-3.42(m, 7H), 2.18-2.08(m, 3H), 1.90-1.84(m, 1H), 1.70-1.55(m, 1H)

5 **Example 64: Preparation of (R)-1-(3-((3-chloro-6-((1-(difluoromethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one**

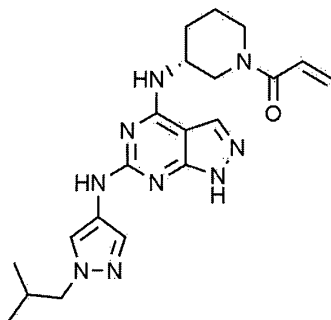


A title compound (24.0 mg, yield: 36.9%) was prepared in the same manner as in
10 Example 42, except that 1- (difluoromethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.40(s, 1H), 7.79(d, 1H), 7.50-7.24(m, 1H), 6.85-6.60(m, 1H), 6.26-6.06(m, 1H), 5.80-5.54(m, 1H), 4.29-4.20(m, 1H), 4.08-4.04(m, 2H), 3.52-3.42(m, 1H), 2.21-2.15(m, 1H), 1.96-1.80(m, 2H), 1.70-1.63(m, 1H)

15

Example 65: Preparation of (R)-1-(3-(((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

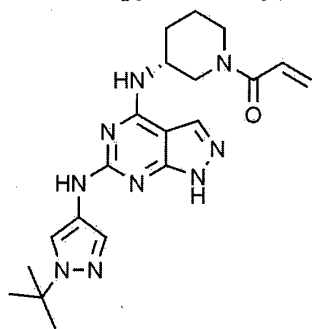


A title compound (26.0 mg, yield: 49.1%) was prepared in the same manner as in

Example 30, except that 1-isobutyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 7.99(m, 2H), 7.58(d, 1H), 6.72-6.60(m, 1H), 6.25-6.13(m, 1H), 5.73-5.61(m, 1H), 4.45-4.27(m, 2H), 3.69-3.94(m, 1H), 3.87(d, 2H), 3.68-3.41(m, 2H), 2.17-2.13(m, 2H), 1.93-1.89(m, 2H), 1.67-1.64(m, 1H), 0.92-0.90(d, 6H)

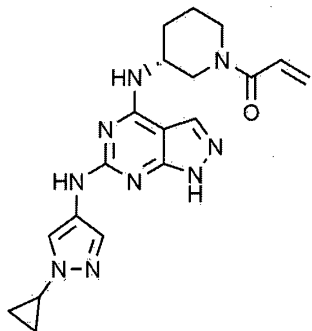
Example 66: Preparation of (R)-1-(3-(((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (27.0 mg, yield: 50.9%) was prepared in the same manner as in Example 30, except that 1-(*tert*-butyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.07(d, 1H), 8.06(d, 1H), 7.90(d, 1H), 6.84-6.57(m, 1H), 6.25-6.08(m, 1H), 5.80-5.59(m, 1H), 4.42-4.34(m, 1H), 3.98-3.83(m, 1H), 3.62-3.50(m, 1H), 2.12-2.09(m, 1H), 1.99-1.85(m, 2H), 1.57(s, 9H), 1.35-1.29(m, 3H)

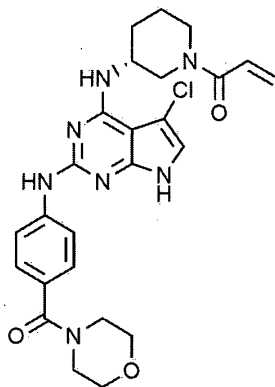
Example 67: Preparation of (R)-1-(3-(((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (26.0 mg, yield: 50.9%) was prepared in the same manner as in Example 30, except that 1-cyclo-1H-propyl-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.02(d, 1H), 8.00(s, 1H), 7.91(d, 1H), 6.81-6.51(m, 1H),
 5 6.27-6.12(m, 1H), 5.78-5.53(m, 1H), 4.26-4.10(m, 1H), 3.98-3.84(m, 1H), 3.56(s, 1H), 3.48(s, 1H), 2.18(s, 1H), 1.89-1.65(m, 2H), 1.67(s, 1H), 1.32-1.21(m, 2H), 1.04(d, 4H)

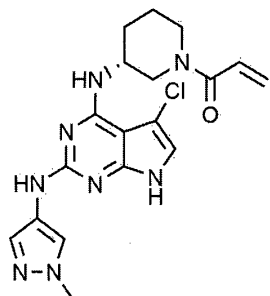
Example 68: Preparation of (R)-1-(3-((5-chloro-2-((4-(morpholine-4-carbonyl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (2.5 mg, yield: 8.7%) was prepared in the same manner as in Example 36, except that (4-aminophenyl)(morpholino)methanone was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.85-7.81(m, 2H), 7.36-7.34(m, 2H), 6.86-6.54(m, 2H),
 15 6.26-6.04(m, 1H), 5.79-5.47(m, 1H), 4.38-4.31(m, 1H), 3.93-3.46(12H), 2.10-1.95(m, 2H), 1.88-1.86(m, 1H), 1.67-1.59(m, 1H)

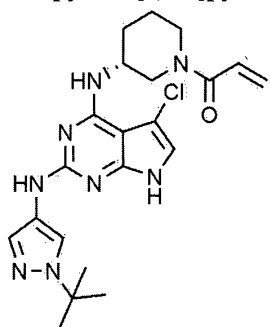
Example 69: Preparation of (R)-1-(3-((5-chloro-2-((1-methyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (2.1 mg, yield: 9.3%) was prepared in the same manner as in Example 36, except that 1-methyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

5 ^1H NMR (500MHz, CD_3OD) δ 7.92-7.87(m, 1H), 7.52-7.45(m, 1H), 6.83-6.51(m, 2H), 6.25-6.06(m, 1H), 5.78-5.49(m, 1H), 4.60-4.50(m, 1H), 4.39-4.27(m, 1H), 3.93-3.45(m, 6H), 2.10-2.02(m, 2H), 1.88-1.86(m, 1H), 1.68-1.59(m, 1H)

Example 70: Preparation of (R)-1-(3-((2-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-5-chloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

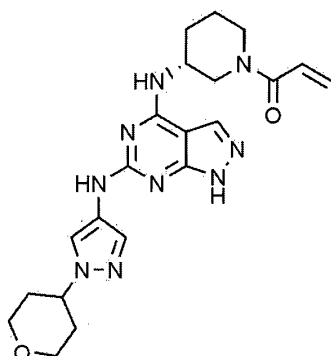


A title compound (5.1 mg, yield: 20.4%) was prepared in the same manner as in Example 36, except that 1-(*tert*-butyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

15 ^1H NMR (500MHz, CD_3OD) δ 8.04-8.02(m, 1H), 7.58-7.56(m, 2H), 6.82-6.55(m, 2H), 6.23-6.05(m, 1H), 5.78-5.50(m, 1H), 4.35-3.62(m, 4H), 3.54-3.44(m, 1H), 2.08-2.00(m, 2H), 1.95-1.78(m, 1H), 1.88-1.85(m, 1H), 1.56(s, 9H)

Example 71: Preparation of (R)-1-(3-((6-((1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-

one



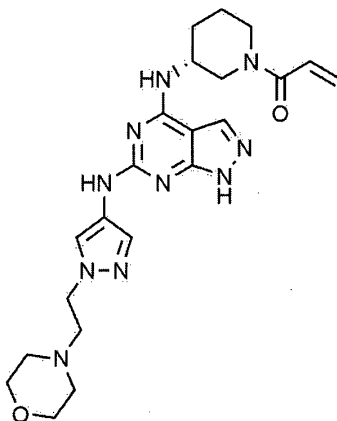
A title compound (22.0 mg, yield: 39.2%) was prepared in the same manner as in Example 30, except that 1-(tetrahydro-2H-pyran-4-yl)-1H-pyrazol-4-amine was used in Example

5 30.

^1H NMR (500MHz, CD_3OD) δ 8.05(d, 1H), 7.91(s, 1H), 7.60(d, 1H), 6.86-6.57(m, 1H), 6.26-6.06(m, 1H), 5.79-5.54(m, 1H), 4.31-4.21(m, 1H), 4.10-4.04(m, 4H), 3.59-3.53(m, 2H), 3.18(t, 1H), 2.20-2.17(m, 1H), 2.03(s, 4H), 1.95-1.93(m, 1H), 1.79-1.70(m, 1H), 1.56-1.52(m, 2H)

10

Example 72: Preparation of (R)-1-(3-((6-((1-(2-morpholinoethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



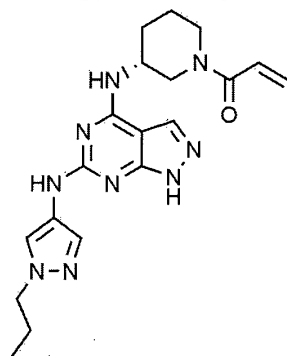
A title compound (21.0 mg, yield: 35.0%) was prepared in the same manner as in Example 30, except that 1-(2-morpholinoethyl)-1H-pyrazol-4-amine was used in Example 30.

15

^1H NMR (500MHz, CD_3OD) δ 8.01(d, 1H), 7.91(s, 1H), 7.60(d, 1H), 6.85-6.58(m, 1H), 6.27-6.07(m, 1H), 5.79-5.54(m, 1H), 4.22-4.21(m, 2H), 4.11-4.05(m, 2H), 3.66(s, 4H), 3.43-3.43(m, 1H), 2.79(t, 1H), 2.48(s, 4H), 2.20-2.17(m, 1H), 1.79-1.67(m, 1H), 1.66-1.64(m, 1H),

1.63-1.61(m, 2H)

Example 73: Preparation of (R)-1-(3-(((6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

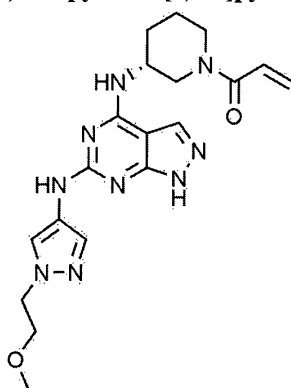


5

A title compound (21.0 mg, yield: 43.1%) was prepared in the same manner as in Example 30, except that 1-propyl-1H-pyrazol-4-amine was used in Example 30.

¹H NMR (500MHz, CD₃OD) δ 7.99(d, 1H), 7.90(s, 1H), 7.57(d, 1H), 6.86-6.59(m, 1H), 6.27-6.08(m, 1H), 5.79-5.55(m, 1H), 4.27-4.21(m, 1H), 4.05-4.03(m, 2H), 3.44-3.16(m, 2H),
 10 2.19-2.17(m, 1H), 1.95-1.93(m, 1H), 1.84-1.80(m, 2H), 1.79-1.67(m, 1H), 1.65-1.60(m, 2H), 0.91-0.90(m, 3H)

Example 74: Preparation of (R)-1-(3-(((6-((1-(2-methoxyethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



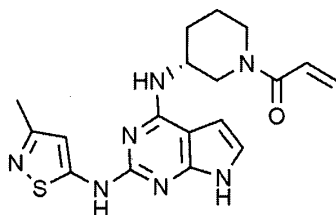
15

A title compound (20.7 mg, yield: 37.7%) was prepared in the same manner as in Example 30, except that 1-(2-methoxyethyl)-1H-pyrazol-4-amine was used in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.02(d, 1H), 7.91(s, 1H), 7.59(d, 1H), 6.85-6.57(m, 1H), 6.26-6.07(m, 1H), 5.79-5.53(m, 1H), 4.26-4.22(m, 3H), 4.07-4.05(m, 3H), 3.72-3.70(m, 2H), 3.48-3.44(m, 1H), 3.40-3.22(m, 2H), 2.19-2.17(m, 1H), 2.01-1.95(m, 1H), 1.80-1.71(m, 1H), 1.67-1.56(m, 2H)

5

Example 75: Preparation of (R)-1-(3-((2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 2,4-dichloro-7-tosyl-7H-pyrrolo [2,3-d]pyrimidine

10 After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (3.0 g, 16.0 mmol) was dissolved in acetone (20.0 mL), 4-methylbenzenesulfonyl chloride (4.6 g, 23.9 mmol) was added thereto. After cooling to 0 °C, 2 M sodium hydroxide solution (12.0 mL) was slowly added dropwise and then stirred at room temperature for 2 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was
15 isolated by column chromatography to obtain a title compound (2.9 g, yield: 80.0%).

¹H NMR (500MHz,CD₃OD) δ 8.12(d, 2H), 7.76(d, 1H), 7.37(d, 2H), 6.68(d, 1H), 2.43(s, 3H)

Step 2: Preparation of *tert*-butyl (R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

20 After 2,4-dichloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine (500.0 mg, 1.5 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (382.0 μL, 2.2 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (322.0 mg, 1.6 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with
25 magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (681.0 mg, yield: 92.0%).

¹H NMR (500MHz, CD₃OD) δ 8.10(d, 1H), 7.39(d, 1H), 7.31-7.23(m, 4H), 4.17-

4.13(m, 1H), 3.70-3.60(m, 1H), 3.45-3.35(m, 3H), 2.40(s, 3H), 1.95-1.85(m, 1H), 1.70-1.65(m, 1H), 1.60-1.55(m, 2H), 1.40-1.37(m, 9H)

Step 3: Preparation of *tert*-butyl (R)-3-((2-((3-methylisothiazol-5-yl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After *tert*-butyl (R)-3-((2-chloro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (300.0 mg, 0.6 mmol) was dissolved in anhydrous *tert*-butanol (2.0 mL), 3-methylisothiazol-5-amine (67.7 mg, 0.6 mmol), tris(dibenzylideneacetone)dipalladium (27 mg, 0.03 mmol), 2'-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl (28.3 mg, 0.06 mmol), and potassium carbonate (163.9 mg, 1.2 mmol) were added thereto. Then, the mixture was reacted at 110 °C for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (223.0 mg, yield: 64.5%).

¹H NMR (500MHz, CD₃OD) δ 8.04(d, 2H), 7.35-7.28(m, 3H), 6.72(s, 1H), 6.59(s, 1H), 4.40-4.28(m, 1H), 4.05-3.80(m, 2H), 3.20-2.80(m, 2H), 2.36(s, 3H), 2.33(s, 3H), 2.15-2.12(m, 1H), 2.00-1.80(m, 1H), 1.75-1.67(m, 2H), 1.10-1.00(m, 9H)

Step 4: Preparation of *tert*-butyl (R)-3-((2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

Tert-butyl (R)-3-((2-((3-methylisothiazol-5-yl)amino)-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (122.0 mg, 0.2 mmol) was dissolved in methanol (1.0 mL). Potassium hydroxide (23.5 mg, 0.4 mmol) was added to the reaction mixture and then stirred at 50 °C for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (60.0 mg, yield: 66.5%).

¹H NMR (500MHz, CD₃OD) δ 6.84(s, 1H), 6.53(s, 1H), 6.48(s, 1H), 4.50-4.25(m, 1H), 4.10-3.90(m, 1H), 3.80-3.70(m, 1H), 3.20-2.70(m, 2H), 2.31(s, 3H), 2.25-2.15(m, 1H), 1.90-1.80(m, 1H), 1.75-1.70(m, 2H), 1.64-1.23(m, 9H)

Step 5: Preparation of (R)-N2-(3-methylisothiazol-5-yl)-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride

To *tert*-butyl (R)-3-((2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (60.0 mg, 0.14 mmol) was added 6 N hydrochloric acid solution (2.0 mL, excess) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

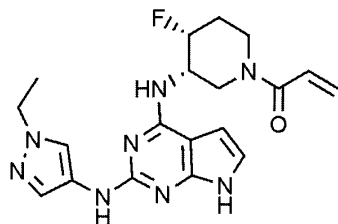
¹H NMR (500MHz, CD₃OD) δ 6.84(s, 1H), 6.53(s, 1H), 6.48(s, 1H), 4.50-4.25(m, 1H), 4.10-3.90(m, 1H), 3.80-3.70(m, 1H), 3.20-2.70(m, 2H), 2.31(s, 3H), 2.25-2.15(m, 1H), 1.90-1.80(m, 1H), 1.75-1.70(m, 2H)

Step 6: Preparation of (R)-1-(3-((2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-N2-(3-methylisothiazol-5-yl)-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride (34.5 mg, 0.07 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (2.5 mL), sodium bicarbonate (16.8 mg, 0.07 mmol) was added thereto at -20 °C and then stirred for 30 minutes. Acryloyl chloride (6.3 μL, 0.07 mmol) was slowly added dropwise to the reaction mixture, and then stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (3.0 mg, yield: 8.6%).

¹H NMR (500MHz, CD₃OD) δ 6.85(s, 1H), 6.84-6.48(m, 3H), 6.27-6.05(m, 1H) 5.78-5.55(m, 1H), 4.59-4.49(m, 1H), 4.17-4.07(m, 1H), 3.25-3.15(m, 2H), 2.31(s, 3H), 2.27-2.15(m, 1H), 2.03-1.89(m, 2H), 1.79-1.59(m, 1H)

Example 76: Preparation of 1-((3S,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one

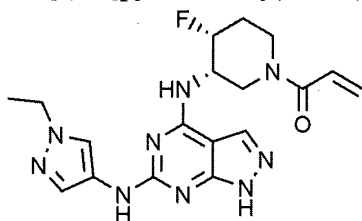


A title compound (15.5 mg, yield: 36.6%) was prepared in the same manner as in Example 1, except using 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.89(d, 1H), 7.50(d, 1H), 6.85-6.75(m, 2H), 6.46(s, 1H), 6.28-6.14(m, 1H), 5.81-5.62(m, 1H), 5.12-5.02(m, 1H), 4.80-4.30(m, 2H), 4.09-4.03(m, 3H), 3.51-3.43(m, 1H), 3.14-3.05(m, 1H), 2.21-2.18(m, 1H), 1.96-1.85(m, 1H), 1.43-1.39(m, 3H)

10

Example 77: Preparation of 1-((3S,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one



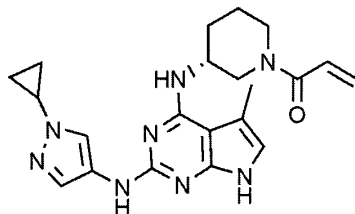
A title compound (7.1 mg, yield: 20.0%) was prepared in the same manner as in Example 30, except using 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 30.

¹H NMR (500MHz, CD₃OD) δ 7.97(m, 2H), 7.54(d, 1H), 6.86-6.60(m, 1H), 6.29-6.15(m, 1H), 5.81-5.79(m, 1H), 5.09(d, 1H), 4.72-4.25(m, 2H), 4.13-4.03(m, 3H), 3.57-3.45(m, 1H), 3.21-3.08(m, 1H), 2.22-2.18(m, 1H), 1.94-1.86(m, 1H), 1.50-1.40(m, 3H)

20

Example 78: Preparation of (R)-1-(3-((2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

one



Step 1: Preparation of *tert*-butyl (R)-3-((2-chloro-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

5 After 2,4-dichloro-5-methyl-7H-pyrrolo[2,3-d]pyrimidine (300.0 mg, 1.5 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695.0 μ L, 2.2 mmol) and *tert*-butyl-(R)-3-aminopiperidine-1-carboxylate (356.9 mg, 1.8 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was
10 isolated by column chromatography to obtain a title compound (461.6 mg, yield: 85.3%).

¹H NMR (500MHz, CD₃OD) δ 6.84(s, 1H), 4.57-4.50(m, 1H), 3.63-3.61(m, 1H), 3.44-3.34(m, 1H), 3.03-2.93(m, 2H), 2.42(s, 3H), 2.17-2.15(m, 1H), 2.09-2.06(m, 1H), 1.98-1.86(m, 2H), 1.50-1.30(m, 9H)

Step 2: Preparation of (R)-2-chloro-5-methyl-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride

To *tert*-butyl (R)-3-((2-chloro-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (461.6 mg, 1.3 mmol) was added 6 N hydrochloric acid solution (2.0 mL, excess) dissolved in methanol. After stirring at room temperature for 30
20 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 6.84(s, 1H), 4.57-4.50(m, 1H), 3.63-3.61(m, 1H), 3.44-3.34(m, 1H), 3.03-2.93(m, 2H), 2.42(s, 3H), 2.17-2.15(m, 1H), 2.09-2.06(m, 1H), 1.98-1.86(m, 2H)

25

Step 3: Preparation of (R)-1-(3-((2-chloro-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-2-chloro-5-methyl-N-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-4-amine hydrochloride (387.8 mg, 1.3 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4 mL), sodium bicarbonate (323.4 mg, 3.9 mmol) was added thereto at -20 °C and then stirred for 30 minutes. Acryloyl chloride (121.4 µL, 1.4 mmol) was slowly added dropwise to the reaction mixture, and then stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (210.4 mg, yield: 61.0%).

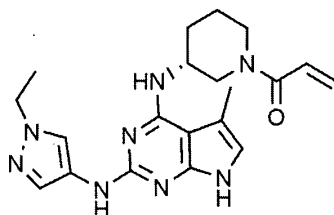
¹H NMR (500MHz, CD₃OD) δ 6.88-6.76(m, 2H), 6.24-6.14(m, 1H), 5.78-5.65(m, 1H), 4.34-4.27(m, 1H), 4.20-3.62(m, 3H), 3.26-3.16(m, 1H), 2.37(d, 3H), 2.11-2.00(m, 1H), 1.89-1.84(m, 2H), 1.70-1.65(m, 1H)

Step 4: Preparation of (R)-1-(3-((2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-yl-1-one

(R)-1-(3-((2-chloro-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (30.0 mg, 0.09 mmol) and 1-cyclopropyl-1H-pyrazol-4-amine (9.6 mg, 0.08 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (5.8 µL, 0.08 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 3 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to obtain a title compound (3.6 mg, yield: 11.3%).

¹H NMR (500MHz, CD₃OD) δ 7.95(d, 1H), 7.49(s, 1H), 6.90-6.48(m, 2H), 6.26-6.05(m, 1H), 5.49-5.50(m, 1H), 4.33-4.31(m, 1H), 4.03-3.80(m, 2H), 3.54-3.38(m, 3H), 2.32(s, 3H), 2.13-2.06(m, 1H), 1.95-1.85(m, 2H), 1.75-1.67(m, 1H), 1.04-0.92(m, 4H)

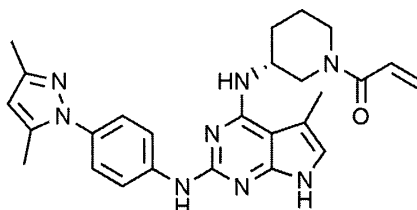
Example 79: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (2.8 mg, yield: 8.43%) was prepared in the same manner as in Example 78, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 78.

¹H NMR (500MHz, CD₃OD) δ 7.91(d, 1H), 7.51(d, 1H), 6.90-6.48(m, 2H), 6.27-6.05(d, 1H), 5.80-5.50(d, 1H), 4.36-4.32(d, 2H), 4.15-4.09(m, 2H), 4.00-3.56(m, 3H), 2.32(s, 3H), 2.15-2.11(m, 1H), 1.94-1.85(m, 2H), 1.70-1.65(m, 1H), 1.44-1.40(m, 3H)

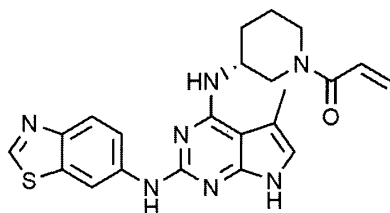
Example 80: Preparation of (R)-1-(3-((2-((4-(3,5-dimethyl-1H-pyrazol-1-yl)phenyl)amino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.1 mg, yield: 25.1%) was prepared in the same manner as in Example 78, except that 4-(3,5-dimethyl-1H-pyrazol-1-yl)aniline was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 78.

¹H NMR (500MHz, CD₃OD) δ 7.86-7.82(m, 2H), 7.25-7.23(m, 2H), 6.85-6.54(m, 2H), 6.26-6.01(m, 2H), 5.78-5.48(m, 1H), 4.36-4.34(m, 2H), 3.84-3.82(m, 1H), 3.48-3.39(m, 2H), 2.35(s, 3H), 2.22(s, 6H), 2.18-2.15(m, 1H), 1.92-1.83(m, 2H), 1.70-1.65(m, 1H)

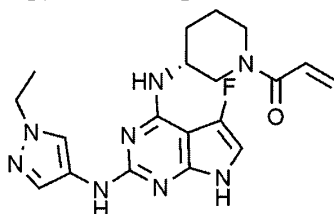
Example 81: Preparation of (R)-1-(3-((2-(benzo[d]thiazol-6-ylamino)-5-methyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (2.6 mg, yield: 10.0%) was prepared in the same manner as in Example 78, except that benzo[d]thiazol-6-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 78.

¹H NMR (500MHz, CD₃OD) δ 8.98(s, 1H), 8.79-8.73(m, 1H), 7.88-7.86(m, 1H), 7.68-7.57(m, 1H), 6.85-6.56(m, 3H), 6.30-5.97(m, 1H), 5.79-5.34(m, 1H), 4.37-4.27(m, 1H), 4.02-3.85(m, 1H), 3.60-3.48(m, 2H), 2.89-2.77(m, 1H), 2.35(s, 3H), 2.15-2.00(m, 1H), 1.92-1.86(m, 1H), 1.77-1.67(m, 2H)

Example 82: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 2,4-dichloro-5-fluoro-7H-pyrrolo[2,3-d]pyrimidine

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (200 mg, 1.06 mmol) was dissolved in acetonitrile (5.0 mL), 1-chloromethyl-4-fluoro-1,4-diazoniabicyclo[2.2.2]octane bis(tetrafluoroborate) (561.6 mg, 1.6 mmol) and acetic acid (1 mL) were added thereto. The mixture was heated at 80 °C and stirred for 24 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (170.0 mg, yield: 80.1%).

¹H NMR (500MHz, CD₃OD) δ 7.36(s, 1H)

Step 2: Preparation of 2,4-dichloro-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidine

After 2,4-dichloro-5-fluoro-7H-pyrrolo[2,3-d]pyrimidine (100.0 mg, 0.5 mmol) was dissolved in N,N-dimethylformamide (2.0 mL), sodium hydride (29.1 mg, 0.7 mmol) was added thereto and then stirred for 30 minutes. To the reaction mixture was added 2-(chloromethoxy)ethyltrimethylsilane (127.0 μ L, 0.7 mmol) and stirred at room temperature for 2 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (138.7 mg, yield: 85.1%).

^1H NMR (500MHz, CD_3OD) δ 7.56(s, 1H), 5.59(s, 2H), 3.57(t, 2H), 0.90(t, 2H), 0.00(s, 9H)

Step 3: Preparation of *tert*-butyl (R)-3-((2-chloro-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After 2,4-dichloro-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidine (138.7 mg, 0.4 mmol) was dissolved in ethanol (100 mL), N,N-diisopropylethylamine (107.8 μ L, 0.6 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (123.9 mg, 0.6 mmol) were added thereto. The reaction mixture was stirred at 110 $^\circ\text{C}$ for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (109.2 mg, yield: 53.2%).

^1H NMR (500MHz, CD_3OD) δ 7.01(s, 1H), 5.44(s, 2H), 4.25-4.20(m, 1H), 3.93-3.90(m, 1H), 3.66-3.60(m, 1H), 3.53(t, 2H), 3.20-3.13(m, 2H), 2.05-2.00(m, 1H), 1.80-1.77(m, 2H), 1.60-1.57(m, 1H), 1.50-1.38(m, 9H), 0.88(t, 2H), 0.00(s, 9H)

Step 4: Preparation of *tert*-butyl (R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

After *tert*-butyl (R)-3-((2-chloro-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (109.2 mg, 0.2 mmol) was dissolved in anhydrous *tert*-butanol (2.0 mL), 1-ethyl-1H-pyrazol-4-amine (20.2 mg, 0.2

mmol), *tris*(dibenzylideneacetone)dipalladium (8.3 mg, 0.01 mmol), 2-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl (8.7 mg, 0.02 mmol) and potassium carbonate (50.3 mg, 0.4 mmol) were added thereto. The mixture was reacted at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced
5 pressure. The residue was isolated by column chromatography to obtain a title compound (67.0 mg, yield: 65.0%).

¹H NMR (500MHz, CD₃OD) δ 7.96(s, 1H), 7.58(s, 1H) 6.63(s, 1H), 5.42(s, 2H), 4.59-4.55(m, 1H), 4.22-4.11(m, 3H), 3.82-3.80(m, 1H), 3.70-3.41(m, 3H), 3.20-2.92(m, 1H), 2.04-2.00(m, 1H), 1.77-1.70(m, 2H), 1.57-1.25(m, 13H), 0.90(t, 2H), 0.00(s, 9H)

10

Step 5: Preparation of (R)-N2-(1-ethyl-1H-pyrazol-4-yl)-5-fluoro-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride

To *tert*-butyl (R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-
15 carboxylate (67.0 mg, 0.15 mmol) was added 6 N hydrochloric acid solution (3.0 mL, excess) dissolved in methanol. After stirring at 60 °C to 70 °C for 2 hours, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 8.05-7.40(m, 3H), 6.90-6.40(m, 1H), 6.30-6.03(m, 1H), 5.85-5.55(m, 1H), 4.70-2.90(m, 7H), 2.15-2.00(m, 1H), 1.95-1.70(m, 2H), 1.69-1.55(m,
20 1H), 1.50-1.35(m, 3H)

Step 6: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

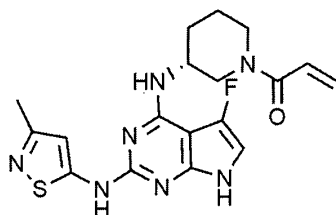
After (R)-N2-(1-ethyl-1H-pyrazol-4-yl)-5-fluoro-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-
25 d]pyrimidin-2,4-diamine hydrochloride (67.5 mg, 0.2 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4.0 ml), sodium bicarbonate (44.7 mg, 0.5 mmol) was added thereto at -20 °C and the mixture was stirred for 30 minutes. Acryloyl chloride (15.1 μL, 0.2 mmol) was slowly added dropwise to the reaction mixture and was stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then
30 concentrated under reduced pressure. The residue was isolated by column chromatography to

obtain a title compound (4.3 mg, yield: 6.1%).

¹H NMR (500MHz, CD₃OD) δ 8.00-7.97(m, 1H), 7.57-7.45(m, 1H), 6.90-6.40(m, 1H), 6.27-6.05(m, 1H), 5.79-5.48(m-2H), 4.23-4.01(m, 4H), 3.93-3.83(m, 2H), 2.99-2.97(m, 1H), 2.15-2.00(m, 1H), 1.91-1.75(m, 2H), 1.70-1.61(m, 1H), 1.44-1.37(m, 3H)

5

Example 83: Preparation of (R)-1-(3-((5-2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

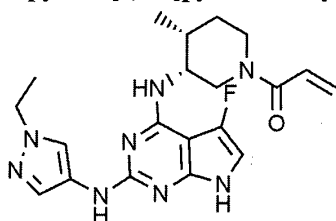


A title compound (2.0 mg, yield: 1.4%) was prepared in the same manner as in Example 82, except that 3-methylisothiazol-5-amine was used instead of 1-ethyl-1H-pyrazol-4-amine in Example 82.

¹H NMR (500MHz, CD₃OD) δ 7.75-7.6(m, 1H), 6.85-6.63(m, 2H), 6.30-6.05(m, 1H), 5.80-5.50(m, 1H), 4.57-4.45(m, 1H), 4.27-4.15(m, 1H), 3.90-3.86(m, 2H), 3.70-3.54(m, 1H), 2.32(s, 3H), 2.25-2.20(m, 1H), 2.16-2.00(m, 1H), 1.89-1.72(m, 2H), 1.65-1.50(m, 3H)

15

Example 84: Preparation of 1-((3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of 2,4-dichloro-5-fluoro-7H-pyrrolo [2,3-d]pyrimidine

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (200 mg, 1.06 mmol) was dissolved in acetonitrile (5.0 mL), 1-chloromethyl-4-fluoro-1,4-diazoniabicyclo[2.2.2]octane bis(tetrafluoroborate) (561.6 mg, 1.6 mmol) and acetic acid (1 mL) were added thereto. The mixture was heated at 80 °C and stirred for 24 hours, and then the organic layer was isolated,

treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (170.0 mg, yield: 80.1%).

^1H NMR (500MHz, CD_3OD) δ 7.36(s, 1H)

5

Step 2: Preparation of 2,4-dichloro-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (369.4 mg, 1.8 mmol) was dissolved in dichloromethane (30.0 mL), 4-dimethylaminopyridine (43.8 mg, 0.4 mmol) and triethylamine (499.8 mg, 3.6 mmol) were added thereto and then stirred for 30 minutes. 4-Methylbenzenesulfonyl chloride (393.1 mg, 2.1 mmol) was added to the reaction mixture, and the mixture was stirred at room temperature for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (363.0 mg, yield 56.2%).

10

^1H NMR (500MHz, CD_3OD) δ 8.07(d, 1H), 7.92(s, 1H), 7.45(d, 1H), 2.42(s, 3H)

15

Step 3: Preparation of *tert*-butyl (3R,4R)-3-((2-chloro-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carboxylate

After 2,4-dichloro-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidine (100.0 mg, 0.3 mmol) was dissolved in ethanol (3 mL), N,N-diisopropylethylamine (72.5 μL , 2.2 mmol) and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate (89.3 mg, 0.3 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (121.0 mg, yield: 81.7%).

20

^1H NMR (500MHz, CD_3OD) δ 8.01(d, 1H), 7.44-7.40(m, 3H), 4.59-4.50(m, 1H), 4.48-4.03(m, 3H), 2.98-2.82(m, 2H), 2.41(s, 3H), 2.10-2.00(m, 1H), 1.70-1.61(m, 1H), 1.58-1.36(m, 4H), 1.00-0.80(m, 9H)

25

Step 4: Preparation of *tert*-butyl (3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-

30

carboxylate

After *tert*-butyl (3R,4R)-3-((2-chloro-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carboxylate (100.0 mg, 0.2 mmol) was dissolved in anhydrous *tert*-butanol (2.0 mL), 1-ethyl-1H-pyrazol-4-amine (21.3 mg, 0.2 mmol),
5 *tris*(dibenzylideneacetone)dipalladium (8.5 mg, 0.01 mmol), 2-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl (8.9 mg, 0.2 mmol) and potassium carbonate (51.5 mg, 0.4 mmol) were added thereto, followed by reacting at 100 °C for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (58.3 mg, yield:
10 51.1%).

¹H NMR (500MHz, CD₃OD) δ 8.55(s, 1H), 7.94(d, 1H), 7.62(s, 1H), 7.30(d, 1H), 7.02(s, 1H), 4.60-4.55(m, 1H), 4.21-4.07(m, 4H), 2.95-2.78(m, 2H), 2.34(s, 3H), 2.06-2.00(m, 1H), 1.57-1.32(m, 8H), 0.98-0.86(m, 9H)

15 **Step 5: Preparation of *tert*-butyl (3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carboxylate**

Tert-butyl (3R,4R)-3-[(2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7-tosyl-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carboxylate (58.3 mg, 0.1 mmol) was dissolved in methanol (2.0 mL). Potassium hydroxide (10.7 mg, 0.2 mmol) was added to
20 the reaction mixture and then stirred at 50 °C for 5 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (28.6 mg, yield: 65.5%).

¹H NMR (500MHz, CD₃OD) δ 7.90(s, 1H), 7.55(s, 1H), 6.50(s, 1H), 4.59-4.50(m, 1H), 4.27-4.07(m, 3H), 3.16-2.85(m, 2H), 2.07-2.00(s, 1H), 1.63-1.61(m, 1H), 1.46-1.28(m,
25 7H), 1.15-1.00(m, 9H)

Step 6: Preparation of N2-(1-ethyl-1H-pyrazol-4-yl)-5-fluoro-N4-((3R,4R)-4-methylpiperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride

To *tert*-butyl (3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carboxylate (28.6 mg, 0.06 mmol)
30

was added 6 N hydrochloric acid solution (2.0 mL, excess) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

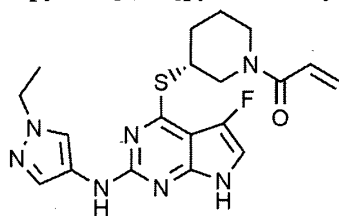
¹H NMR (500MHz, CD₃OD) δ 7.96-7.88(m, 1H), 7.69-7.63(m, 2H), 4.22-4.20(m, 2H), 4.10-4.05(m, 1H), 3.66-3.63(m, 1H), 3.16-3.11(m, 1H), 2.32-2.20(m, 1H), 1.98-1.94(m, 1H), 1.78-1.73(m, 1H) 1.48-1.46(m, 3H), 1.30-1.25(m, 1H), 1.15-1.05(m, 4H)

Step 7: Preparation of 1-((3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

After N2-(1-ethyl-1H-pyrazol-4-yl)-5-fluoro-N4-((3R,4R)-4-methylpiperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride (22.5 mg, 0.06 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4.0 ml), sodium bicarbonate (14.4 mg, 0.2 mmol) was added thereto at -20 °C and the mixture was stirred for 30 minutes. Acryloyl chloride (5.1 µL, 0.07 mmol) was slowly added dropwise to the reaction mixture and then stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (3.5 mg, yield: 14.9%).

¹H NMR (500MHz, CD₃OD) δ 8.01-7.80(m, 1H), 7.72-7.58(m, 2H), 6.86-6.35(m, 1H), 6.20-6.00(m, 1H), 5.85-5.45(m, 1H), 4.58-4.30(m, 2H), 4.22-4.14(m, 2H), 3.84-3.83(m, 2H), 2.95-2.78(m, 1H), 2.18-2.12(m, 1H), 1.70-1.65(m, 1H), 1.46-1.41(m, 3H), 1.31-1.28(m, 1H), 0.95-0.88(m, 3H)

Example 85: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



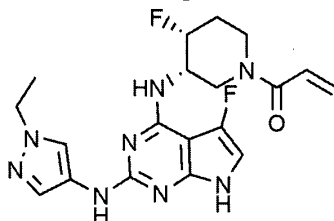
A title compound (4.1 mg, yield: 19.9%) was prepared in the same manner as in Example 84, except that *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate was used instead of

tert-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate in Example 84.

¹H NMR (500MHz, CD₃OD) δ 8.09-8.05(m, 1H), 7.85-7.50(m, 2H), 6.85-6.60(m, 1H), 6.28-6.11(m, 1H), 5.80-5.58(m, 1H), 4.60-4.00(m, 4H), 3.80-3.50(m, 2H), 3.48-3.38(m, 1H), 2.10-2.00(m, 1H), 1.90-1.75(m, 2H), 1.72-1.58(m, 1H), 1.50-1.40(m, 3H)

5

Example 86: Preparation of 1-((3S,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one

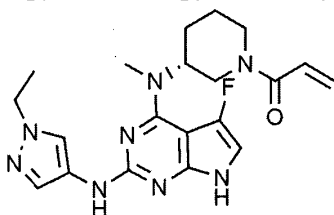


A title compound (7.9 mg, yield: 22.2%) was prepared in the same manner as in Example 84, except that *tert*-butyl (3R, 4R)-3-amino-4-fluoropiperidine-1-carboxylate was used instead of *tert*-butyl (3R, 4R)-3-amino-4-methylpiperidine-1-carboxylate in Example 84.

¹H NMR (500MHz, CD₃OD) δ 8.00-7.98(m, 1H), 7.57-7.41(m, 2H), 6.95-6.85(m, 1H), 6.27-6.15(m, 1H), 5.89-5.81(m, 1H), 5.07-4.97(m, 1H), 4.58-4.40(m, 1H), 4.22-4.06(m, 2H), 3.87-3.84(m, 1H), 3.44-3.34(m, 1H), 3.00-2.98(m, 1H), 2.20-2.15(m, 1H), 1.93-1.85(m, 1H), 1.46-1.33(m, 3H)

15

Example 87: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one

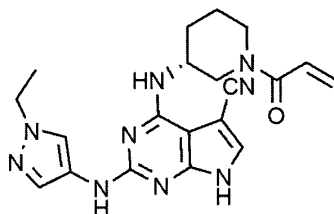


A title compound (7.9 mg, yield: 22.2%) was prepared in the same manner as in Example 84, except that *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate was used instead of *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate in Example 84.

¹H NMR (500MHz, CD₃OD) δ 7.95-7.61(m, 3H), 6.85-6.80(m, 1H), 6.25-6.05(m,

1H), 5.75-5.60(m, 1H), 4.68-4.58(m, 1H), 4.22-4.12(m, 3H), 3.85-3.75(m, 1H), 3.16-3.07(m, 3H), 2.70-2.55(m, 1H), 2.00-1.97(m, 1H), 1.69-1.65(m, 2H), 1.44-1.33(m, 3H)

**Example 88: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-((1-ethyl-
5 1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile**



**Step 1: Preparation of *tert*-butyl (R)-3-((2-chloro-5-cyano-7H-pyrrolo[2,3-
d]pyrimidin-4-yl)amino)piperidine-1-carboxylate**

After 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile (300.0 mg, 1.4 mmol)
10 was dissolved in ethanol (3.0 mL), N,N-diisopropylethylamine (369.8 μ L, 2.1 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (425.2 mg, 2.1 mmol) was added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (308 mg, yield:
15 57.7%).

^1H NMR (500MHz, CD_3OD) δ 7.83(m, 1H), 4.35-4.20(m, 1H), 3.73-3.59(m, 2H), 3.56-3.52(m, 1H), 2.13-2.02(m, 1H), 1.89-1.78(m, 2H), 1.70-1.60(m, 1H), 1.35-1.15(m, 9H)

**Step 2: Preparation of (R)-2-chloro-4-(piperidin-3-ylamino)-7H-pyrrolo[2,3-
20 d]pyrimidine-5-carbonitrile hydrochloride**

To *tert*-butyl (R)-3-((2-chloro-5-cyano-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (308.0 mg, 0.8 mmol) was added 6 N hydrochloric acid solution (5.0 mL, excess) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out
25 without isolation.

^1H NMR (500MHz, CD_3OD) δ 7.83(m, 1H), 4.35-4.20(m, 1H), 3.73-3.59(m, 2H), 3.56-3.52(m, 1H), 2.13-2.02(m, 1H), 1.89-1.78(m, 2H), 1.70-1.60(m, 1H)

Step 3: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-chloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile

After (R)-2-chloro-4-(piperidin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile hydrochloride (252.8 mg, 0.8 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4.0 ml), sodium bicarbonate (203.4 mg, 2.4 mmol) was added thereto at -20 °C and the mixture was stirred for 30 minutes. Acryloyl chloride (68.9 µL, 0.9 mmol) was slowly added dropwise to the reaction mixture and stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (130.0 mg, yield: 48.6%).

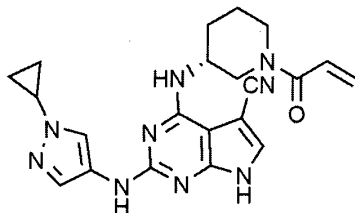
¹H NMR (500MHz, CD₃OD) δ 7.83(d, 1H), 6.84-6.75(m, 1H), 6.20-6.13(m, 1H), 5.74-5.64(m, 1H), 4.41-4.03(m, 2H), 3.94-3.65(m, 2H), 3.44-3.34(m, 1H), 2.15-2.10(m, 1H), 1.89-1.83(m, 2H), 1.80-1.75(m, 1H)

Step 4: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile

(R)-4-((1-acryloylpiperidin-3-yl)amino)-2-chloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile (30.0 mg, 0.09 mmol) and 1-ethyl-1H-pyrazol-4-amine (7.8 mg, 0.07 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (6.4 µL, 0.08 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 3 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to obtain a title compound (5.5 mg, yield: 19.6%).

¹H NMR (500MHz, CD₃OD) δ 7.95(d, 1H), 7.56-7.50(m, 2H), 6.90-6.52(m, 1H) 6.25-6.04(m, 1H), 5.78-5.50(m, 1H), 4.35-4.25(m, 1H), 4.14-4.11(m, 2H), 3.93-3.85(m, 1H), 3.80-3.65(m, 1H), 3.59-3.28(m, 2H), 2.23-2.13(m, 1H), 1.90-1.79(m, 2H), 1.68-1.66(m, 1H), 1.44-1.36(m, 3H)

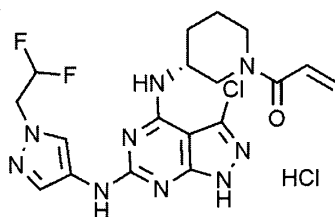
Example 89: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)amino)-2-((1-

cyclopropyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile

A title compound (2.8 mg, yield: 8.43%) was prepared in the same manner as in Example 88, except that 1-cyclopropyl-1H-pyrazol-4-amine was used instead of 1-ethyl-1H-pyrazol-4-amine in Example 88.

¹H NMR (500MHz, CD₃OD) δ 8.00(s, 1H), 7.53-7.51(m, 2H), 6.85-6.51(m, 1H), 6.24-6.04(m, 1H), 5.77-5.50(m, 1H), 4.41-3.95(m, 2H), 3.85-3.45(m, 2H), 2.15-2.10(m, 1H), 1.87-1.80(m, 3H), 1.70-1.67(m, 1H), 1.05-1.04(m, 4H)

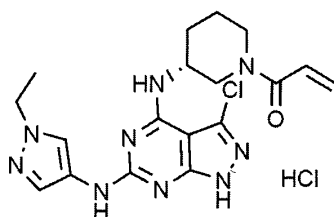
Example 90: Preparation of (R)-1-(3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one hydrochloride



After (R)-1-(3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (Example 44) (30.0 mg, 0.07 mmol) was dissolved in ethyl acetate, 2.0 N hydrochloric acid solution (3.0 mL, excess) dissolved in diethyl ether was added thereto. After stirring at room temperature for 1 hour, the reaction mixture was concentrated to obtain a title compound (28.0 mg, yield: 82.3%).

¹H NMR (500MHz, CD₃OD) δ 8.10-8.04(m, 1H), 7.68-7.63(m, 1H), 6.87-6.50(m, 1H), 6.28-6.15(m, 2H), 6.04-5.80(m, 1H), 4.58-4.53(m, 3H), 4.36-4.30(m, 1H), 4.05-3.90(m, 2H), 3.44-3.40(m, 1H), 2.15-2.13(m, 1H), 1.98-1.86(m, 2H), 1.66-1.64(m, 1H)

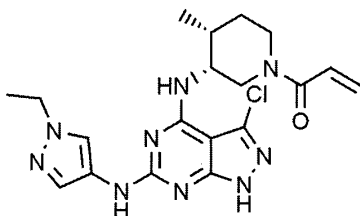
Example 91: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

hydrochloride

A title compound (2.8 mg, yield: 8.43%) was prepared in the same manner as in Example 90, except that (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (Example 46) was used instead of (R)-1-(3-(3-chloro-6-(1-(2,2-difluoroethyl)-1H-pyrazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-ylamino)piperidin-1-yl)prop-2-en-1-one in Example 90.

¹H NMR (500MHz, CD₃OD) δ 8.07(s, 1H), 7.96(s, 1H), 6.86-6.81(m, 1H), 6.27-6.15(m, 1H) 5.81-5.58(m, 1H) 4.37-4.30(m, 2H), 4.11-4.06(m, 2H), 3.92-3.82(m, 1H), 3.43-3.39(m, 2H), 2.15-2.10(m, 1H), 1.90-1.80(m, 2H), 1.70-1.63(m, 1H), 1.46-1.40(m, 3H)

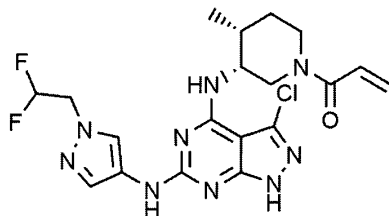
Example 92: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (5.7 mg, yield: 26.6%) was prepared in the same manner as in Example 42, except using 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.96(s, 1H), 7.59(s, 1H), 6.90-6.46(m, 1H), 6.20-6.01(m, 1H), 5.77-5.42(m, 1H), 4.68-4.58(m, 2H), 4.46-4.39(m, 2H), 4.16-4.13(m, 2H), 3.16-3.12(m, 1H), 2.89-2.85(m, 1H), 2.25-2.15(m, 1H), 1.76-1.74(m, 1H), 1.46-1.44(m, 3H), 1.05(d, 3H)

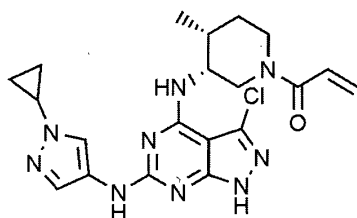
Example 93: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



5 A title compound (31.5 mg, yield: 66.4%) was prepared in the same manner as in Example 42, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.05(s, 1H), 7.65(s, 1H), 6.90-6.45(m, 1H), 6.30-6.01(m, 2H), 5.77-5.42(m, 1H), 4.67-4.39d(m, 5H), 3.16-2.84(m, 1H), 2.25-2.15(m, 1H), 1.76-1.73(m, 1H), 1.49-1.41(m, 1H), 1.30-1.25(m, 1H), 1.04(d, 3H)

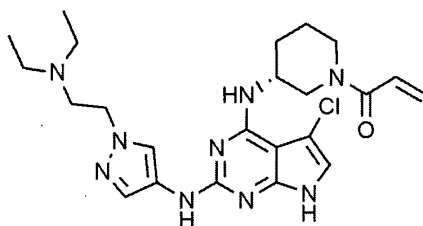
Example 94: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



15 A title compound (30.0 mg, yield: 66.1%) was prepared in the same manner as in Example 42, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and cyclopropyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.99(s, 1H), 7.57(s, 1H), 6.89-6.43(m, 1H), 6.20-6.01(m, 1H), 5.77-5.42(m, 1H), 4.69-4.10(m, 3H), 3.60-3.58(m, 1H), 3.39-3.37(m, 1H), 3.15-2.85(m, 1H), 2.23-2.19(m, 1H), 1.76-1.74(m, 1H), 1.46-1.41(m, 1H), 1.08-1.00(m, 7H)

Example 95: Preparation of (R)-1-(3-((5-chloro-2-((1-(2-(diethylamino)ethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



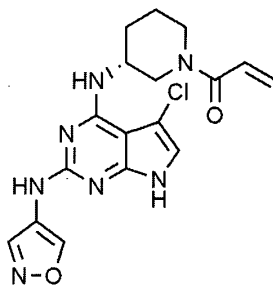
5

A title compound (4.5 mg, yield: 18.6%) was prepared in the same manner as in Example 36, except that 1-(2-(diethylamino)ethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500 MHz, CD₃OD) δ 7.98-7.92(m, 1H), 7.58-7.56(m, 1H), 6.82-6.55(m, 2H), 6.25-6.06(m, 1H), 5.78-5.50(m, 1H), 4.34-3.40(m, 7H), 3.00-2.97(m, 2H), 2.68-2.61(m, 4H), 2.08-2.02(m, 2H), 1.90-1.87(m, 1H), 1.67-1.59(m, 1H), 1.08-1.02(m, 6H)

10

Example 96: Preparation of (R)-1-(3-((5-chloro-2-(isoxazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



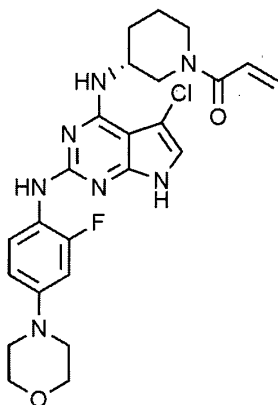
15

A title compound (8.3 mg, yield: 42.8%) was prepared in the same manner as in Example 36, except that isoxazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500 MHz, CD₃OD) δ 9.03(s, 1H), 8.48(s, 1H), 6.90-6.53(m, 2H), 6.25-6.08(m, 1H), 5.78-5.50(m, 1H), 4.40-4.20(m, 2H), 3.90-3.70(m, 2H), 3.60-3.40(m, 1H), 2.13-2.00(m, 2H), 1.92-1.73(m, 1H), 1.72-1.55(m, 1H)

20

Example 97: Preparation of (R)-1-(3-((5-chloro-2-(isoxazol-4-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

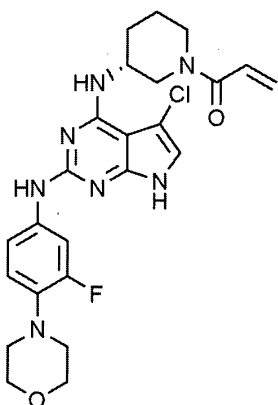


A title compound (11.4 mg, yield: 45.8%) was prepared in the same manner as in Example 36, except that 2-fluoro-4-morpholinoaniline was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 8.00-7.90(m, 1H), 7.08-6.51(m, 4H), 6.25-6.05(m, 1H), 5.80-5.45(m, 1H), 4.35-4.10(m, 2H), 3.85-3.70(m, 4H), 3.65-3.35(m, 3H), 3.25-3.00(m, 4H), 2.15-1.95(m, 1H), 1.90-1.70(m, 1H), 1.63-1.54(m, 2H)

10

Example 98: Preparation of (R)-1-(3-((5-chloro-2-((3-fluoro-4-morpholinophenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

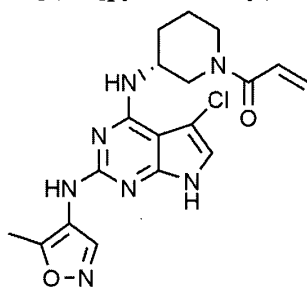


A title compound (6.9 mg, yield: 27.8%) was prepared in the same manner as in Example 36, except that 3-fluoro-4-morpholinoaniline was used instead of 4-(4-

methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.84-7.72(m, 1H), 7.27-7.18(m, 1H), 6.98-6.90(m, 1H), 6.87-6.50(m, 2H), 6.28-6.00(m, 1H), 5.80-5.43(m, 1H), 4.48-3.90(m, 2H), 3.85-3.77(m, 4H), 3.76-3.58(m, 2H), 3.52-3.33(m, 1H), 3.02-2.95(m, 4H), 2.15-2.00(m, 1H), 1.90-1.75(m, 2H), 1.70-1.55(m, 1H)

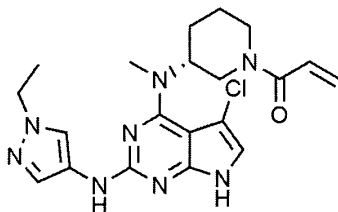
Example 99: Preparation of (R)-1-(3-((5-chloro-2-((5-methylisoxazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.6 mg, yield: 37.7%) was prepared in the same manner as in Example 36, except that 5-methylisoxazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 8.75-8.65(m, 1H), 6.90-6.50(m, 2H), 6.25-6.04(m, 1H), 5.80-5.52(m, 1H), 4.35-4.20(m, 1H), 3.90-3.60(m, 2H), 3.55-3.35(m, 2H), 2.39 (s, 3H), 2.16-2.05(m, 1H), 1.90-1.75(m, 2H), 1.70-1.60(m, 1H)

Example 100: Preparation of (R)-1-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one

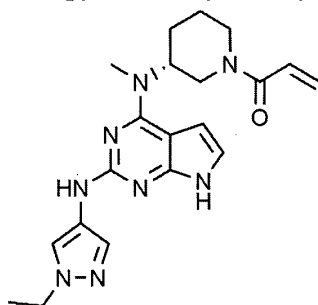


A title compound (9.1 mg, yield: 42.6%) was prepared in the same manner as in Example 36, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of

tert-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.90-7.75(m, 1H), 7.48(s, 1H), 6.90-6.60(m, 1H), 6.40-6.10(m, 2H), 5.80-5.60(m, 1H), 4.70-4.50(m, 2H), 4.20-4.00(m, 3H), 3.70-3.60(m, 1H),
 5 3.30-3.20(m, 3H), 2.20-2.10(m, 3H), 2.10-1.90(m, 2H), 1.65-1.50(m, 2H), 1.45-1.35(m, 1H)

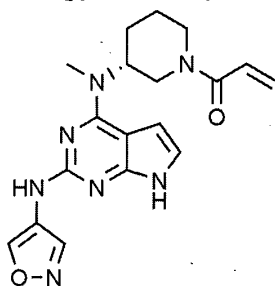
Example 101: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one



10 A title compound (8.2 mg, yield: 41.4%) was prepared in the same manner as in Example 1, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CDCl₃) δ 9.09(s, 1H), 7.77(s, 1H), 7.46(s, 1H), 6.66-6.61(m, 2H),
 15 6.34-6.31(m, 3H), 5.67-5.65(m, 1H), 4.76-4.75(m, 2H), 4.15-4.07(m, 3H), 3.28(s, 3H), 3.07-2.98(m, 1H), 2.81-2.50(m, 1H), 2.10-1.87(m, 5H), 1.69-1.63(m, 1H)

Example 102: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one



20

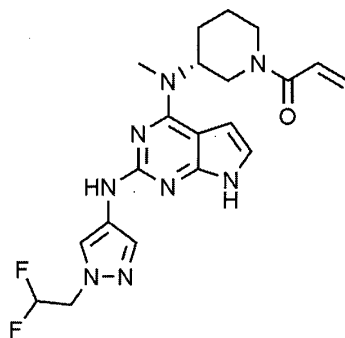
A title compound (8.7 mg, yield: 47.3%) was prepared in the same manner as in

Example 1, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and isoxazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 8.94(s, 1H), 8.47(s, 1H), 6.83-6.74(m, 2H), 6.52-6.51(m, 1H), 6.27-6.17(m, 1H), 5.79-5.66(m, 1H), 4.60(s, 1H), 4.14(s, 1H), 3.67 (s, 3H), 2.03-2.02(m, 3H), 1.61-1.60(m, 4H)

Example 103: Preparation of (R)-1-(3-((2-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-

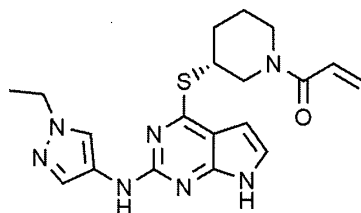
10 one



A title compound (8.7 mg, yield: 40.6%) was prepared in the same manner as in Example 1, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.97-7.96(m, 1H), 7.57-7.56(m, 1H), 6.85-6.79(m, 2H), 6.49-6.48(m, 1H), 6.27-6.10(m, 2H), 5.80-5.62(m, 1H), 4.73-4.60(m, 1H), 4.45-4.44(m, 2H), 4.17-4.14(m, 1H), 3.30(s, 3H), 3.11-2.93(m, 2H), 2.03-2.02(m, 3H), 1.70-1.60(m, 2H)

20 **Example 104: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one**

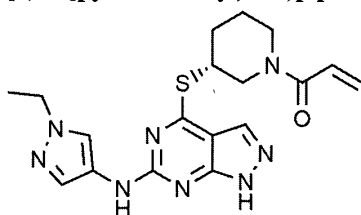


A title compound (5.9 mg, yield: 29.9%) was prepared in the same manner as in Example 1, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.59-7.58(m, 1H), 6.92-6.90(m, 1H), 6.85-6.45(m, 1H), 6.28-6.02(m, 2H), 5.80-5.43(m, 1H), 4.66-4.25(m, 1H), 4.20-4.00(m, 3H), 3.90-3.45(m, 2H), 3.40-3.20(m, 1H), 2.25-2.15(m, 1H), 1.90-1.75(m, 2H), 1.70-1.60(m, 1H), 1.50-1.40(m, 3H)

10

Example 105: Preparation of (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

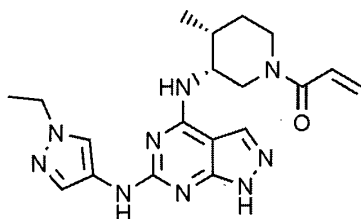


A title compound (8.4 mg, yield: 42.2%) was prepared in the same manner as in Example 30, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.20-8.10(m, 1H), 7.83(s, 1H), 7.64(s, 1H), 6.90-6.50(m, 1H), 6.30-6.10(m, 1H), 5.80-5.50(m, 1H), 4.60-6.35(m, 1H), 4.30-4.00(m, 3H), 3.90-3.35(m, 3H), 2.28-2.20(m, 1H), 1.97-1.80(m, 2H), 1.77-1.65(m, 1H), 1.50-1.40(m, 3H)

20

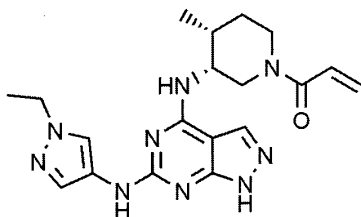
Example 106: Preparation of 1-((3R,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (8.2 mg, yield: 41.6%) was prepared in the same manner as in Example 30, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.05-7.90(m, 2H), 7.60-7.55(m, 1H), 6.88-6.33(m, 1H), 6.20-5.90(m, 1H), 5.75-5.30(m, 1H), 4.70-4.35(m, 3H), 4.20-4.10(m, 2H), 4.04-2.80(m, 1H), 2.25-2.10(m, 1H), 1.90-1.75(m, 1H), 1.70-1.40(m, 5H), 1.10-1.00(m, 3H)

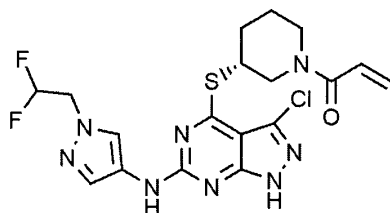
Example 107: Preparation of 1-((3R,4R)-3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (5.7 mg, yield: 28.8%) was prepared in the same manner as in Example 1, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 1.

¹H NMR (500MHz, CD₃OD) δ 7.89(s, 1H), 7.56(s, 1H), 6.88-6.30(m, 3H), 6.30-5.30(m, 2H), 4.60-4.30(m, 3H), 4.20-4.10(m, 2H), 3.95-3.35(m, 1H), 2.98-2.13(m, 2H), 1.88-1.55(m, 2H), 1.50-1.40(m, 3H), 1.10-1.00(m, 3H)

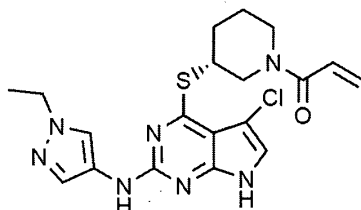
Example 108: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



A title compound (9.3 mg, yield: 39.7%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.20-8.05(m, 1H), 7.75-7.68(m, 1H), 6.88-6.45(m, 1H), 6.30-6.00(m, 2H), 5.80-5.50(m, 1H), 4.65-4.30(m, 3H), 4.28-4.00(m, 1H), 3.92-3.60(m, 2H), 3.50-3.31(m, 1H), 2.25-2.18(m, 1H), 1.95-1.80(m, 2H), 1.73-1.65(m, 1H)

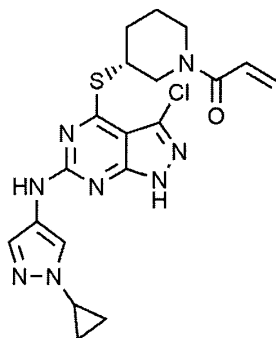
Example 109: Preparation of (R)-1-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



A title compound (8.2 mg, yield: 37.8%) was prepared in the same manner as in Example 36, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 36.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.60-7.58(m, 1H), 6.90-6.88(m, 1H), 6.87-6.50(m, 1H), 6.26-6.08(m, 1H), 5.80-5.50(m, 1H), 4.70-4.30(m, 1H), 4.25-4.15(m, 3H), 3.92-3.80(m, 1H), 3.70-3.15(m, 2H), 2.25-2.18(m, 1H), 1.95-1.75(m, 2H), 1.73-1.65(m, 1H), 1.50-1.40(m, 3H)

Example 110: Preparation of (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

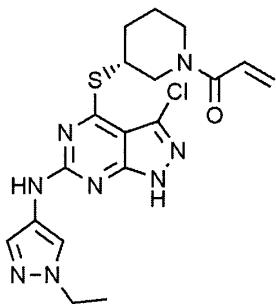


A title compound (4.4 mg, yield: 19.7%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-cyclopropyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.13-8.00(m, 1H), 7.65-7.55(m, 1H), 6.88-6.45(m, 1H), 6.30-6.10(m, 1H), 5.82-5.50(m, 1H), 4.68-4.50(m, 1H), 4.40-4.20(m, 1H), 4.10-3.97(m, 1H), 3.92-3.80(m, 1H), 3.48-3.31(m, 1H), 2.28-2.15(m, 1H), 2.08-2.00(m, 1H), 1.95-1.80(m, 2H), 1.75-1.60(m, 1H), 1.13-1.00(m, 4H)

10

Example 111: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

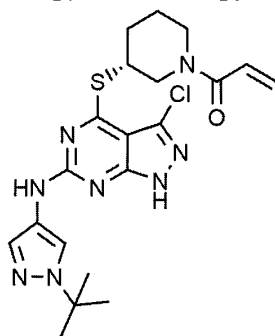


A title compound (7.5 mg, yield: 34.5%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.10-7.93(m, 1H), 7.70-7.60(m, 1H), 6.88-6.50(m, 1H), 6.30-6.10(m, 1H), 5.80-5.50(m, 1H), 4.65-3.30(m, 6H), 2.28-2.20(m, 1H), 2.08-2.00(m,

1H), 1.95-1.80(m, 2H), 1.78-1.62(m, 1H), 1.50-1.40(m, 3H)

Example 112: Preparation of (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



5

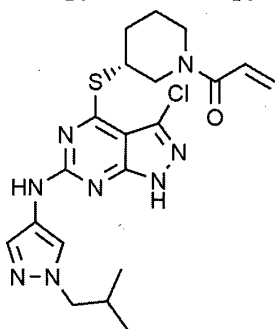
A title compound (10.3 mg, yield: 44.7%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(*tert*-butyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.20-8.05(m, 1H), 7.72-7.65(m, 1H), 6.88-6.50(m, 1H), 6.30-6.10(m, 1H), 5.80-5.50(m, 1H), 4.70-3.30(m, 4H), 2.28-2.20(m, 1H), 2.08-1.98(m, 1H), 1.97-1.80(m, 2H), 1.78-1.66(m, 1H), 1.59(s, 9H)

10

Example 113: Preparation of (R)-1-(3-((3-chloro-6-((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

15



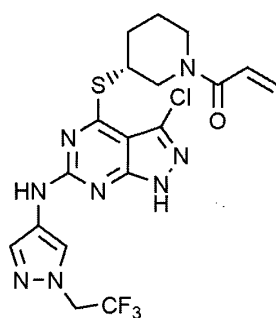
A title compound (4.7 mg, yield: 20.4%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-isobutyl-1H-pyrazol-4-amine instead of 4-(4-

methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.10-7.95(m, 1H), 7.72-7.65(m, 1H), 6.88-6.50(m, 1H), 6.30-6.10(m, 1H), 5.80-5.50(m, 1H), 4.70-3.35(m, 6H), 2.28-2.10(m, 2H), 2.08-1.98(m, 1H), 1.97-1.80(m, 2H), 1.78-1.66(m, 1H), 0.92-0.85(m, 6H)

5

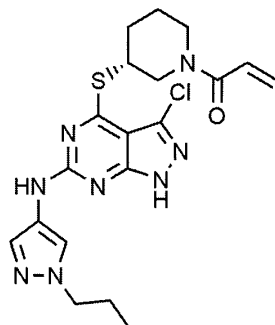
Example 114: Preparation of (R)-1-(3-((3-chloro-6-((1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



10 A title compound (10.1 mg, yield: 41.5%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

15 ¹H NMR (500MHz, CD₃OD) δ 8.22-8.10(m, 1H), 7.80-7.70(m, 1H), 6.88-6.50(m, 1H), 6.30-6.10(m, 1H), 5.82-5.50(m, 1H), 4.90-4.85(m, 2H), 4.45-3.30(m, 4H), 2.28-2.15(m, 1H), 2.08-1.99(m, 1H), 1.97-1.80(m, 2H), 1.78-1.65(m, 1H)

Example 115: Preparation of (R)-1-(3-((3-chloro-6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

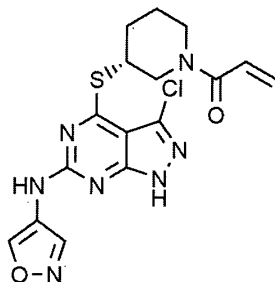


A title compound (9.8 mg, yield: 43.8%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-propyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.10-7.95(m, 1H), 7.70-7.60(m, 1H), 6.90-6.50(m, 1H), 6.30-6.05(m, 1H), 5.80-5.50(m, 1H), 4.42-4.20(m, 1H), 4.10-4.00(m, 2H), 3.95-3.60(m, 2H), 3.50-3.30(m, 1H), 2.28-2.20(m, 1H), 2.08-1.80(m, 4H), 1.77-1.50(m, 2H), 0.96-0.81(m, 3H)

10

Example 116: Preparation of (R)-1-(3-((3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one

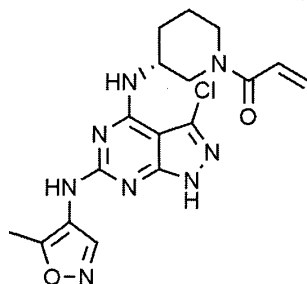


A title compound (9.8 mg, yield: 48.1%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and isoxazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 9.20-9.10(m, 1H), 8.56(s, 1H), 6.90-6.53(m, 1H), 6.30-6.10(m, 1H), 5.80-5.50(m, 1H), 4.70-3.62(m, 5H), 2.00-1.80(m, 1H), 1.79-1.65(m, 2H), 1.63-1.55(m, 1H)

20

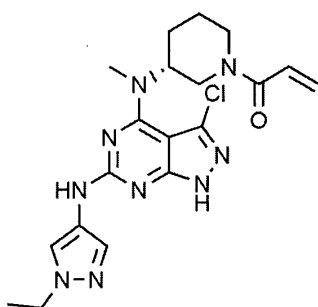
Example 117: Preparation of (R)-1-(3-((3-chloro-6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidin-1-yl)prop-2-en-1-one



5 A title compound (4.2 mg, yield: 20.8%) was prepared in the same manner as in Example 42, except that 5-methylisoxazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.70-8.66(m, 1H), 6.84-6.61(m, 1H), 6.26-6.11(m, 1H), 5.79-5.59(m, 1H), 4.32-4.28(m, 1H), 3.96-3.84(m, 2H), 3.63-3.43(m, 2H), 2.40(s, 3H),
10 2.10-2.04(m, 1H), 1.92-1.83(m, 2H), 1.65-1.63(m, 1H)

Example 118: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-one



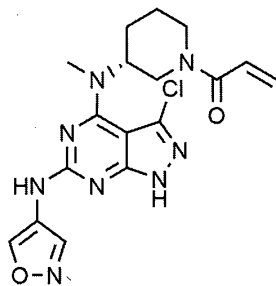
15 A title compound (9.5 mg, yield: 44.3%) was prepared in the same manner as in Example 42, except that *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate was used instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

20 ¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.57-7.55(m, 1H), 6.83-6.67(m, 1H), 6.24-6.13(m, 1H), 5.77-5.64(m, 1H), 4.74-4.52(m, 2H), 4.11-4.09(m, 3H), 3.24(s, 3H), 3.10-

2.62(m, 2H), 2.00-1.92(m, 3H), 1.63-1.62(m, 1H), 1.43(t, 3H)

Example 119: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidin-1-yl)prop-2-en-1-

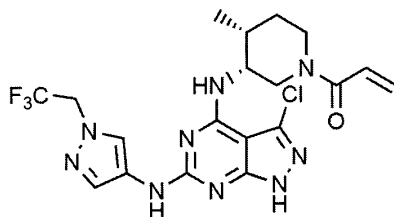
5 one



A title compound (8.7 mg, yield: 43.3%) was prepared in the same manner as in Example 42, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and isoxazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 9.03(s, 1H), 8.51(s, 1H), 6.84-6.72(m, 1H), 6.24-6.16(m, 1H), 5.78-5.67(m, 1H), 4.82-4.56(m, 2H), 4.21-4.13(m, 1H), 3.31(s, 3H), 3.11-2.99(m, 1H), 2.05-1.94(m, 3H), 1.61-1.60(m, 2H)

Example 120: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

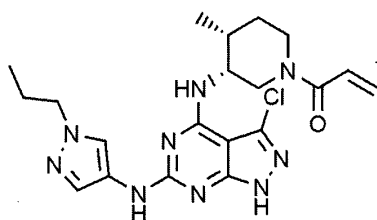


A title compound (8.1 mg, yield: 33.4%) was prepared in the same manner as in Example 42, except using 1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.11(s, 1H), 7.67(s, 1H), 6.86-6.42(m, 1H), 6.20-6.02(m, 1H), 5.77-5.42(m, 1H), 4.66-4.38(m, 2H), 4.21-4.11(m, 1H), 3.39-3.35(m, 1H), 3.12-2.85(m, 1H), 2.19-2.14(m, 2H), 2.08-2.00(m, 1H), 1.76-1.74(m, 1H), 1.61-1.56(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H)

5

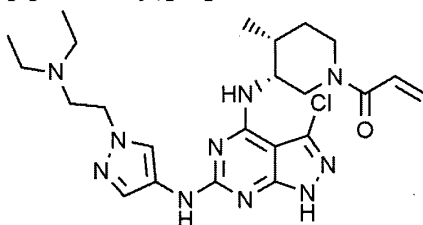
Example 121: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-propyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



10 A title compound (7.1 mg, yield: 32.1%) was prepared in the same manner as in Example 42, except using 1-propyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.95(s, 1H), 7.57(s, 1H), 6.86-6.42(m, 1H), 6.19-6.01(m, 1H), 5.77-5.42(m, 1H), 4.68-4.12(m, 3H), 4.08-4.05(m, 2H), 3.40-3.36(m, 1H), 3.12-2.84(m, 1H), 2.20-2.17(m, 1H), 1.87-1.83(m, 2H), 1.76-1.74(m, 1H), 1.61-1.56(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H), 0.95-0.89(m, 3H)

Example 122: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-(2-(diethylamino)ethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

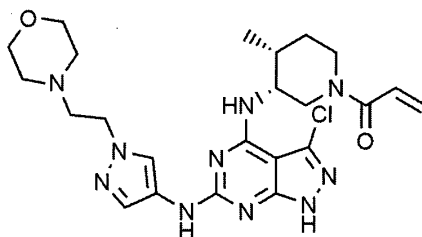


A title compound (6.4 mg, yield: 25.7%) was prepared in the same manner as in

Example 42, except using 1-(2-(diethylamino)ethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.02(s, 1H), 7.64(s, 1H), 6.87-6.43(m, 1H), 6.20-6.02(m, 1H), 5.77-5.42(m, 1H), 4.71-4.20(m, 5H), 3.39-3.35(m, 1H), 3.15-3.00(m, 3H), 2.90-2.85(m, 1H), 2.78-2.63(m, 4H), 2.20-2.17(m, 1H), 1.78-1.75(m, 1H), 1.61-1.58(m, 1H), 1.45-1.41(m, 1H), 1.13-1.10(m, 4H), 1.05-1.04(d, *J* = 5 Hz, 3H)

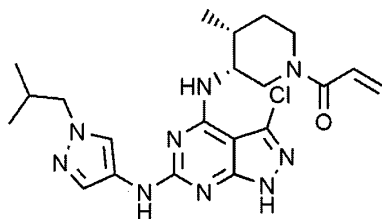
Example 123: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-(2-morpholinoethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (12.4 mg, yield: 48.2%) was prepared in the same manner as in Example 42, except using 1-(2-morpholinoethyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.00(s, 1H), 7.63(s, 1H), 6.87-6.44(m, 1H), 6.21-6.02(m, 1H), 5.76-5.42(m, 1H), 4.70-4.24(m, 5H), 3.69-3.63(m, 4H), 3.40-3.38(m, 1H), 3.12-2.85(m, 1H), 2.81-2.75(m, 2H), 2.50-2.45(m, 4H), 2.19-2.17(m, 1H), 2.03-2.02(m, 1H), 1.76-1.74(m, 1H), 1.44-1.40(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H)

Example 124: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-isobutyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

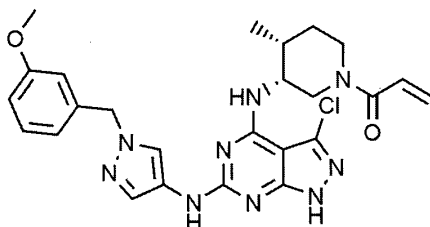


A title compound (8.3 mg, yield: 36.2%) was prepared in the same manner as in Example 42, except using 1-isobutyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.94(s, 1H), 7.60(s, 1H), 6.87-6.43(m, 1H), 6.20-6.02(m, 1H), 5.78-5.42(m, 1H), 4.69-4.11(m, 3H), 3.90-3.89(d, *J* = 5 Hz, 2H), 3.40-3.35(m, 1H), 3.14-2.83(m, 1H), 2.18-2.11(m, 2H), 1.75-1.73(m, 1H), 1.48-1.41(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H), 0.94-0.93(d, *J* = 5 Hz, 6H)

10

Example 125: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-(3-methoxybenzyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

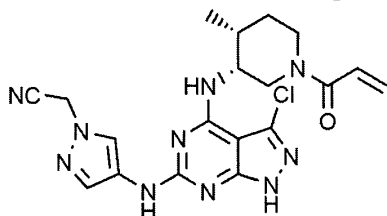


A title compound (9.8 mg, yield: 37.7%) was prepared in the same manner as in Example 42, except using 1-(3-methoxybenzyl)-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.99(s, 1H), 7.60(s, 1H), 7.28-7.25(m, 1H), 6.88-6.80(m, 3H), 6.87-6.43(m, 1H), 6.19-6.00(m, 1H), 5.75-5.38(m, 1H), 5.27(s, 2H), 4.71-4.11(m, 3H), 3.76(s, 3H), 3.32-3.30(m, 1H), 3.14-2.83(m, 1H), 2.18-2.06(m, 1H), 1.75-1.72(m, 1H), 1.60-1.58(m, 1H), 1.02-1.01(d, *J* = 5 Hz, 3H)

20

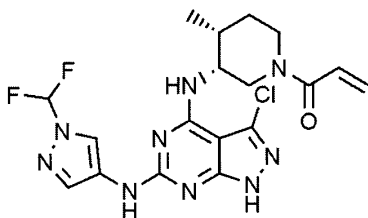
Example 126: Preparation of 2-(4-(((3R,4R)-1-acryloyl-4-methylpiperidin-3-yl)amino)-3-chloro-1H-pyrazolo[3,4-d]pyrimidin-6-yl)amino)-1H-pyrazol-1-yl)acetonitrile



A title compound (4.1 mg, yield: 18.7%) was prepared in the same manner as in Example 42, except using 2-(4-amino-1H-pyrazol-1-yl) acetonitrile instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.12(s, 1H), 7.69(s, 1H), 6.87-6.42(m, 1H), 6.18-6.01(m, 1H), 5.77-5.41(m, 1H), 5.28(s, 2H), 4.76-4.10(m, 3H), 3.41-3.37(m, 1H), 3.15-2.84(m, 1H), 2.20-2.19(m, 1H), 1.76-1.74(m, 1H), 1.50-1.41(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H)

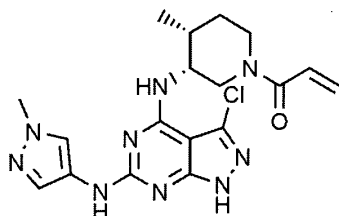
Example 127: Preparation of 1-(((3R,4R)-3-((3-chloro-6-((1-(difluoromethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (10.2 mg, yield: 45.1%) was prepared in the same manner as in Example 42, except using 1-difluoromethyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline, and *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 42.

¹H NMR (500MHz, CD₃OD) δ 8.40(s, 1H), 7.79(s, 1H), 7.40(t, *J* = 60 Hz, 1H), 6.87-6.41(m, 1H), 6.19-6.00(m, 1H), 5.76-5.41(m, 1H), 4.75-4.11(m, 3H), 3.41-3.37(m, 1H), 3.13-2.82(m, 1H), 2.20-2.19(m, 1H), 1.76-1.74(m, 1H), 1.47-1.40(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H)

Example 128: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-methyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

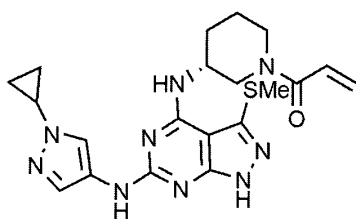


5

A title compound (8.0 mg, yield: 38.5%) was prepared in the same manner as in Example 42, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-methyl-1H-pyrazol-4-amine instead of 4-(4-methylpiperazin-1-yl)aniline in Example 42.

¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.58(s, 1H), 6.87-6.41(m, 1H), 6.19-6.00(m, 1H), 5.77-5.40(m, 1H), 4.69-4.38(m, 3H), 3.85(s, 3H), 3.40-3.37(m, 1H), 3.12-2.83(m, 1H), 2.20-2.17(m, 1H), 2.03-2.02(m, 1H), 1.75-1.73(m, 1H), 1.50-1.40(m, 1H), 1.05-1.04(d, *J* = 5 Hz, 3H)

Example 129: Preparation of (R)-1-3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



Step 1: Preparation of *tert*-butyl (R)-3-((6-chloro-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino) piperidine-1-carboxylate

20

After 4,6-dichloro-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidine (300.0 mg, 1.3 mmol) was dissolved in ethanol (3 mL), *N,N*-diisopropylethylamine (333.4 μL, 1.9 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (383.4 mg, 1.9 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hour. The organic layer was isolated, treated with

magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (403.2 mg, yield: 79.4%).

¹H NMR (500MHz, CD₃OD) δ 4.77-4.59(m, 1H), 4.30-4.25(m, 1H), 3.86-3.65(m, 2H), 3.54-3.51(m, 1H), 2.59(s, 3H), 2.05-1.92(m, 2H), 1.75-1.66(m, 2H), 1.43-1.22(m, 9H)

5

Step 2: Preparation of (R)-6-chloro-3-(methylthio)-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine hydrochloride

To *tert*-butyl (R)-3-((6-chloro-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino) piperidine-1-carboxylate (403.2 mg, 1.0 mmol) was added 6 N hydrochloric acid solution (2.0 mL, excessive amount) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 4.60-4.57(m, 1H), 3.64-3.61(m, 1H), 3.44-3.34(m, 1H), 3.16-2.97(m, 1H), 2.62(s, 3H), 2.20-2.07(m, 2H), 1.98-1.80(m, 2H)

15

Step 3: Preparation of (R)-1-(3-((6-chloro-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-6-chloro-3-(methylthio)-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4-amine hydrochloride (339.6 mg, 1.0 mmol) was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4 ml), sodium bicarbonate (255.3 mg, 1.1 mmol) was added thereto at -20 °C and the mixture was stirred for 30 minutes. Acryloyl chloride (86.4 μL, 1.1 mmol) was slowly added dropwise to the reaction mixture and then stirred at -20 °C for 1 hour. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (207.0 mg, yield 58.1%).

25

¹H NMR (500MHz, CD₃OD) δ 6.83-6.72(m, 1H), 6.22-6.12(m, 1H), 5.78-5.62(m, 1H), 4.79-4.26(m, 1H), 4.10-4.02(m, 1H), 3.80-3.60(m, 3H), 2.57(d, 3H), 2.11-1.93(m, 2H), 1.88-1.72(m, 2H)

30

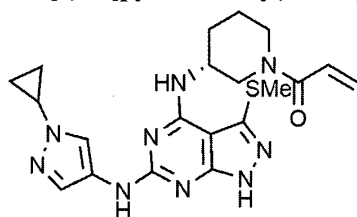
Step 4: Preparation of (R)-1-(3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-

(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

(R)-1-(3-((6-chloro-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (30.0 mg, 0.09 mmol) and 1-cyclopropyl-1H-pyrazol-4-amine (9.6 mg, 0.08 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (5.2 μ L, 0.07 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 3 hours, and then the solvent was concentrated. This reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol. The residue was isolated by column chromatography to obtain a title compound (16.9 mg, yield: 58.9%).

¹H NMR (500MHz, CD₃OD) δ 8.00(s, 1H), 7.54(s, 1H), 6.86-6.52(m, 1H), 6.25-6.05(m, 1H), 5.78-5.50(m, 1H), 4.36-4.30(m, 1H), 4.20-4.11(m, 1H), 3.90-3.74(m, 2H), 3.65-3.58(m, 2H), 2.52(s, 3H), 2.08-2.05(m, 1H), 1.93-1.85(m, 2H), 1.70-1.65(m, 1H), 1.07-1.00(m, 4H)

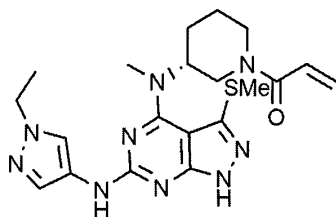
Example 130: Preparation of (R)-1-(3-((6-(isoxazol-4-ylamino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (9.8 mg, yield: 48.9%) was prepared in the same manner as in Example 129, except that isoxazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 9.10-9.00(m, 1H), 8.52-8.45(m, 1H), 6.90-6.50(m, 1H), 6.30-6.02(m, 1H), 5.80-5.50(m, 1H), 4.40-4.25(m, 1H), 4.20-3.40(m, 4H), 2.51(s, 3H), 2.10-1.98(m, 1H), 1.97-1.60(m, 3H)

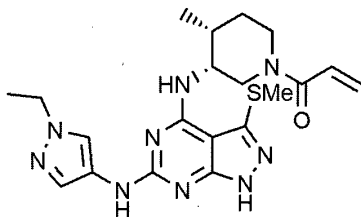
Example 131: Preparation of (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methylamino)piperidin-1-yl)prop-2-en-1-one



A title compound (6.4 mg, yield: 28.9%) was prepared in the same manner as in Example 129, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 7.98-7.90(m, 1H), 7.60-7.50(m, 1H), 6.85-6.57(m, 1H), 6.22-6.05(m, 1H), 5.77-5.55(m, 1H), 4.70-4.40(m, 2H), 4.20-4.05(m, 3H), 3.40-2.60(m, 5H), 2.54(s, 3H), 2.10-1.85(m, 3H), 1.70-1.50(m, 1H), 1.45-1.35(m, 3H)

Example 132: Preparation of 1-((3R,4R)-3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

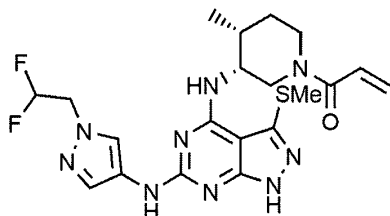


A title compound (10.1 mg, yield: 45.8%) was prepared in the same manner as in Example 129, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-ethyl-1H-pyrazol-4-amine instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 8.15-7.90(m, 1H), 7.62-7.55(m, 1H), 6.90-6.38(m, 1H), 6.20-5.90(m, 1H), 5.80-5.30(m, 1H), 4.80-4.35(m, 3H), 4.20-4.07(m, 2H), 3.40-3.20(m, 1H), 3.10-2.70(m, 1H), 2.54-2.40(m, 3H), 2.20-2.10(m, 1H), 1.80-1.70(m, 1H), 1.50-1.38(m, 4H), 1.10-0.98(m, 3H)

Example 133: Preparation of 1-((3R,4R)-3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-

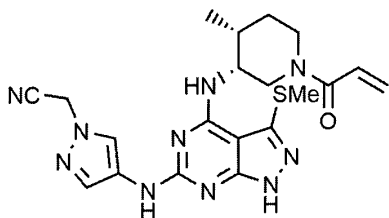
4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



A title compound (5.2 mg, yield: 21.6%) was prepared in the same manner as in
 5 Example 129, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 8.10-8.00(m, 1H), 7.70-7.62(m, 1H), 6.90-6.35(m,
 1H), 6.30-5.90(m, 2H), 5.80-5.30(m, 1H), 4.75-4.35(m, 5H), 4.20-3.30(m, 1H), 3.10-2.80(m,
 10 1H), 2.50-2.40(m, 3H), 2.20-2.10(m, 1H), 1.80-1.40(m, 2H), 1.10-1.00(m, 3H)

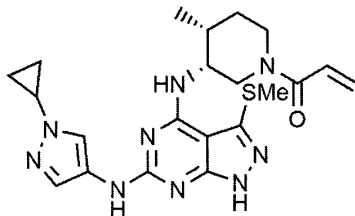
Example 134: Preparation of 2-(4-(((3R,4R)-1-acryloyl-4-methylpiperidin-3-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-6-yl)amino)-1H-pyrazol-1-yl)acetonitrile



15 A title compound (10.3 mg, yield: 45.5%) was prepared in the same manner as in Example 129, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate, and 2-(4-amino-1H-pyrazol-1-yl)acetonitrile instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

20 ¹H NMR (500MHz, CD₃OD) δ 8.20-8.10(m, 1H), 7.72-7.62(m, 1H), 6.90-6.35(m, 1H), 6.20-5.90(m, 1H), 5.80-5.35(m, 1H), 5.28(s, 2H), 4.80-4.35(m, 3H), 4.20-3.30(m, 1H), 3.13-2.75(m, 1H), 2.50-2.40(m, 3H), 2.25-2.18(m, 1H), 1.80-1.40(m, 2H), 1.20-1.10(m, 3H)

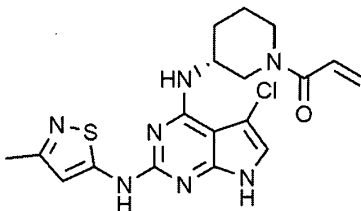
Example 135: Preparation of 1-((3R,4R)-3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



5 A title compound (5.2 mg, yield: 21.6%) was prepared in the same manner as in Example 129, except that *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate was used instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 129.

¹H NMR (500MHz, CD₃OD) δ 8.10-7.90(m, 1H), 7.60-7.50(m, 1H), 6.90-6.35(m, 1H), 6.20-5.95(m, 1H), 5.80-5.35(m, 1H), 4.80-4.10(m, 3H), 3.60-3.50(m, 1H), 3.40-2.75(m, 10 2H), 2.50-2.40(m, 3H), 2.25-2.18(m, 1H), 1.80-1.40(m, 2H), 1.10-1.00(m, 7H)

Example 136: Preparation of 3-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropenenitrile



15

Step 1: Preparation of 2,4,5-trichloro-1H-pyrrolo[2,3-d]pyrimidine

2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine (5.0 g, 26.6 mmol) and N-chlorosuccinimide (5.3 g, 39.9 mmol) were dissolved in N,N-dimethylformamide (50.0 mL) and then stirred at room temperature for 24 hours. The organic layer was isolated, treated with 20 magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (5.5 g, yield: 93.4%).

¹H NMR (500MHz, CD₃OD) δ 7.54(s, 1H)

Step 2: Preparation of *tert*-butyl (R)-3-((2,5-dichloro-7H-pyrrolo[2,3-

d]pyrimidin-4-yl)amino)-piperidine-1-carboxylate

After 2,4,5-trichloro-7H-pyrrolo[2,3-d]pyrimidine (2.2 g, 9.9 mmol) was dissolved in N,N-dimethylformamide, sodium hydride (262.0 mg, 10.9 mmol) was added dropwise thereto at 0 °C. After stirring for 30 minutes, (2-(chloromethoxy)ethyl)trimethylsilane (1.7 mL, 9.89 mmol) was added dropwise and the mixture was stirred at room temperature for 3 hours and 30 minutes. Then, the organic layer was isolated, treated with sodium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (2.0 g, yield: 58.3%).

¹H NMR (500MHz, CD₃OD) δ 7.74(s, 1H), 5.60-5.59(m, 2H), 3.59-3.58(m, 2H), 0.91-0.89(m, 2H), 0.01(m, 9H)

Step 3: Preparation of (R)-tert-butyl 3-((2,5-dichloro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

Tert-butyl (R)-3-((2,5-dichloro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)-piperidine-1-carboxylate (2.0 g, 5.7 mmol), (R)-*tert*-butyl 3-aminopiperidine-1-carboxylate (1.4 g, 3.97 mmol) and N,N-diisopropylethylamine (1.5 mL, 5.96 mmol) were dissolved in ethanol(30.0 mL), and then stirred at 105°C for 7 hours. Subsequently, the residue was isolated by column chromatography to obtain a title compound (2.2 g, yield: 74.4%).

¹H NMR (500MHz, CD₃OD) δ 7.24(s, 1H), 5.48-5.43(m, 2H), 4.30-4.10(m, 1H), 3.80-3.50(m, 5H), 2.08-1.90(m, 1H), 1.90-1.82 (m, 1H), 1.79-1.70 (m, 1H), 1.69-1.60(m, 1H), 1.57-1.01(m, 10H), 0.90-0.81(m, 2H), 0.01(m, 9H)

Step 4: Preparation of tert-butyl (R)-3-((5-chloro-2-((3-methylisothiazol-5-yl)amino)-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

(R)-*tert*-butyl 3-((2,5-dichloro-7-((2-(trimethylsilyl)ethoxy)methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate(200.0 mg, 0.39 mmol), 3-methylisothiazol-5-amine(39.8 mg, 0.35 mmol), tris(dibenzylideneacetone)dipalladium(18.0 mg, 0.002 mmol), 2-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl (18.5 mg, 0.39 eq), and potassium

carbonate(118.0 mg, 0.86 eq) were dissolved in t-butanol, and the mixture was stirred at room temperature for 45 minutes, followed by stirring at 105°C for 24 hours. Then, the reaction solution was filtered through celite while dissolving it in methanol and then isolated by column chromatography to obtain the title compound (120.0 mg, yield: 52.0%).

5 ¹H NMR (500MHz, CD₃OD) δ 7.00(s, 1H), 6.56(s, 1H), 6.52-6.48(m, 2H), 4.55-4.20(m, 1H), 4.08-3.85(m, 1H), 3.70-3.50(m, 4H), 2.33 (s, 3H), 2.10-2.00(m, 2H), 1.98-1.55 (m, 3H), 1.50-1.10 (m, 9H), 0.91-0.83(m, 2H), 0.01(m, 9H)

Step 5: Preparation of (R)-5-chloro-N2-(3-methylisothiazol-5-yl)-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine-2,4-diamine

10 After *tert*-Butyl (R)-3-((5-chloro-2-((3-methylisothiazol-5-yl)amino)-7-((2-(trimethylsilyl)ethoxy)-methyl)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (567.0 mg, 1.76 mmol) was dissolved in dichloromethane, trifluoro acetic acid (1.5 mL) was added dropwise thereto and the mixture was stirred at 60 °C for 12 hours. After
15 disappearance of the starting material was identified by TLC, the reaction mixture was neutralized with sodium bicarbonate and extracted with ethyl acetate and H₂O. The organic layer was dried with sodium sulfate and then concentrated under reduced pressure to obtain a title compound (100.0 mg, yield : 29.0%).

¹H NMR (500MHz, CD₃OD) δ 7.05-6.92(m, 1H), 6.60-6.50(m, 1H), 4.40-4.20(m, 20 1H), 3.33-3.25(m, 1H), 2.98-2.90(m, 1H), 2.70-2.55 (m, 1H), 2.40-2.30 (m, 3H), 2.10-2.00(m, 1H), 1.90-1.72(m, 2H), 1.71-1.54(m, 2H)

Step 6: Preparation of (R)-1-(3-((5-chloro-2-((3-methylisothiazol-5-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

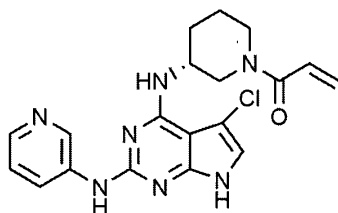
25 After (R)-5-chloro-N2-(3-methylisothiazol-5-yl)-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidine-2,4-diamine (40.0 mg, 0.11 mmol) was dissolved in a 20:3 mixed solution of tetrahydrofuran : H₂O, sodium bicarbonate (27.7 mg, 0.33 eq) was added thereto at 0 °C and then stirred for 15 minutes. Acryloyl chloride (8.9 μL, 0.11 eq) was added to the reaction mixture and then stirred at 0 °C for 15 minutes. The organic layer was isolated, treated with
30 magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was

isolated by column chromatography to obtain a title compound (4.2 mg, yield 9.2%).

¹H NMR (500MHz, CD₃OD) δ 6.88-6.50(m, 3H), 6.35-6.02(m, 1H), 5.80-5.45(m, 1H), 4.60-4.50(m, 1H), 3.90-3.75(m, 2H), 3.65-3.50(m, 1H), 2.3 (s, 3H), 2.20-2.10 (m, 1H), 1.99-1.60(m, 4H)

5

Example 137: Preparation of (R)-1-(3-((5-chloro-2-(pyridin-3-ylamino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

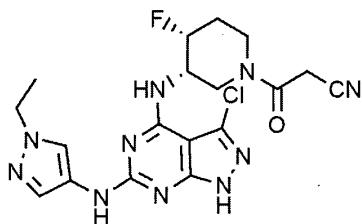


A title compound (33.1 mg, yield: 23.5%) was prepared in the same manner as in Example 136, except that pyridin-3-amine was used instead of 3-methylisothiazol-5-amine in Example 136.

¹H NMR (500MHz, CD₃OD) δ 8.95-8.88(m, 1H), 8.35-8.23(m, 1H), 8.10-8.00(m, 1H), 7.38-7.27(m, 1H), 7.02-6.50(m, 2H), 6.30-6.00(m, 1H), 5.80-5.40(m, 1H), 4.70-4.55(m, 1H), 4.40-4.27(m, 1H), 3.90-3.40(m, 3H), 2.15-1.55(m, 4H)

15

Example 138: Preparation of 3-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropanenitrile



20

Step 1: Preparation of 3,4,6-trichloro-1H-pyrazolo [3,4-d]pyrimidine

4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine (5.0 g, 26.5 mmol) and N-chlorosuccinimide (5.3 g, 39.7 mmol) were dissolved in N,N-dimethylformamide (50.0 mL), and then stirred at room temperature for 24 hours. The organic layer was isolated, treated with

magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (3.3 g, yield 56.0%).

^1H NMR (500MHz, DMSO- d_6) δ 13.07(s, 1H), 7.94(s, 1H)

5 **Step 2: Preparation of *tert*-butyl (3S,4R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate**

After 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine (350.0 mg, 1.5 mmol) was dissolved in ethanol (50 mL), N,N-diisopropylethylamine (391.8 μL , 2.3 mmol) and *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate (490.9 mg, 2.3 mmol) were added thereto. 10 The reaction mixture was stirred at 110 °C for 12 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (350.0 mg, yield: 57.7%).

^1H NMR (500MHz, CD_3OD) δ 5.10-5.00(m, 1H), 4.58-4.50(m, 1H), 3.85-3.80(m, 1H), 3.26-3.16(m, 3H), 2.10-1.89(m, 2H), 1.46(s, 9H)

15

Step 3: Preparation of (3S,4R)-3-(3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate

Tert-butyl (3S,4R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate (150.0 mg, 0.4 mmol) and 1-ethyl-1H-pyrazol-4-amine (31.6 20 mg, 0.3 mmol) were dissolved in 2-butanol (3.0 mL). Trifluoroacetic acid (26.2 μL , 0.3 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 5 hours, and then the solvent was concentrated. This reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol. The residue was isolated by column chromatography to obtain a title compound (49.8 mg, yield: 36.8%).

25 ^1H NMR (500MHz, CD_3OD) δ 7.97(s, 1H) 7.56(s, 1H), 5.10-5.00(m, 1H), 4.55-4.35(m, 1H), 4.16-4.09(m, 2H), 3.90-3.50(m, 2H), 3.16-2.95(m, 1H), 2.20-1.85(m, 3H), 1.46-1.22(m, 12H)

30 **Step 4: Preparation of 3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-((3S,4R)-4-fluoropiperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4,6-diamine hydrochloride**

To (3S,4R)-3-(3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate (45.0 mg, 0.09 mmol) was added 6 N hydrochloric acid solution (2.0 mL, excess) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction
 5 was carried out without isolation.

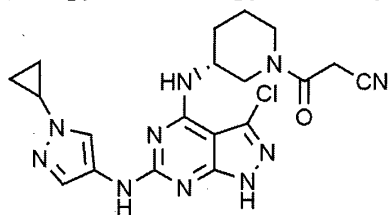
¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H) 7.56(s, 1H), 5.10-5.00(m, 1H), 4.55-4.35(m, 1H), 4.16-4.09(m, 2H), 3.90-3.50(m, 2H), 3.16-2.95(m, 1H), 2.20-1.85(m, 3H), 1.46-1.39(m, 3H)

10 **Step 5: Preparation of 3-((3S,4R)-3-(3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropanenitrile**

After 2-cyanocetic acid (20.9 mg, 0.2 mmol) was dissolved in N,N-dimethylformamide (2.0 mL), 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate (93.5 mg, 0.3 mmol), N,N-diisopropylethylamine
 15 (51.6 μL, 0.6 mmol) and 3-chloro-N6-(1-ethyl-1H-pyrazolo[3,4-d]pyrimidin-4,6-diamine hydrochloride (85.3 mg, 0.2 mmol) were added thereto. The reaction mixture was stirred at room temperature for 24 hours. The organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated in reduced pressure. The residue was isolated by column
 20 chromatography to obtain a title compound (60.0 mg, yield: 65.5%).

¹H NMR (500MHz, CD₃OD) δ 7.95(s, 1H), 7.61(s, 1H), 5.07-5.05(m, 1H), 4.49-4.37(m, 1H), 4.17-4.13(m, 2H), 3.59-3.40(m, 2H), 3.16-2.94(m, 1H), 2.80(s, 2H), 2.25-2.15(m, 1H), 2.10-1.90(m, 2H), 1.46-1.43(m, 3H)

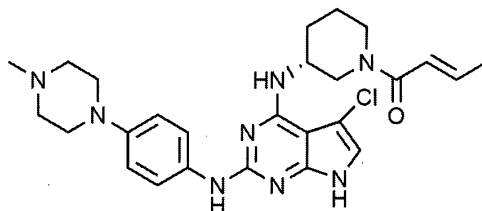
25 **Example 139: Preparation of (R)-3-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile**



A title compound (7.9 mg, yield: 35.7%) was prepared in the same manner as in Example 138, except using *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine in Example 138.

5 ¹H NMR (500MHz, CD₃OD) δ 8.00 (s, 1H), 7.58-7.54 (m, 1H), 4.59 (s, 1H), 4.50-4.00 (m, 3H), 3.64-3.59 (m, 2H), 3.19-3.14 (m, 2H), 2.20-2.01 (m, 2H), 1.91-1.80 (m, 3H), 1.72-7.60 (m, 3H)

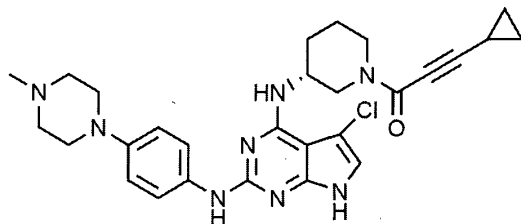
Example 140: Preparation of (R,E)-1-(3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)but-2-en-1-one



A title compound (11.6 mg, yield: 45.7%) was prepared in the same manner as in Example 138, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, 4-(4-methylpiperazin-1-yl)aniline instead of 1-ethyl-1H-pyrazol-4-amine, and (E)-but-2-enoic acid instead of 2-cyanoacetic acid in Example 138.

¹H NMR (500MHz, CD₃OD) δ 7.65-7.50(m, 2H), 6.90-6.80(m, 2H), 6.78-6.70(m, 1H), 6.65-6.45(m, 1H), 6.20-6.10(m, 1H), 4.40-4.20(m, 1H), 3.85-3.40(m, 3H), 3.20-3.05(m, 4H), 2.70-2.55(m, 4H), 2.35(s, 3H), 2.10-1.50(m, 8H)

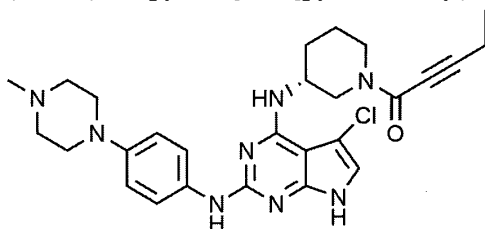
Example 141: Preparation of (R)-1-(3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-cyclopropylprop-2-yn-1-one



A title compound (7.8 mg, yield: 29.3%) was prepared in the same manner as in Example 138, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, 4-(4-methylpiperazin-1-yl)aniline instead of 1-ethyl-1H-pyrazol-4-amine, and 3-cyclopropylpropionic acid instead of 2-cyanoacetic acid in Example 138.

¹H NMR (500MHz, CD₃OD) δ 7.60-7.50(m, 2H), 7.00-6.90(m, 2H), 6.80-6.70(m, 1H), 4.40-4.20(m, 2H), 4.10-3.80(m, 2H), 3.60-3.35(m, 1H), 3.20-3.10(m, 4H), 2.70-2.60(m, 4H), 2.36(s, 3H), 2.10-1.50(m, 4H), 1.15-0.40(m, 5H)

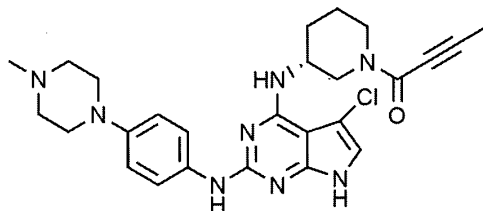
Example 142: Preparation of (R)-1-(3-((5-chloro-2-((4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)pent-2-yn-1-one



A title compound (7.5 mg, yield: 28.9%) was prepared in the same manner as in Example 138, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, 4-(4-methylpiperazin-1-yl)aniline instead of 1-ethyl-1H-pyrazol-4-amine, and pent-2-ynoic acid instead of 2-cyanoacetic acid in Example 138.

¹H NMR (500MHz, CD₃OD) δ 7.60-7.50(m, 2H), 6.97-6.90(m, 2H), 6.80-6.70(m, 1H), 4.40-3.80(m, 4H), 3.60-3.40(m, 1H), 3.20-3.10(m, 4H), 2.70-2.60(m, 4H), 2.30(s, 3H), 2.12-1.45(m, 6H), 1.25-0.80(m, 3H)

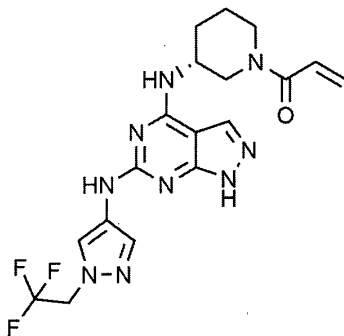
Example 143: Preparation of (R)-1-(3-((5-chloro-2-((4-(4-methylpiperazin-1-yl)phenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)but-2-yn-1-one



5 A title compound (6.2 mg, yield: 24.3%) was prepared in the same manner as in Example 138, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, 4-(4-methylpiperazin-1-yl)aniline instead of 1-ethyl-1H-pyrazol-4-amine, and but-2-ynoic acid instead of 2-cyanoacetic acid in
10 Example 138.

¹H NMR (500MHz, CD₃OD) δ 7.60-7.50(m, 2H), 7.00-6.90(m, 2H), 6.78-6.70(m, 1H), 4.40-3.70(m, 4H), 3.60-3.40(m, 1H), 3.20-3.10(m, 4H), 2.70-2.60(m, 4H), 2.37(s, 3H), 2.15-1.55(m, 7H)

15 **Example 144: Preparation of (R)-1-(3-((6-((1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one**

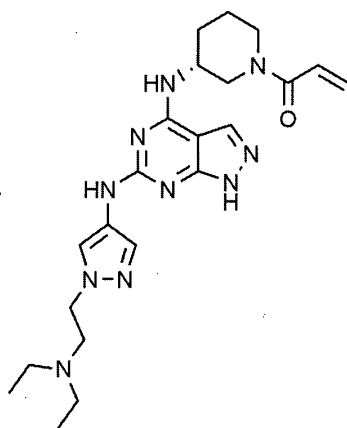


A title compound (5.3 mg, yield: 24.5%) was prepared in the same manner as in Example 30, except that 1-(2,2,2-trifluoroethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.
20

¹H NMR (500MHz, CD₃OD) δ 8.20-8.01 (m, 1H), 7.92 (s, 1H), 7.69-7.60 (m, 1H),

6.86-6.80 (m, 1H), 6.28-6.07 (m, 1H), 5.80-5.51 (m, 1H), 4.27-4.02 (m, 4H), 3.50-3.16 (m, 1H),
2.67-2.15 (m, 2H), 1.97-1.92 (m, 1H), 1.79-1.72 (m, 1H), 1.61-1.60 (m, 1H), 1.31-1.27 (m, 1H)

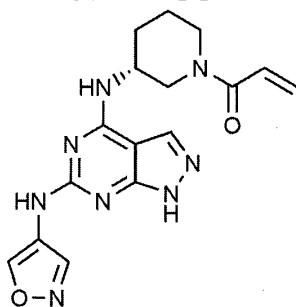
Example 145: Preparation of (R)-1-(3-((6-((1-(2-(diethylamino)ethyl)-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (4.9 mg, yield: 21.7%) was prepared in the same manner as in Example 30, except that 1-(2-(diethylamino)ethyl)-1H-pyrazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline in Example 30.

¹H NMR (500MHz, CD₃OD) δ 8.03-7.99(m, 2H), 7.61-7.60(m, 1H), 6.85-6.58(m, 1H), 6.26-6.07(m, 1H), 5.79-5.54(m, 1H), 4.25-4.20(m, 3H), 4.08-4.05(m, 2H), 3.27-3.17(m, 1H), 3.06-2.99(m, 2H), 2.72-2.66(m, 4H), 2.20-2.15(m, 2H), 2.03-1.95(m, 1H), 1.80-1.66(m, 2H), 1.10(t, 6H)

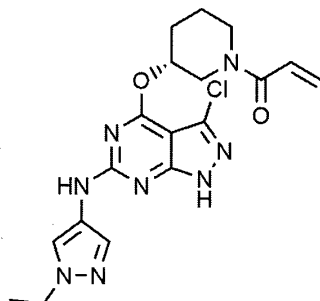
Example 146: Preparation of (R)-1-(3-((6-(isoxazol-4-ylamino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (7.1 mg, yield: 39.9%) was prepared in the same manner as in Example 30, except that isoxazol-4-amine was used instead of 4-(4-methylpiperazin-1-yl)aniline.

¹H NMR (500MHz, CD₃OD) δ 9.07(s, 1H), 8.51-8.50(m, 1H), 7.94-7.92(m, 1H),
 5 6.85-6.55(m, 1H), 6.27-6.09(m, 1H), 5.79-5.56(m, 1H), 4.27-4.04(m, 3H), 3.48-3.20(m, 2H),
 2.20-2.16(m, 2H), 1.80-1.61(m, 2H)

Example 147: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one



10

Step 1: Preparation of 3,4,6-trichloro-1H-pyrazolo [3,4-d]pyrimidine

4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine (2.5 g, 13.3 mmol) and N-chlorosuccinimide (2.7 g, 19.9 mmol) were dissolved in N,N-dimethylformamide (30.0 mL), and then stirred at room temperature for 24 hours. The organic layer was isolated, treated with
 15 magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was
 isolated by column chromatography to obtain a title compound (1.6 g, yield: 56.0%).

Step 2: Preparation of 3,4,6-trichloro-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidine

20 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine(1.3 g, 5.8 mmol) was dissolved in N,N-dimethylformamide (10.0 mL), to which sodium hydride(207.1 mg, 8.6 eq) was added and then stirred for 30 minutes. To this reaction mixture was added (2-(chloromethoxy)ethyl)trimethylsilane (840.0 μL, 5.8 mmol), followed by reacting at room temperature for 3 hours, to which water was added and extracted with ethyl acetate. The isolated
 25 organic layer was dried over anhydrous magnesium sulfate and then concentrated under

reduced pressure. The obtained residue was isolated by column chromatography to obtain a title compound (1.6 g, yield: 81.0%).

^1H NMR (500MHz, CDCl_3) δ 5.69(s, 2H), 3.66(t, 2H), 0.92(t, 2H), -0.05(s, 9H)

5 **Step 3: Preparation of *tert*-butyl (R)-3-((3,6-dichloro-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate**

Tert-butyl (R)-3-hydroxypiperidine-1-carboxylate (800.0 mg, 2.26 mmol) was dissolved in tetrahydrofuran(10.0 mL) and then cooled to 0 °C in an ice bath. Sodium hydride
10 (303.2 mg, 4.52 mmol) was added to the reaction mixture and then stirred for 30 minutes. Subsequently, 3,4,6-trichloro-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidine (1.3 g, 2.0 eq) was added and then stirred for 2 hours, to which water was added and extracted with ethyl acetate. The isolated organic layer was dried over anhydrous magnesium sulfate and then concentrated under reduced pressure. The obtained residue was
15 isolated by column chromatography to obtain a title compound (1.3 g, yield: 67.7%).

^1H NMR (500MHz, CDCl_3) δ 5.64(s, 2H), 5.47-5.35(m, 1H), 4.02-4.00(m, 1H), 3.77-3.74(m, 2H), 3.64(t, 2H), 3.51-3.40(m, 1H), 3.21(s, 1H), 2.02-1.98(m, 2H), 1.61(s, 1H), 1.44-1.21(m, 10H), 0.92(t, 2H), -0.05(s, 9H)

20 **Step 4: Preparation of *tert*-butyl (R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate**

After *tert*-butyl (R)-3-((3,6-dichloro-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate(250.0 mg, 0.48 mmol) was
25 dissolved in *tert*-butanol (3.0 mL), 1-ethyl-1H-pyrazol-4-amine (64. mg, 0.53 mmol), tris(dibenzylideneacetone)dipalladium(44.9 mg, 0.024 mmol), 2-dicyclohexylphosphino-2',4',6'-triisopropylbiphenyl(23.8 mg, 0.048 mmol), and potassium carbonate (135.4 mg, 0.96 mmol) were added thereto and then reacted at 110 °C for 12 hours, to which water was added and extracted with ethyl acetate. The isolated organic layer was dried over anhydrous
30 magnesium sulfate and then concentrated under reduced pressure. The obtained residue was

isolated by column chromatography to obtain a title compound (182.0 mg, yield: 62.5%).

¹H NMR (500MHz, CDCl₃) δ 7.54(s, 1H), 7.30-7.06(m, 1H), 5.54(s, 1H), 5.21(s, 1H), 4.17-4.12(m, 2H), 3.98-3.68(m, 2H), 3.64(t, 2H), 3.46-3.45(m, 1H), 3.21-3.20(m, 1H), 2.12-1.96(m, 3H), 1.60-1.22(m, 13H), 0.92(t, 2H), -0.05(s, 9H)

5

Step 5: Preparation of *tert*-butyl (R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate

After *tert*-butyl (R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1-((2-(trimethylsilyl)ethoxy)methyl)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate (182.0 mg, 0.30 mmol) was dissolved in tetrahydrofuran (2.0 mL), tetrabutylammonium fluoride (2.0 mL, excessive amount) was added thereto and then stirred for 20 hours. Water was then added and extracted with ethyl acetate. The isolated organic layer was dried over anhydrous magnesium sulfate and then concentrated under reduced pressure. The obtained residue was isolated by column chromatography to obtain a title compound (126.0 mg, yield: 87.5%).

¹H NMR (500MHz, CDCl₃) δ 7.56 (s, 1H), 7.29-7.05(m, 1H), 5.19(s, 1H), 3.95-3.65(m, 2H), 3.64(t, 2H), 3.46-3.45(m, 1H), 3.23-3.20(m, 1H), 2.10-1.94(m, 3H), 1.59-1.20(m, 13H)

Step 6: Preparation of (R)-3-chloro-N-(1-ethyl-1H-pyrazol-4-yl)-4-(piperidin-3-yloxy)-1H-pyrazolo[3,4-d]pyrimidin-6-amine hydrochloride

Tert-butyl (R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carboxylate (126.0 mg, 0.27 mmol) was dissolved in 1,4 dioxane (0.5 mL), followed by the addition of 4 N HCl in dioxane (3.0 mL, excessive amount). The mixture was then stirred at room temperature for 2 hours. The reaction product was concentrated to obtain a title compound (98.1 mg, yield: 100%).

¹H NMR (500MHz, CDCl₃) δ 7.58 (s, 1H), 7.25-7.06(m, 1H), 5.24(s, 1H), 3.94-3.64(m, 5H), 3.50-3.45(m, 1H), 3.25-3.20(m, 1H), 2.12-1.96(m, 3H), 1.60-1.22(m, 4H)

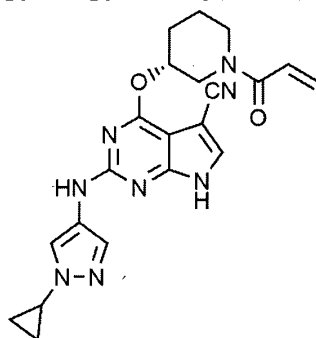
Step 7: Preparation of (R)-1-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-

1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one

After (R)-3-chloro-N-(1-ethyl-1H-pyrazol-4-yl)-4-(piperidin-3-yloxy)-1H-pyrazolo[3,4-d]pyrimidin-6-amine hydrochloride (30.0 mg, 0.07 mmol) was dissolved in THF: H₂O = 3:1 (1.0/0.3 mL), sodium bicarbonate (20.7 mg, 0.22 mmol) was added thereto at -20 °C and then stirred for 30 minutes. Acryloyl chloride (8.0 µL, 0.8 mmol) was added to the reaction mixture, followed by stirring at -20 °C for 1 hour, to which water was added and extracted with ethyl acetate. The isolated organic layer was dried over anhydrous magnesium sulfate and then concentrated under reduced pressure. The obtained residue was isolated by column chromatography to obtain a title compound (4.9 mg, yield: 15.3%).

¹H NMR (500MHz, CD₃OD) δ 8.05(s, 1H), 7.62(s, 1H), 6.85-6.55(m, 1H), 6.17-6.04(m, 1H), 6.01-5.72(m, 1H), 5.61-5.44(m, 2H), 4.30-4.28(m, 1H), 4.17-4.13(m, 2H), 3.75-3.68(m, 2H), 2.09-2.01(m, 3H), 1.67-1.65(m, 1H), 1.47-1.44(m, 3H)

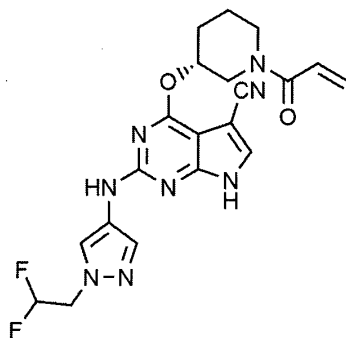
Example 148: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile



A title compound (7.8 mg, yield: 37.5%) was prepared in the same manner as in Example 147, except using 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine, and 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 8.02-8.01(m, 1H), 7.58-7.55(m, 1H), 6.80-6.40(m, 1H), 6.13-5.92(m, 1H), 5.72-5.36(m, 2H), 4.70-4.10(m, 2H), 3.73-3.57(m, 3H), 2.13-2.07(m, 3H), 1.66-1.65(m, 1H), 1.09-1.01(m, 4H)

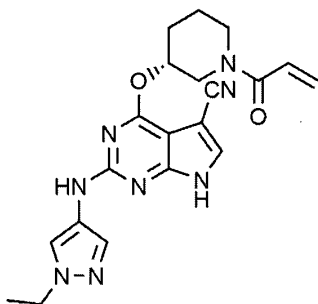
Example 149: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile



A title compound (9.0 mg, yield: 40.5%) was prepared in the same manner as in
 5 Example 147, except using 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine in Example 147, and 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine in Step 2.

¹H NMR (500MHz, CD₃OD) δ 8.06(s, 1H), 7.67-7.58(m, 2H), 6.80-6.39(m, 1H),
 6.14-6.11(m, 1H), 5.94-5.40(m, 1H), 5.37-5.35(m, 1H), 4.53-4.47(m, 2H), 4.23-4.17(m, 1H),
 10 3.78-3.55(m, 2H), 3.30-3.23(m, 2H), 2.11-2.00(m, 3H), 1.65-1.64(m, 1H)

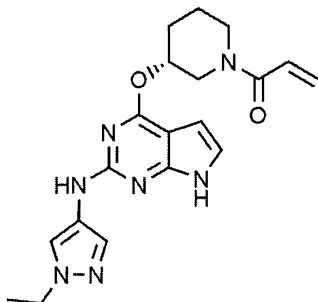
Example 150: Preparation of (R)-4-((1-acryloylpiperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile



15 A title compound (4.3 mg, yield: 21.1%) was prepared in the same manner as in Example 147, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 7.98(s, 1H), 7.60-7.57(m, 2H), 6.81-6.41(m, 1H),
 6.13-5.92(m, 1H), 5.72-5.39(m, 2H), 4.25-4.09(m, 4H), 3.85-3.55(m, 3H), 2.20-2.00(m, 3H),
 20 1.66-1.65(m, 1H), 1.46-1.44(m, 3H)

Example 151: Preparation of (R)-1-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one

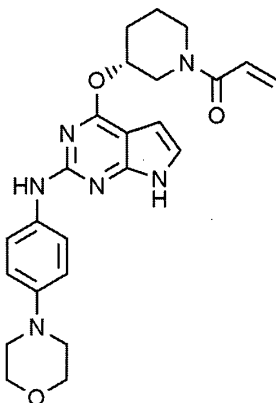


5 A title compound (5.9 mg, yield: 30.9%) was prepared in the same manner as in Example 147, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 7.98-7.96(m, 1H), 7.57-7.55(m, 1H), 6.84-6.50(m, 2H), 6.25-6.05(m, 2H), 5.70-5.47(m, 2H), 4.16-4.12(m, 4H), 3.77-3.59(m, 2H), 2.03-1.92(m, 3H), 1.66-1.64(m, 1H), 1.47-1.44(m, 3H)

10

Example 152: Preparation of (R)-1-(3-((2-((4-morpholinophenyl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one

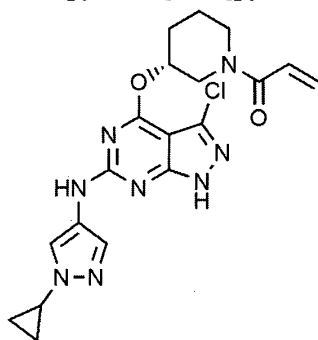


15 A title compound (7.2 mg, yield: 32.3%) was prepared in the same manner as in Example 147, except using 4-morpholinoaniline instead of 1-ethyl-1H-pyrazol-4-amine, and 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 7.57-7.55(m, 2H), 6.93-6.91(m, 2H), 6.84-6.83(m, 1H), 6.26-6.03(m, 2H), 6.00-5.75(m, 1H), 5.61-5.44(m, 1H), 5.35-5.34(m, 1H), 4.08-4.05(m, 2H), 3.83-3.81(m, 4H), 3.75-3.72(m, 2H), 3.05-3.04(m, 4H), 2.07-1.94(m, 3H), 1.70-1.60(m, 1H).

5

Example 153: Preparation of (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one

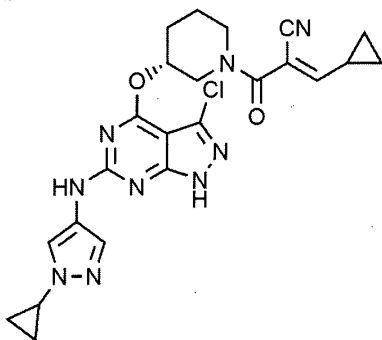


A title compound (7.5 mg, yield: 35.2%) was prepared in the same manner as in
 10 Example 147, except that 1-cyclopropyl-1H-pyrazol-4-amine was used instead of 1-ethyl-1H-pyrazol-4-amine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 8.01(s, 1H), 7.58-7.57(m, 1H), 7.80-7.50(m, 1H), 6.17-6.01(m, 2H), 5.74-5.58(m, 1H), 4.32-4.29(m, 2H), 3.69-3.58(m, 3H), 2.07-1.99(m, 3H), 1.66-1.64(m, 1H), 1.09-1.00(m, 4H)

15

Example 154: Preparation of (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile



Step 1: Preparation of (R)-3-chloro-N-(1-cyclopropyl-1H-pyrazol-4-yl)-4-(piperidin-3-yloxy)-1H-pyrazolo[3,4-d]pyrimidin-6-amine hydrochloride

A title compound (102.5 mg, yield: 45.8%) was prepared in the same manner as in Example 147, except that 1-cyclopropyl-1H-pyrazol-4-amine was used instead of 1-ethyl-1H-pyrazol-4-amine in Example 147.

¹H NMR (500MHz, CD₃OD) δ 8.25(s, 1H), 7.88(s, 1H), 5.61(s, 1H), 3.73-3.71(m, 2H), 3.68-3.58(m, 1H), 3.52-3.49(m, 2H), 3.24-3.18(m, 1H), 2.26-2.19(m, 1H), 1.96-1.93(m, 1H), 1.19-1.13(m, 4H)

Step 2: Preparation of (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one

After (R)-3-chloro-N-(1-cyclopropyl-1H-pyrazol-4-yl)-4-(piperidin-3-yloxy)-1H-pyrazolo[3,4-d]pyrimidin-6-amine hydrochloride (85.3 mg, 0.24 mmol), and 2-cyanocetic acid (20.0 mg, 0.24 mmol) were dissolved in N,N-dimethylformamide (2.0 mL), 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate (112.2 mg, 0.36 mmol) was added thereto. The reaction mixture was stirred at room temperature for 24 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (72.0 mg, yield 65.5%).

¹H NMR (500MHz, CD₃OD) δ 8.21(s, 1H), 7.84(s, 1H), 5.60(s, 1H), 3.73-3.71(m, 2H), 3.68-3.58(m, 1H), 3.51-3.48(m, 2H), 3.31(s, 2H), 3.23-3.15(m, 1H), 2.24-2.18(m, 1H), 1.94-1.90(m, 1H), 1.15-1.10(m 4H)

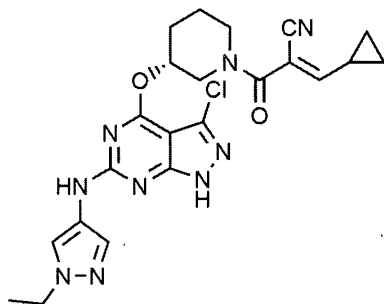
Step 3: Preparation of (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile

After (R)-1-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidin-1-yl)prop-2-en-1-one (50.0 mg, 0.12 mmol) was dissolved in methanol, piperidine (23.0 μL, 0.23 mmol) and cyclopropanecarbaldehyde (13.2 μL, 0.18 mmol) were added thereto. After stirring at room temperature for 5 hours, the organic layer was

isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (9.6 mg, yield 33.2%).

¹H NMR (500MHz, CD₃OD) δ 8.07(s, 1H), 7.60(s, 1H), 6.27-6.24(m, 1H), 5.48(s, 1H), 4.57(s, 2H), 3.61-3.51(m, 2H), 2.12-2.01(m, 3H), 1.76-1.70(m, 3H), 1.13-1.03(m, 6H), 0.99-0.89(m, 3H)

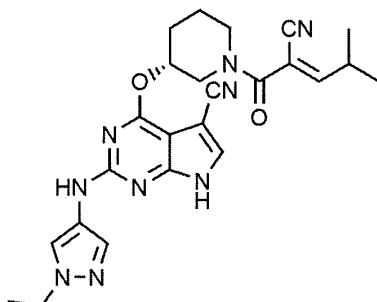
Example 155: Preparation of (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile



A title compound (11.2 mg, yield: 46.3%) was prepared in the same manner as in Example 154, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 154.

¹H NMR (500MHz, CD₃OD) δ 8.03(s, 1H), 7.62(s, 1H), 6.27-6.25(m, 1H), 5.49(s, 1H), 4.63-4.41(m, 2H), 3.65-3.45(m, 2H), 2.13-2.03(m, 4H), 1.69-1.59 (m, 3H), 1.32(s, 3H), 1.13-0.89(m, 4H)

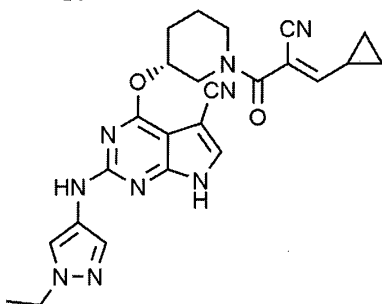
Example 156: Preparation of (R,E)-4-((1-(2-cyano-4-methylpent-2-enoyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile



A title compound (4.5 mg, yield: 15.3%) was prepared in the same manner as in Example 154, except using (R)-4-((1-(2-cyanoacetyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile instead of (R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile, and isobutyraldehyde instead of cyclopropanecarbaldehyde in Example 154.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.62-7.61(m, 2H), 6.92-6.52(m, 1H), 5.51(s, 1H), 4.50-3.90(m, 4H), 3.63-3.54(m, 1H), 3.02-2.81(m, 1H), 2.55-2.03(m, 4H), 1.72-1.71(m, 1H), 1.45(t, 3H), 1.16-1.15(m, 3H), 0.89-0.88(m, 3H)

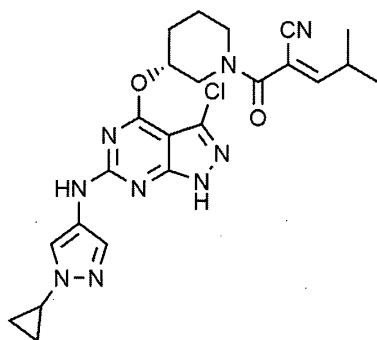
Example 157: Preparation of (R,E)-4-((1-(2-cyano-3-cyclopropylacryloyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile



A title compound (5.3 mg, yield: 16.8%) was prepared in the same manner as in Example 154, except that (R)-4-((1-(2-cyanoacetyl)piperidin-3-yl)oxy)-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidine-5-carbonitrile was used instead of ((R,E)-2-(3-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile in Example 154.

¹H NMR (500MHz, CD₃OD) δ 7.98(s, 1H), 7.60-7.59(m, 2H), 6.69-6.22(m, 1H), 5.49-5.45(m, 1H), 4.56-4.44(m, 1H), 4.37-4.00(m, 3H), 3.51-3.00(m, 2H), 2.15-2.13(m, 3H), 1.33-1.32(m, 1H), 1.50-1.54(m, 3H), 1.01-0.89(m, 5H)

5 **Example 158: Preparation of (R,E)-2-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)oxy)piperidine-1-carbonyl)-4-methylpent-2-enenitrile**

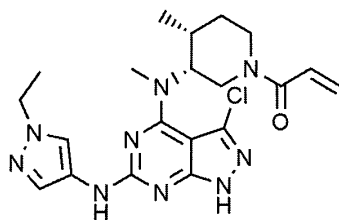


A title compound (11.8 mg, yield: 47.4%) was prepared in the same manner as in
10 Example 154, except that isobutyraldehyde was used instead of cyclopropanecarbaldehyde in Example 154.

¹H NMR (500MHz, CD₃OD) δ 8.07(s, 1H), 7.58-7.57(m, 1H), 6.80-6.60(m, 1H), 5.54(s, 1H), 4.38-3.90(m, 2H), 3.61-3.45(m, 3H), 3.02-2.60(m, 1H), 2.13-2.03(m, 3H), 1.73-1.72(m, 1H), 1.15-1.13(m, 4H), 1.07-1.04(m, 6H)

15

Example 159: Preparation of (1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one



20

Step 1: Preparation of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine

4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine (1.0 g, 5.3 mmol), and N-

chlorosuccinimide (1.0 g, 7.9 mmol) were dissolved in N,N-dimethylformamide (15.0 mL), and then stirred at room temperature for 8 hours. Purified water was added until crystals were formed, and then stirred for 10 minutes. Thereafter, purified water was added until a large amount of crystals was shown, followed by filtering while washing with purified water. The
5 filtered solid was dissolved in excessive amount of ethyl acetate and dichloromethane, treated with sodium sulfate, washed with ethyl acetate and filtered. The organic layer was concentrated under reduced pressure to obtain a title compound (1.1 g, yield: 93.4%).

**Step 2: Preparation of N-((3R,4R)-1-benzyl-4-methylpiperidin-3-yl)-3,6-dichloro-
10 N-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-amine**

After (3R,4R)-1-benzyl-N,4-dimethylpiperidin-3-amine dihydrochloride (195.5 mg, 0.6 mmol) was dissolved in ethanol (5.0 ml), N,N-diisopropylethylamine (350.8 μ L, 2.0 mmol) was added dropwise thereto, and the mixture was stirred at room temperature for 10 minutes. 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine (50.0 mg, 0.4 mmol) was added, the temperature
15 was raised to 100 °C and stirring was further carried out for 2 hours. Thereafter, the solution was filtered under reduced pressure, and the obtained residue was isolated by column chromatography to obtain a title compound (45.1 mg, yield: 24.9%).

^1H NMR (500MHz, CD_3OD) δ 7.32-7.21(m, 5H), 5.11-5.07(m, 1H), 3.69(s, 3H), 3.52-3.48(m, 2H), 2.98-2.70(m, 2H), 2.62-2.59(m, 1H), 2.25-2.14(m, 2H), 1.73-1.72(m, 2H),
20 0.95-0.94(d, J = 5 Hz, 3H)

**Step 3: Preparation of N4-((3R,4R)-1-benzyl-4-methylpiperidin-3-yl)-3-chloro-
N6-(1-ethyl-1H-pyrazol-4-yl)-N4-methyl-1H-pyrazolo[3,4-d]pyrimidine-4,6-diamine**

N-((3R,4R)-1-benzyl-4-methylpiperidin-3-yl)-3,6-dichloro-N-methyl-1H-
25 pyrazolo[3,4-d]pyrimidin-4-amine (54.5 mg, 0.1 mmol) and 1-ethyl-1H-pyrazol-4-amine (11.5 mg, 0.1 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (9.5 μ L, 0.1 mmol) was added to the reaction mixture, followed by reacting at 190 °C for 15 hours, and then the reactant was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to
30 obtain a title compound (10.4 mg, yield: 16.1%).

¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.57(s, 1H), 7.35-7.21(m, 5H), 5.04-5.01(m, 1H), 4.16-4.11(m, 2H), 3.63(s, 3H), 3.54-3.50(m, 2H), 3.01-2.77(m, 2H), 2.68-2.59(m, 1H), 2.28-2.12(m, 2H), 1.76-1.74(m, 2H), 1.47-1.42(m, 3H), 0.95-0.94(d, *J* = 5 Hz, 3H)

5 **Step 4: Preparation of 3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-methyl-N4-((3R,4R)-4-methylpiperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidine-4,6-diamine**

N4-((3R,4R)-1-benzyl-4-methylpiperidin-3-yl)-3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-methyl-1H-pyrazolo[3,4-d]pyrimidine-4,6-diamine (10.4 mg, 0.02 mmol) was dissolved in methanol(1.0 mL), to which Pd/C (3.0 mg) was added and H₂ gas was added. The reaction
10 mixture was stirred at room temperature for 3 hours and filtered through celite. The filtrate was concentrated to obtain a title compound (9.0 mg, yield 100%).

¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.57(s, 1H), 5.04-5.01(m, 1H), 4.49-4.41(m, 2H), 3.54(s, 3H), 3.12-2.72(m, 2H), 2.68-2.59(m, 1H), 2.20-2.01(m, 2H), 1.60-1.58(m, 2H), 1.48-1.43(m, 3H), 0.90-0.89 (d, *J* = 5 Hz, 3H)

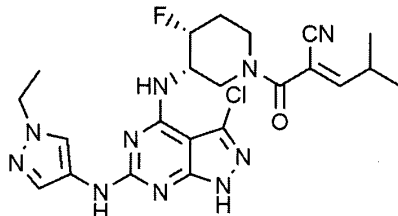
15

Step 5: Preparation of 1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)-4-methylpiperidin-1-yl)prop-2-en-1-one

3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-methyl-N4-((3R,4R)-4-methylpiperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidine-4,6-diamine (9.0 mg, 0.02 mmol) was dissolved in a 3:1
20 mixed solution of tetrahydrofuran: H₂O (1.0 mL), sodium bicarbonate (5.8 mg, 0.07 mmol) was added thereto at 0 °C and then stirred for 10 minutes. Acryloyl chloride (1.9 μL, 0.02 mmol) was slowly added dropwise to the reaction mixture, and then stirred at 0 °C for 10 minutes. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under
25 reduced pressure. The residue was isolated by column chromatography to obtain a title compound (4.7 mg, yield: 45.9%).

¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.57(s, 1H), 6.88-6.78(m, 1H), 6.24-6.20(m, 1H), 5.78-5.71(m, 1H), 4.99-4.92(m, 1H), 4.24-4.21(m, 1H), 4.16-4.11(m, 2H), 3.92-3.87(m, 1H), 3.60-3.56(m, 1H), 3.37(s, 3H), 2.44-2.42(m, 1H), 2.20-2.19(m, 1H), 1.81-1.78(m,
30 2H), 1.46-1.44(m, 3H), 1.07-1.06(d, *J* = 5 Hz, 3H)

Example 160: Preparation of (E)-2-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carbonyl)-4-methylpent-2-enenitrile



5

Step 1: Preparation of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine

4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine (10.0 g, 53.0 mmol) and N-chlorosuccinimide (10.6 g, 79.4 mmol) were dissolved in N,N-dimethylformamide (100.0 mL) and then stirred at room temperature for 24 hours. The organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (6.6 g, yield: 56.0%).

10

^1H NMR (500MHz, DMSO- d_6) δ 13.07(s, 1H), 7.94(s, 1H)

Step 2: Preparation of *tert*-butyl (3S,4R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate

15

After 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine (700.0 mg, 3.0 mmol) was dissolved in ethanol (100 mL), N,N-diisopropylethylamine (783.6 μL , 4.6 mmol) and *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate (981.8 mg, 4.6 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (700.0 mg, yield: 57.7%).

20

^1H NMR (500MHz, CD_3OD) δ 5.10-5.00(m, 1H), 4.58-4.50(m, 1H), 3.85-3.80(m, 1H), 3.26-3.16(m, 3H), 2.10-1.89(m, 2H), 1.46(s, 9H)

25

Step 3: Preparation of (3S,4R)-3-(3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate

(3S,4R)-3-((3,6-dichloro-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate (300.0 mg, 0.8 mmol) and 1-ethyl-1H-pyrazol-4-amine (63.2 mg, 0.6 mmol) were dissolved in 2-butanol (6.0 mL). Trifluoroacetic acid (52.4 μ L, 0.6 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 5 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to obtain a title compound (99.6 mg, yield: 36.8%).

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H) 7.56(s, 1H), 5.10-5.00(m, 1H), 4.55-4.35(m, 1H), 4.16-4.09(m, 2H), 3.90-3.50(m, 2H), 3.16-2.95(m, 1H), 2.20-1.85(m, 3H), 1.46-1.22(m, 12H)

Step 4: Preparation of 3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-((3S,4R)-4-fluoropiperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4,6-diamine hydrochloride

To (3S,4R)-3-(3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carboxylate (90.0 mg, 0.18 mmol) was added 6 N hydrochloric acid solution (4.0 mL, excessive amount) dissolved in methanol. After stirring at room temperature for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H) 7.56(s, 1H), 5.10-5.00(m, 1H), 4.55-4.35(m, 1H), 4.16-4.09(m, 2H), 3.90-3.50(m, 2H), 3.16-2.95(m, 1H), 2.20-1.85(m, 3H), 1.46-1.39(m, 3H)

Step 5: Preparation of 3-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropenenitrile

After 2-cyanoacetic acid (41.8 mg, 0.4 mmol) was dissolved in N,N-dimethylformamide (4.0 mL), 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate (187.0 mg, 0.6 mmol), N,N-diisopropylethylamine (103.2 μ L, 1.2 mmol) and 3-chloro-N6-(1-ethyl-1H-pyrazol-4-yl)-N4-((3S,4R)-4-fluoropiperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidin-4,6-diamine hydrochloride (170.6 mg, 0.4

mmol) were added thereto. The reaction mixture was stirred at room temperature for 24 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (120.0 mg, yield: 65.5%).

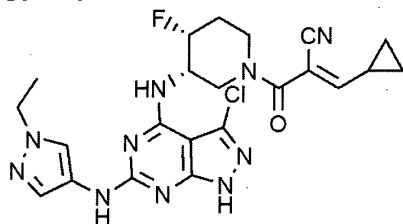
5 ^1H NMR (500MHz, CD_3OD) δ 7.95(s, 1H), 7.61(s, 1H), 5.07-5.05(m, 1H), 4.49-4.37(m, 1H), 4.17-4.13(m, 2H), 3.59-3.40(m, 2H), 3.16-2.94(m, 1H), 2.80(s, 2H), 2.25-2.15(m, 1H), 2.10-1.90(m, 2H), 1.46-1.43(m, 3H)

Step 6: Preparation of (E)-2-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carbonyl)-4-methylpent-2-enenitrile

After 3-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)-3-oxopropenenitrile (25.0 mg, 0.06 mmol) was dissolved in methanol, piperidine (11.1 μL , 0.08 mmol) and isobutylaldehyde (67.7 mg, 0.08 mmol) were added thereto. After stirring at room temperature for 5 hours, the organic layer was isolated, treated with magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (4.8 mg, yield: 17.1%).

20 ^1H NMR (500MHz, CD_3OD) δ 7.98(s, 1H), 7.56(s, 1H), 6.71(d, 1H), 5.18-5.09(m, 1H), 4.82-4.60(m, 1H), 4.35-4.25(m, 1H), 4.16-4.12(m, 2H), 4.08-3.80(m, 1H), 3.70-3.50(m, 1H), 3.22-3.20(m, 1H), 2.85-2.65(m, 1H), 2.30-2.00(m, 1H), 1.89-1.74(m, 1H), 1.58-1.54(m, 3H), 1.19-1.13(m, 6H).

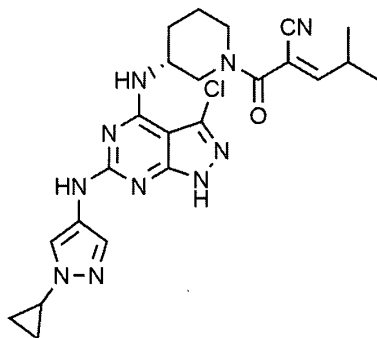
Example 161: Preparation of (E)-2-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidine-1-carbonyl)-3-cyclopropylacrylonitrile



A title compound (3.6 mg, yield: 12.9%) was prepared in the same manner as in Example 160, except that cyclopropanecarbaldehyde was used instead of isobutylaldehyde in Example 160.

¹H NMR (500MHz, CD₃OD) δ 8.01(s, 1H), 7.60(s, 1H), 6.70-6.55(m, 1H), 4.65-4.58(m, 1H), 4.16-4.11(m, 2H), 3.94-3.90(m, 1H), 3.51-3.44(m, 1H), 2.22-2.00(m, 2H), 1.89-1.58(m, 2H), 1.46-1.43(m, 3H), 1.28-1.18(m, 1H), 0.98-0.90(m, 4H)

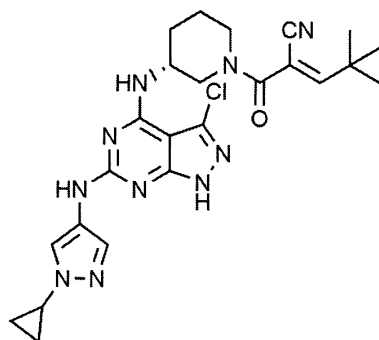
Example 162: Preparation of (R,E)-2-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile



A title compound (7.0 mg, yield: 28.2%) was prepared in the same manner as in Example 160, except using 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine, and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.96(s, 1H), 7.54(s, 1H), 6.96-6.65(m, 1H), 3.64-3.60(m, 2H), 3.59-3.48(m, 1H), 2.20(s, 1H), 1.92(s, 1H), 1.91-1.88(m, 2H), 1.71-1.69(m, 2H), 1.606-1.59 (m, 2H), 1.65-1.05(m, 6H), 0.92-0.89(m, 4H)

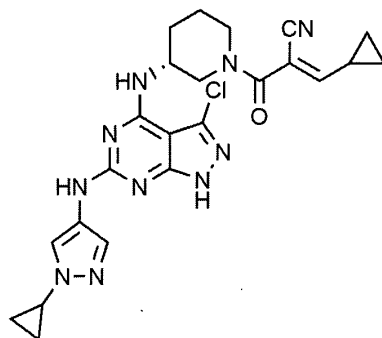
Example 163: Preparation of (R,E)-2-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4,4-dimethylpent-2-enenitrile



A title compound (9.7 mg, yield: 38.1%) was prepared in the same manner as in Example 160, except using *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, pivalaldehyde instead of isobutyraldehyde, and 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.98(s, 1H), 7.56(s, 1H), 6.92-6.62(m, 1H), 4.38(s, 1H), 3.61-3.55(m, 2H), 2.15(s, 1H), 2.19-1.99(m, 2H), 1.98-1.94(m, 1H), 1.89-1.82(m, 2H), 1.76-1.68(m, 1H), 1.39-1.31(m, 9H), 1.09-0.99(m, 4H)

Example 164: Preparation of (R,E)-2-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile

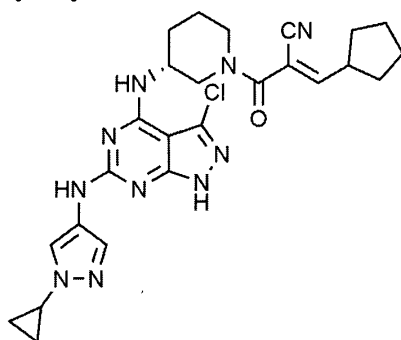


A title compound (11.6 mg, yield: 47.2%) was prepared in the same manner as in Example 160, except using *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, cyclopropanecarbaldehyde instead of isobutyraldehyde, and 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.99(s, 1H), 7.55(s, 1H), 6.65-6.34(m, 1H), 4.57(s,

1H), 4.35(s, 1H), 2.07(s, 1H), 2.06-1.99(m, 1H), 1.92-1.88(m, 2H), 1.85-1.83(m, 1H), 1.76(s, 1H), 1.52-1.48(m, 2H), 1.20-1.15(m, 2H), 1.08-1.03(m, 4H), 0.99-0.87(m, 4H)

Example 165: Preparation of (R,E)-2-((3-chloro-6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopentylacrylonitrile

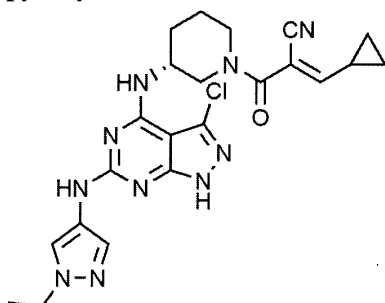


A title compound (8.7 mg, yield: 33.5%) was prepared in the same manner as in Example 160, except using *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, cyclopentanecarbaldehyde instead of isobutyraldehyde, and 1-cyclopropyl-1H-pyrazol-4-amine instead of 1-ethyl-1H-pyrazol-4-amine in Example 160.

¹H NMR (500MHz, CD₃OD) δ 8.00 (s, 1H), 7.55 (s, 1H), 7.00-6.65 (m, 1H), 4.39 (s, 1H), 3.62-3.57 (m, 5H), 2.19-1.88 (m, 5H), 1.74-1.28 (m, 8H), 0.91-0.88 (m, 5H)

15

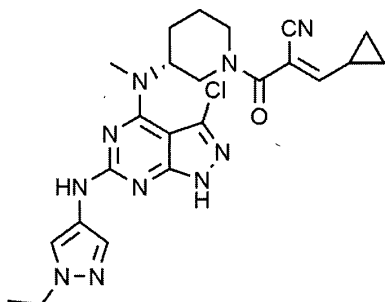
Example 166: Preparation of (R,E)-2-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile



A title compound (8.9 mg, yield: 36.9%) was prepared in the same manner as in Example 160, except using *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

5 ^1H NMR (500MHz, CD_3OD) δ 7.97(s, 1H), 7.58(s, 1H), 6.60-6.30(m, 1H), 4.36-4.35(m, 1H), 4.14-4.13(m, 2H), 3.79-3.47(m, 3H), 2.13-2.02(m, 2H), 1.95-1.89(m, 2H), 1.73-1.72(m, 2H), 1.48-1.40(m, 3H), 1.20-0.80(m, 4H)

Example 167: Preparation of (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile

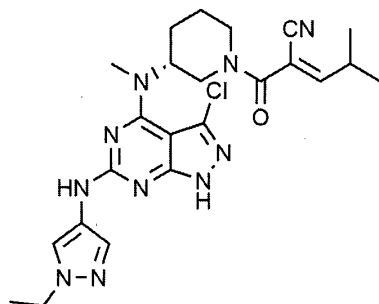


A title compound (6.1 mg, yield: 24.8%) was prepared in the same manner as in Example 160, except using *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

15 ^1H NMR (500MHz, CD_3OD) δ 7.94(s, 1H), 7.58(s, 1H), 6.50(s, 1H), 4.61-4.55(m, 2H), 4.14-3.98(m, 3H), 3.30-3.26(m, 3H), 2.19-2.03(m, 5H), 1.71-1.60(m, 2H), 1.45(t, 3H), 0.89-0.88(m, 4H)

20

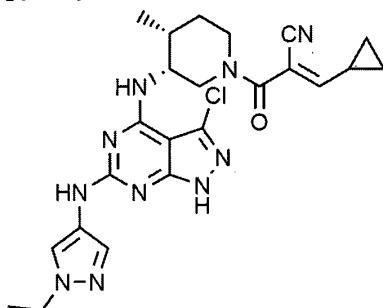
Example 168: Preparation of (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)(methyl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile



A title compound (4.9 mg, yield: 19.7%) was prepared in the same manner as in Example 160, except that *tert*-butyl (R)-3-(methylamino)piperidine-1-carboxylate was used instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.95(s, 1H), 7.63-7.59(m, 1H), 6.90-6.60(m, 1H), 4.58-4.56(m, 2H), 4.15-4.12(m, 2H), 3.34(s, 3H), 3.23-3.20(m, 1H), 2.05-2.01(m, 3H), 1.97-1.94(m, 1H), 1.85-1.60(m, 2H), 1.45(t, 3H), 1.14-1.00(m, 6H)

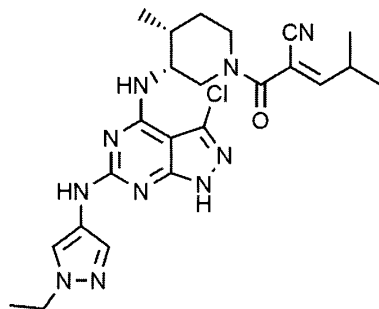
Example 169: Preparation of (E)-2-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carbonyl)-3-cyclopropylacrylonitrile



A title compound (11.1 mg, yield: 44.9%) was prepared in the same manner as in Example 160, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.96(s, 1H), 7.60(s, 1H), 6.30-6.24(m, 1H), 4.64-4.63(m, 1H), 4.32-4.31(m, 1H), 4.15-4.13(m, 2H), 3.33-3.32(m, 1H), 2.98-2.96(m, 1H), 2.22-2.02(m, 3H), 1.90-1.57(m, 4H), 1.45(t, 3H), 1.08-1.00(m, 6H)

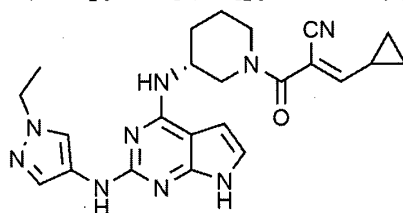
Example 170: Preparation of (E)-2-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidine-1-carbonyl)-4-methylpent-2-enenitrile



5 A title compound (10.7 mg, yield: 43.1%) was prepared in the same manner as in Example 160, except that *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate was used instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.60-7.58(m, 1H), 6.70-6.40(m, 1H), 4.44-4.40(m, 2H), 4.15-4.13(m, 2H), 3.46-3.37(m, 1H), 3.09-3.08(m, 1H), 2.56-2.55(m, 1H),
10 2.23-2.20(m, 1H), 2.19-2.17(m, 1H), 1.89-1.80(m, 1H), 1.70-1.51(m, 2H), 1.45-1.40(m, 3H), 1.08-1.01(m, 5H), 0.91-0.88(m, 3H)

Example 171: Preparation of (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile

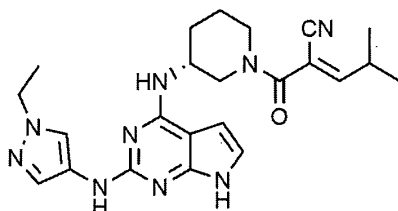


15 A title compound (8.9 mg, yield: 40.1%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde
20 instead of isobutyraldehyde in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.91(s, 1H), 7.54(s, 1H), 6.74(d, *J* = 3.5 Hz, 1H), 6.57-6.14(m, 2H), 4.27-4.24(m, 1H), 4.14-4.11(m, 2H), 4.05-3.59(m, 3H), 2.15-2.10(m, 1H), 2.03-

1.98(m, 2H), 1.82-1.59(m, 4H), 1.45-1.42(m, 3H), 1.51-1.32(m, 1H), 0.66-0.24(m, 2H)

Example 172: Preparation of (R,E)-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile



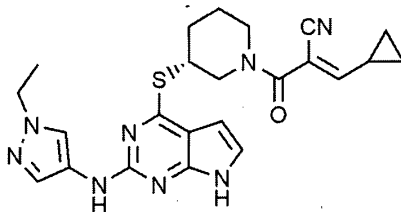
5

A title compound (4.3 mg, yield: 19.1%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.90(s, 1H), 7.53(s, 1H), 6.86-6.53(m, 2H), 6.41(d, *J* = 3.5 Hz, 1H), 4.69-4.64(m, 1H), 4.32-4.23(m, 1H), 4.15-4.11(m, 2H), 3.96-3.52(m, 3H), 2.14-2.11(m, 1H), 2.04-1.95(m, 2H), 1.86-1.58(m, 3H), 1.46-1.42(m, 3H), 1.18-1.13(m, 2H), 1.11-1.00(m, 2H), 0.74-0.72(m, 1H)

10

Example 173: Preparation of (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)acrylonitrile



15

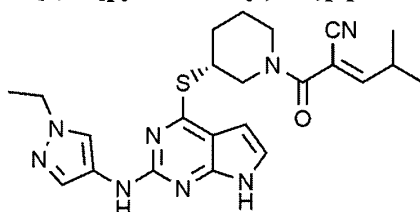
A title compound (10.8 mg, yield: 46.9%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

20

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.61(s, 1H), 6.92(d, *J* = 3.5 Hz, 1H), 6.54-6.02(m, 2H), 4.31-4.29(m, 1H), 4.15-4.12(m, 2H), 4.05-3.47(m, 3H), 2.25-2.18(m, 1H), 2.03-

1.57(m, 5H), 1.46-1.43(m, 3H), 1.16-1.01(m, 2H), 0.84-0.12(m, 2H)

Example 174: Preparation of (R,E)-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-4-methylpent-2-enenitrile



5

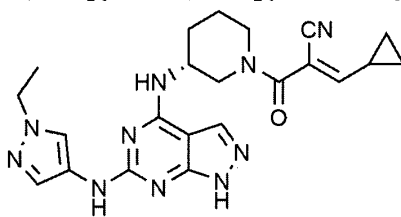
A title compound (10.8 mg, yield: 46.3%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 6.12(s, 1H), 7.00-6.90(m, 1H), 6.60-6.10(m, 2H), 4.40-4.20(m, 1H), 4.18-4.10(m, 2H), 3.90-3.70(m, 2H), 3.50-3.32(m, 1H), 2.30-2.15(m, 1H), 2.10-1.80(m, 3H), 1.79-1.50(m, 2H), 1.50-1.40(m, 3H), 1.20-0.90(m, 2H), 0.85-0.10(m, 2H)

10

15

Example 175: Preparation of (R,E)-3-cyclopropyl-2-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile



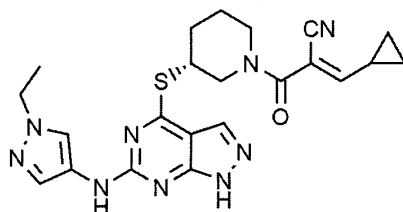
A title compound (7.3 mg, yield: 32.7%) was prepared in the same manner as in Example 160, except using 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

20

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.92(s, 1H), 7.57(s, 1H), 6.57-6.10(m, 1H), 4.27-4.24(m, 1H), 4.15-4.12(m, 2H), 4.02-3.44(m, 3H), 2.15-2.11(m, 1H), 2.04-1.97(m,

2H), 1.84-1.57(m, 4H), 1.46-1.43(m, 3H), 1.18-1.14 (m, 1H), 0.74-0.28(m, 2H)

Example 176: Preparation of (R,E)-3-cyclopropyl-2-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)acrylonitrile



5

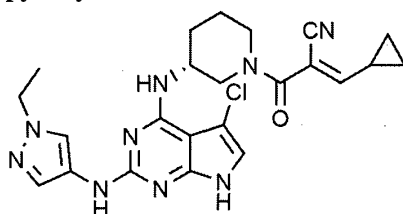
A title compound (11.1 mg, yield: 47.8%) was prepared in the same manner as in Example 160, except using 4,6-dichloro-1H-pyrazolo[3,4-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde in Example 160.

10

¹H NMR (500MHz, CD₃OD) δ 8.10-7.60(m, 3H), 6.40-6.00(m, 1H), 4.40-4.25(m, 2H), 4.20-4.10(m, 2H), 4.05-3.30(m, 3H), 2.30-2.20(m, 1H), 2.05-1.50(m, 4H), 1.49-1.40(m, 3H), 0.80-0.10(m, 4H)

15

Example 177: Preparation of (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile



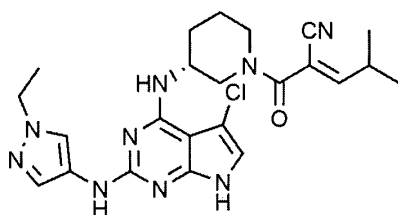
A title compound (8.5 mg, yield: 35.4%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde,

20

¹H NMR (500MHz, CD₃OD) δ 7.95-7.88(m, 1H), 7.60-7.50(m, 1H), 6.80-6.70(m,

1H), 6.50-6.00(m, 1H), 4.40-4.30(m, 1H), 4.35-4.00(m, 3H), 4.00-3.40(m, 2H), 2.15-1.50(m, 6H), 1.50-1.40(m, 3H), 1.20-0.95(m, 2H), 0.80-0.10(m, 2H)

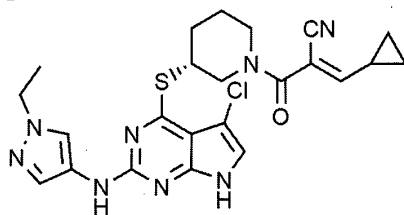
Example 178: Preparation of (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)-4-methylpent-2-enenitrile



A title compound (4.5 mg, yield: 18.8%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.95-7.88(m, 1H), 7.60-7.50(m, 1H), 6.90-6.50(m, 2H), 4.40-4.30(m, 1H), 4.25-4.00(m, 2H), 3.98-3.40(m, 3H), 2.70-2.50(m, 1H), 2.15-1.50(m, 5H), 1.48-1.40(m, 3H), 1.20-0.55(m, 6H)

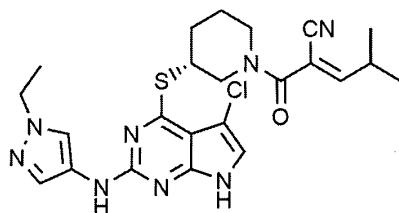
Example 179: Preparation of (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile



A title compound (6.4 mg, yield: 25.7%) was prepared in the same manner as in Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde instead of isobutyraldehyde.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.61(s, 1H), 6.93-6.88(m, 1H), 6.45-5.98(m, 1H), 4.40-4.30(m, 1H), 4.20-4.10(m, 2H), 4.04-3.60(m, 3H), 2.30-2.17(m, 1H), 2.05-1.50(m, 5H), 1.49-1.40(m, 3H), 1.20-0.93(m, 2H), 0.75-0.20(m, 2H)

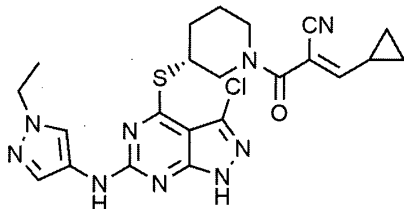
5 **Example 180: Preparation of (R,E)-2-(3-((5-chloro-2-((1-ethyl-1H-pyrazol-4-yl)amino)-7H-pyrrolo[2,3-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-4-methylpent-2-enenitrile**



A title compound (10.4 mg, yield: 41.7%) was prepared in the same manner as in
10 Example 160, except using 2,4-dichloro-7H-pyrrolo[2,3-d]pyrimidine instead of 3,4,6-trichloro-1H-pyrazolo[3,4-d]pyrimidine, and *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate in Example 160.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.61(s, 1H), 6.93-6.89(m, 1H), 6.70-6.48(m, 1H), 4.50-4.38(m, 1H), 4.20-4.10(m, 2H), 4.04-3.85(m, 2H), 3.50-3.35(m, 1H), 2.30-2.20(m, 1H), 2.08-1.70(m, 5H), 1.62-1.50(m, 1H), 1.49-1.40(m, 3H), 1.25-0.98(m, 3H), 0.75-0.55(m, 2H)

20 **Example 181: Preparation of (R,E)-2-(3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)thio)piperidine-1-carbonyl)-3-cyclopropylacrylonitrile**



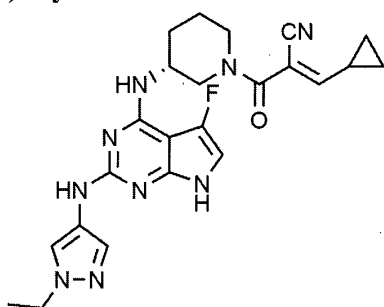
A title compound (7.6 mg, yield: 30.6%) was prepared in the same manner as in
Example 160, except using *tert*-butyl (R)-3-mercaptopiperidine-1-carboxylate instead of *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and cyclopropanecarbaldehyde

instead of isobutyraldehyde in Example 160.

¹H NMR (500MHz, CD₃OD) δ 8.05(s, 1H), 7.65(s, 1H), 6.55-6.20(m, 1H), 4.45-4.25(m, 2H), 4.20-4.10(m, 2H), 4.04-3.65(m, 2H), 2.30-2.20(m, 1H), 2.08-1.55(m, 8H), 1.20-0.90(m, 2H), 0.80-0.20(m, 2H)

5

Example 182: Preparation of (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile



Step 1: Preparation of (R)-3-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile

After 2-cyanoacetic acid (83.6 mg, 0.8 mmol) was dissolved in N,N-dimethylformamide (8.0 mL), 1-[bis(dimethylamino)methylene]-1H-1,2,3-triazolo[4,5-b]pyridinium 3-oxide hexafluorophosphate (374.0 mg, 1.2 mmol), N,N-diisopropylethylamine (206.4 μL, 2.4 mmol) and (R)-N2-(1-ethyl-1H-pyrazol-4-yl)-5-fluoro-N4-(piperidin-3-yl)-7H-pyrrolo[2,3-d]pyrimidin-2,4-diamine hydrochloride (Step 5 of Example 82) (341.2 mg, 0.8 mmol) were added thereto. The reaction mixture was stirred at room temperature for 24 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (240.0 mg, yield: 64.9%).

20

¹H NMR (500MHz, CD₃OD) δ 7.97-7.45(m, 3H), 4.30-4.20(m, 1H), 4.18-4.10(m, 2H), 3.90-3.79(m, 2H), 3.70-3.60(m, 3H), 2.18-1.75(m, 4H), 1.73-1.50(m, 2H), 1.45-1.38(m, 2H)

25

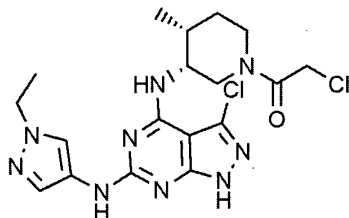
Step 2: Preparation of (R,E)-3-cyclopropyl-2-(3-((2-((1-ethyl-1H-pyrazol-4-

yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidine-1-carbonyl)acrylonitrile

After (R)-3-(3-((2-((1-ethyl-1H-pyrazol-4-yl)amino)-5-fluoro-7H-pyrrolo[2,3-d]pyrimidin-4-yl)amino)piperidin-1-yl)-3-oxopropanenitrile (50.0 mg, 0.12 mmol) was dissolved in methanol, piperidine (22.2 μ L, 0.16 mmol) and cyclopropanecarbaldehyde (135.4 mg, 0.16 mmol) were added thereto. The reaction mixture was stirred at room temperature for 5 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (9.6 mg, yield: 17.1%).

¹H NMR (500MHz, CD₃OD) δ 7.97-7.45(m, 3H), 6.55-6.25(m, 1H), 4.30-4.20(m, 1H), 4.18-4.10(m, 2H), 3.90-3.79(m, 2H), 3.70-3.60(m, 1H), 2.18-1.75(m, 4H), 1.73-1.50(m, 2H), 1.45-1.38(m, 3H), 1.20-1.00(m, 4H)

Example 183: Preparation of 2-chloro-1-((3R,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-methylpiperidin-1-yl)ethan-1-one



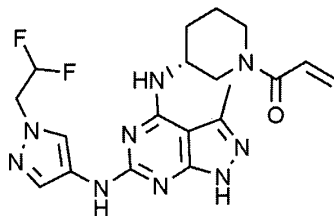
A title compound (4.7 mg, yield: 15.5%) was prepared in the same manner as in Example 138, except using *tert*-butyl (3R,4R)-3-amino-4-methylpiperidine-1-carboxylate instead of (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate, and 2-chloroacetic acid instead of 2-cyanoacetic acid in Example 138.

¹H NMR (500MHz, CD₃OD) δ 8.00(s, 1H), 7.59(s, 1H), 4.79-4.39(m, 3H), 4.23-4.12(m, 3H), 4.03-3.98(m, 2H), 3.16-2.92(m, 1H), 2.25-2.15(m, 1H), 1.89-1.77(m, 1H), 1.62-1.60(m, 1H), 1.46-1.44(m, 3H), 1.07-1.04(m, 3H)

25

Example 184: Preparation of (R)-1-(3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-

one



Step 1: Preparation of *tert*-butyl (R)-3-((6-chloro-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate

5 After 4,6-dichloro-3-methyl-1H-pyrazolo[3,4-d]pyrimidine (300.0 mg, 1.5 mmol) was dissolved in ethanol (10 mL), N,N-diisopropylethylamine (695.0 μ L, 2.2 mmol) and *tert*-butyl (R)-3-aminopiperidine-1-carboxylate (355.0 mg, 1.8 mmol) were added thereto. The reaction mixture was stirred at 110 °C for 12 hours, and then the organic layer was isolated, treated with magnesium sulfate, filtered and then concentrated under reduced pressure. The residue was
10 isolated by column chromatography to obtain a title compound (414.5 mg, yield: 76.3%).

^1H NMR (500MHz, CD_3OD) δ 4.59-4.50(m, 1H), 4.30-4.24(m, 1H), 3.93-3.89(m, 1H), 3.72-3.67(m, 1H), 3.19-3.17(m, 1H), 2.59(s, 3H), 2.10-2.00(m, 1H), 1.88-1.77(m, 2H), 1.63-1.30(m, 10H)

Step 2: Preparation of (R)-6-chloro-3-methyl-N- (piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidine-4-amine hydrochloride

To *tert*-butyl (R)-3-((6-chloro-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidine-1-carboxylate (414.5 Mg, 1.1 mmol) was added 6 N hydrochloric acid solution (2.0 mL, excessive amount) dissolved in methanol. After stirring at room temperature
20 for 30 minutes, the reaction mixture was concentrated and the subsequent reaction was carried out without isolation.

^1H NMR (500MHz, CD_3OD) δ 4.71-4.65(m, 2H), 3.61-3.59(m, 1H), 3.40-3.34(m, 1H), 3.07-2.97(m, 2H), 2.69-2.66(m, 2H), 2.17-2.09(m, 2H), 1.98-1.87(m, 2H)

Step 3: Preparation of (R)-1-(3-((6-chloro-3-methyl-7H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

After (R)-6-chloro-3-methyl-N-(piperidin-3-yl)-1H-pyrazolo[3,4-d]pyrimidine-4-

amine hydrochloride was dissolved in a 3:1 mixed solution of tetrahydrofuran: distilled water (4 mL), sodium bicarbonate (349.9 mg, 4.2 mmol) was added thereto at -20 °C, and then stirred for 30 minutes. Acryloyl chloride (131.4 μ L, 1.5 mmol) was slowly added dropwise to the reaction mixture, and then stirred at -20 °C for 1 hour. The organic layer was isolated, treated with
 5 magnesium sulfate, filtered, and then concentrated under reduced pressure. The residue was isolated by column chromatography to obtain a title compound (265.4 mg, yield: 56.4%).

^1H NMR (500MHz, CD_3OD) δ 6.91-6.79(m, 1H), 6.25-6.17(m, 1H), 5.79-5.70(m, 1H), 4.37-4.30(m, 2H), 4.18-4.15(m, 1H), 3.54-3.48(m, 1H), 3.20-2.96(m, 1H), 2.58(s, 3H), 2.15-2.13(m, 1H), 2.10-1.82(m, 2H), 1.70-1.65(m, 1H)

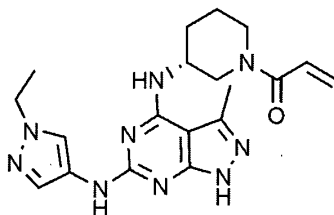
10

Step 4: Preparation of (R)-1-(3-(((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

(R)-1-(3-((6-chloro-3-methyl-7H-pyrazolo[3,4-d] pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one (30.0 mg, 0.09 mmol) and 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine (10.6 mg, 0.07 mmol) were dissolved in 2-butanol (2.0 mL). Trifluoroacetic acid (6.6 μ L, 0.09 mmol) was added to the reaction mixture, followed by reacting at 120 °C for 3 hours, and then the solvent was concentrated. The reaction mixture was neutralized by adding 7 N ammonia solution dissolved in methanol, and the residue was isolated by column chromatography to
 20 obtain a title compound (16.7 mg, yield: 55.3%).

^1H NMR (500MHz, CD_3OD) δ 8.05(d, 1H), 7.61(d, 1H), 6.87-6.53(m, 1H), 6.28-6.11(m, 2H), 6.08-5.54(m, 1H), 4.58-4.32(m, 4H), 4.05-3.95(m, 2H), 3.50-3.12(m, 1H), 2.5(s, 3H), 2.15-2.11(m, 1H), 1.95-1.86(m, 2H), 1.65-1.63(m, 1H)

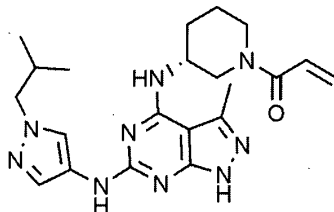
Example 185: Preparation of (R)-1-(3-(((1-ethyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (21.9 mg, yield: 79.3%) was prepared in the same manner as in Example 184, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine in Example 184.

¹H NMR (500MHz, CD₃OD) δ 7.95(d, 1H), 7.54(d, 1H), 6.87-6.55(m, 1H), 6.28-6.08(m, 1H), 5.81-5.54(m, 1H), 4.51-3.92(m, 5H), 3.50-3.18(m, 2H), 2.49(s, 3H), 2.20-2.10(m, 1H), 1.96-1.89(m, 2H), 1.64-1.62(m, 1H)

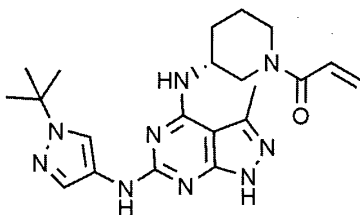
Example 186: Preparation of (R)-1-(3-((6-((1-isobutyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (19.4 mg, yield: 65.5%) was prepared in the same manner as in Example 184, except that 1-isobutyl-1H-pyrazol-4-amine was used instead of 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine in Example 184.

¹H NMR (500MHz, CD₃OD) δ 7.93(d, 1H), 7.55(d, 1H), 6.87-6.55(m, 1H), 6.29-6.09(m, 1H), 5.81-5.56(m, 1H), 4.52-3.93(m, 3H), 3.87-3.85(m, 2H), 3.50-3.15(m, 2H), 2.49(s, 3H), 2.16-2.08(m, 2H), 1.93-1.87(m, 2H), 1.63-1.61(m, 1H), 0.90(s, 6H)

Example 187: Preparation of (R)-1-(3-((6-((1-tert-butyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

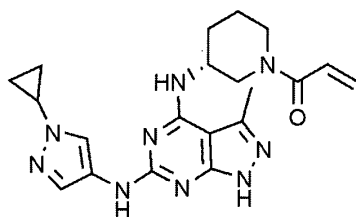


A title compound (17.8 mg, yield: 60.1%) was prepared in the same manner as in Example 184, except that 1-(tert-butyl)-1H-pyrazol-4-amine was used instead of 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine in Example 184.

¹H NMR (500MHz, CD₃OD) δ 8.04(s, 1H), 7.59(s, 1H), 6.85-6.58(m, 1H), 6.27-

6.06(m, 1H), 5.80-5.54(m, 1H), 4.36-4.30(m, 1H), 4.06-4.04(m, 1H), 3.90-3.87(m, 1H), 3.45-3.22(m, 2H), 2.49(s, 3H), 2.12-2.09(m, 1H), 2.00-1.85(m, 2H), 1.63-1.50(m, 10H)

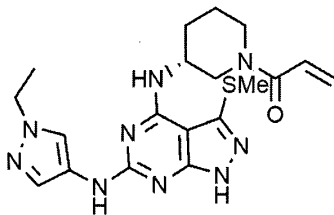
Example 188: Preparation of (R)-1-(3-((6-((1-cyclopropyl-1H-pyrazol-4-yl)amino)-3-methyl-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (21.9 mg, yield: 79.3%) was prepared in the same manner as in Example 184, except that 1-cyclopropyl-1H-pyrazol-4-amine was used instead of 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine in Example 184.

¹H NMR (500MHz, CD₃OD) δ 7.99(s, 1H), 7.51(s, 1H), 6.87-6.55(m, 1H), 6.27-6.08(m, 1H), 5.80-5.54(m, 1H), 4.48-3.91(m, 3H), 3.57-3.56(m, 1H), 3.50-3.22(m, 3H), 2.50(s, 3H), 2.20-2.11(m, 1H), 1.97-1.87(m, 2H), 1.65-1.63(m, 1H), 1.04-1.00(m, 4H)

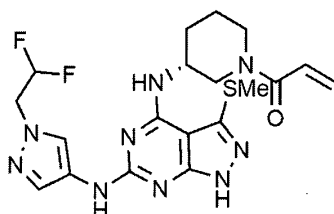
Example 189: Preparation of (R)-1-(3-((6-((1-ethyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (20.4 mg, yield: 79.6%) was prepared in the same manner as in Example 129, except that 1-ethyl-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 7.97(s, 1H), 7.56(d, 1H), 6.90-6.50(m, 1H), 6.26-6.08(m, 1H), 5.80-5.55(m, 1H), 4.38-4.30(m, 1H), 4.15-4.07(m, 2H), 3.95-3.79(m, 2H), 3.59-3.50(m, 1H), 2.51(s, 3H), 2.15-2.08(m, 1H), 1.94-1.85(m, 2H), 1.75-1.69(m, 1H), 1.44-1.40(m, 3H)

Example 190: Preparation of (R)-1-(3-((6-((1-(2,2-difluoroethyl)-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



5

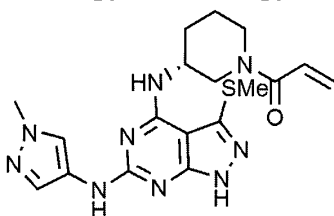
A title compound (16.1 mg, yield: 57.9%) was prepared in the same manner as in Example 129, except that 1-(2,2-difluoroethyl)-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 8.06(d, 1H), 7.63(d, 1H), 6.85-6.50(m, 1H), 6.26-6.04(m, 2H), 5.79-5.52(m, 1H), 4.52-4.47(m, 2H), 4.38-4.33(m, 2H), 3.96-3.90(m, 1H), 3.79-3.77(m, 1H), 3.55-3.48(m, 1H), 2.52(s, 3H), 2.09-2.07(m, 1H), 1.93-1.81(m, 2H), 1.75-1.70(m, 1H)

10

Example 191: Preparation of (R)-1-(3-((6-((1-methyl-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

15



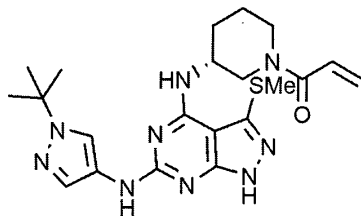
A title compound (10.6 mg, yield: 42.7%) was prepared in the same manner as in Example 129, except that 1-methyl-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

20

¹H NMR (500MHz, CD₃OD) δ 8.54(s, 1H), 7.92(s, 1H), 6.86-6.52(m, 1H), 6.25-6.04(m, 1H), 5.79-5.49(m, 1H), 4.36-4.19(m, 1H), 3.84-3.75(m, 4H), 3.60-3.54(m, 2H), 2.51(s, 3H), 2.15-2.05(m, 1H), 1.95-1.84(m, 2H), 1.75-1.70(m, 1H)

Example 192: Preparation of (R)-1-(3-((6-((1-(*tert*-butyl)-1H-pyrazol-4-yl)amino)-3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one

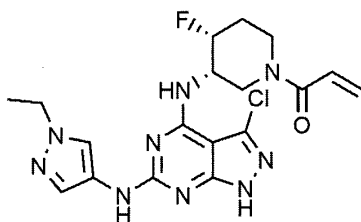
3-(methylthio)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)piperidin-1-yl)prop-2-en-1-one



A title compound (16.8 mg, yield: 61.5%) was prepared in the same manner as in Example 129, except that 1-(*tert*-butyl)-1H-pyrazol-4-amine was used instead of 1-cyclopropyl-1H-pyrazol-4-amine in Example 129.

¹H NMR (500MHz, CD₃OD) δ 8.06(s, 1H), 7.61(s, 1H), 6.85-6.55(m, 1H), 6.24-6.04(m, 1H), 5.79-5.50(m, 1H), 4.37-4.35(m, 1H), 3.88-3.48(m, 4H), 2.51(s, 3H), 2.09-2.00(m, 1H), 1.93-1.84(m, 2H), 1.70-1.65(m, 1H), 1.57(s, 9H)

Example 193: Preparation of 1-((3S,4R)-3-((3-chloro-6-((1-ethyl-1H-pyrazol-4-yl)amino)-1H-pyrazolo[3,4-d]pyrimidin-4-yl)amino)-4-fluoropiperidin-1-yl)prop-2-en-1-one



A title compound (8.2 mg, yield: 37.8%) was prepared in the same manner as in Example 46, except that *tert*-butyl (3S,4R)-3-amino-4-fluoropiperidine-1-carboxylate was used instead of *tert*-butyl (R)-3-aminopiperidine-1-carboxylate in Example 46.

¹H NMR (500MHz, CD₃OD) δ 7.99(s, 1H), 7.57(s, 1H), 6.85-6.70(m, 1H), 6.31-6.27(m, 1H), 5.83-5.65(m, 1H), 5.15-5.05(m, 1H), 4.58-4.25(m, 1H), 4.13-4.03(m, 3H), 3.60-3.43(m, 1H), 3.34-3.05(m, 2H), 2.22-2.15(m, 1H), 2.01-1.90(m, 2H), 1.43-1.42(m, 3H)

Experimental Example 1: Measurement of JAK 3 and BTK Enzyme inhibitory activity

JAK3 and BTK kinase inhibitory activities were measured for the compounds

prepared in the above Examples through in vitro analysis on the ADP Glo (Glo) platform.

Specifically, the inhibitory activities of JAK3 and BTK kinase were measured using a JAK3 kinase assay kit (Promega, V9441) and a BTK kinase assay kit (Promega, V9071) which were purchased from Promega. Recombinant purified human JAK3 and BTK were diluted with 1 x kinase reaction buffer (JAK3: 40 mM Tris-Cl, pH 7.5, 20 mM MgCl₂, 0.1 mg/mL BSA and 50 uM DTT / BTK: 40 mM Tris-Cl, pH 7.5, 20 mM MgCl₂, 0.1 mg/mL BSA, 2 mM MnCl₂ and 50 uM DTT) and added to a 96 well plate (JAK3: final concentration of 4 ng per reaction/BTK: final concentration of 8 ng per reaction). The compounds were treated so as to be finally a 1% DMSO aqueous solution, and a substrate cocktail containing ATP (JAK3: final concentration of 5 uM / BTK: final concentration of 10 uM) and 0.2 µg/µL of Poly(Glu4, Tyr1) peptide (JAK3 and BTK final concentration) in the total 25 µL reactants was added to a 96-well plate to initiate enzymatic reaction. After incubation (30 °C) for 1 hour, equivalent volume (25 µL per reaction) of ADP Glo was added and incubated (30 °C) for 40 minutes at room temperature. Then, a kinase detection reagent (50 µL per reaction) was added and incubated (30 °C) for 30 minutes at room temperature. The kinase activity was measured by chemiluminescence according to the instructions of ADP Glo kinase assay kit, and the inhibitory activity of the compounds according to the present invention was calculated. For the analysis of the results of each compound, Microsoft Excel was used, and IC₅₀ values were calculated by SigmaPlot software. The results are shown in Tables 1 to 5 below.

[Table 1]

Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)	Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)
1	3.73	10.7	21	0.4	4.5
2	8.9	27	22	17.3% inhibition@100nM	8.9% inhibition@100nM
3	28.2% inhibition @100nM	30.0% inhibition @100nM	23	0.9	13.9
4	6.9	26.7	24	0.4	15.3
5	24.6	76.7	25	11.2	39.3% inhibition@100nM
6	4.6	9.6	26	39.2	10.3% inhibition@100nM
7	5	14	27	32.8% inhibition@1uM	48.0% inhibition@1uM

8	1.3	14.3	28	185.5	3.3% inhibition@1uM
9	3.9	20.5	29	66.2	47.1% inhibition@1uM
10	32.8% inhibition@100nM	23.7% inhibition@100nM	30	52.6% inhibition@100nM	6
11	36.0% inhibition@100nM	26.5% inhibition@100nM	31	3.4	2.2
12	1.3	10.6	32	10.7	2.8
13	0.4% inhibition@1uM	1% inhibition@1uM	33	4.9	1.6
14	1.1	19.2	34	3.4	2.1
15	0.3	7.2	35	5.2	2.1
16	0.4	10.1	36	1.1	16.5
17	0.6	11.8	37	0.1	2.5
18	0.3	8.8	38	0.1	4.3
19	0.5	8.7	39	0.1	1.9
20	0.7	9.5	40	0.2	1.8

[Table 2]

Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)	Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)
41	0.5	1.8	61	1.1	1.7
42	5.1	3.2	62	0.2	1.7
43	9.9	6.7	63	0.3	1.7
44	0.3	2.5	64	0.7	1.7
45	0.7	2.9	65	5.5	1.7
46	0.6	2	66	3.7	1.7
47	1.1	5.2	67	5.7	1.7
48	0.4	0.9	68	0.7	12.1
49	0.7	0.9	69	0.3	8.6
50	0.8	1.8	70	0.2	2.3
51	0.8	1.8	71	2.8	1.3
52	0.2	0.8	72	2.2	1.1
53	1	0.9	73	4.4	1.1
54	0.7	1.8	74	5.4	1.7
55	1.3	1.2	75	0.2	4.1
56	0.3	1.8	76	0.3	1.0
57	0.4	1.8	77	6.0	1.1
58	0.5	1.8	78	0.7	11.5
59	1.4	1.8	79	0.4	9.2
60	0.2	1.7	80	8.4	33.3

[Table 3]

Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)	Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)
81	3.8	25.5	103	1.3	21.0
82	0.3	0.9	104	0.3	2.7
83	1.2	3.4	105	3.6	1.9

84	3.8	6.6	106	54.8	1.8
85	1.5	2.2	107	3.7	7.8
86	0.4	1.0	108	0.6	1.3
87	16.4	42.2	109	0.3	2.0
88	0.2	1.1	110	1.7	1.4
89	0.138	0.864	111	1.1	1.3
90	0.5	1.0	112	1.6	1.8
91	1.1	1.1	113	2.8	2.3
92	4.5	2.1	114	0.9	2.3
93	2.0	1.3	115	1.7	2.0
94	4.5	1.7	116	19.5	25.3
95	0.5	5.9	117	> 400	81.2
96	0.7	24.0	118	3.2	1.3
97	> 80	707.7	119	32.5	11.1
98	2.4	14.1	120	7.8	7.5
101	1.8	7.4	121	18.0	4.9
102	13.2	146.1	122	8.1	6.0

[Table 4]

Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)	Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)
123	7.9	5.8	143	19.3	165.1
124	8.4	3.5	144	1.4	1.5
125	20.4	8.2	145	1.7	1.4
126	2.9	1.2	146	7.4	1.4
127	43.1	1.5	147	1.4	2.1
128	14.6	5.8	148	0.3	1.6
129	42.3	2.3	149	0.2	1.6
130	9.0	1.1	150	0.3	1.6
131	24.5	1.6	151	0.4	3.0
132	25.5	2.1	152	2.0	4.5
133	16	1.6	153	1.6	2.2
134	17.7	1.8	154	54.2	2.5
135	42.3	2.3	155	53.7	2.2
136	0.1	1.1	156	0.5	1.1
137	2.6	19.6	157	0.6	1.4
138	> 80	92.7	158	2.9	1.2
139	> 80	142.8	159	27.1	4.5
140	> 80	985.9	160	1.5	3.4
141	38.5	306.5	161	5.0	1.3
142	22.4	210.1	162	3.0	4.8

[Table 5]

Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)	Ex. No.	JAK3 IC ₅₀ (nM)	BTK IC ₅₀ (nM)
163	2.7	1.2	179	1.1	8.3
164	8.8	2.2	180	1.5	3.1
165	2.2	1.9	181	30.2	2.2
166	5.6	1.2	182	2.3	2.1
167	43.1	1.5	183	0.8	1.6
168	18.0	4.3	184	4.4	1.4

169	240.9	2.2	185	15.1	2.1
170	11.4	2.9	186	18.0	1.6
171	0.9	2.2	187	14.6	1.8
172	0.9	1.6	188	23.5	2.9
173	1.0	6.0	189	3.4	0.721
174	0.7	2.0	190	2.0	0.760
175	14.9	1.4	191	4.2	0.760
176	42.1	3.0	192	3.0	0.726
177	0.4	1.2	193	0.5	0.9
178	0.5	0.9			