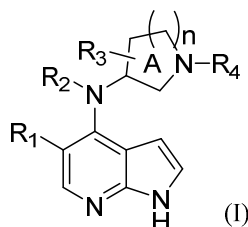


CLAIMS

What is claimed is:

1. A compound, or a pharmaceutically acceptable salt, hydrate or solvate thereof, of Formula (I):



wherein:

R₁ is selected from -CO₂R₅, -C₁-C₅alkyl-CO₂R₅, -C₃-C₆-cycloalkyl-CO₂R₅, -NHCO₂R₅, -N(C₁-C₅ alkyl)-CO₂R₅, -O-CO₂R₅, or -C₁-C₅alkyl-O-CO₂R₅;

R₂ is selected from H, -C₁-C₄alkyl, -C₃-C₆cycloalkyl, or -C₁-C₂alkyl-C₃-C₆cycloalkyl, wherein the alkyl or cycloalkyl groups are optionally substituted with one or more groups selected from halogen, -OH, or -O-C₁-C₅alkyl;

n is 0, 1 or 2;

Ring A is substituted at one or more carbons with one, two, or three R₃ substituents wherein each R₃ group is independently selected from H, halogen, -C₁-C₄alkyl, -C₃-C₆cycloalkyl, -OH, or -O-C₁-C₅alkyl wherein each alkyl or cycloalkyl group is optionally substituted with one or more groups selected from: halogen, -OH, -C₁-C₅alkylalkoxy, or -O-C₁-C₅alkyl;

Two R₃ groups on the same or different carbon atoms of the ring A may be optionally joined to form a spirocyclic or bicyclic ring system with ring A;

R₄ is selected from -C(O)-R₆, -CH₂R₆, -C(O)-CH=CH₂, -C(O)-C(CH₂OCH₃)=CH₂, -C(O)-CH=CHCH₃, -C(O)-CH=CHCH₂NR₇R₈, -C(O)-C₁-C₅alkyl, or -C(O)-C₃-C₆cycloalkyl, wherein the alkyl or cycloalkyl groups may be optionally substituted with one or more groups selected from -OH, halogen, alkyne, or -CN;

R₅ is selected from -C₁-C₅alkyl, or -C₃-C₆cycloalkyl wherein the alkyl or cycloalkyl groups may be optionally substituted by one or more groups selected from halogen, -OH, or -O-C₁-C₅alkyl;

R₆ is selected from -C₁-C₅alkyl, -C₃-C₆cycloalkyl, -C₁-C₅alkyl-C₃-C₆cycloalkyl, -NR₇R₈, -O-aryl, -O-heteroaryl, or heteroaryl wherein the alkyl, cycloalkyl, or heteroaryl groups can be optionally substituted by one or more groups selected from halogen, -CN, alkyne, -OH, trifluoromethyl, -O-C₁-C₅alkyl, or -O-C₃-C₆cycloalkyl;

R₇ and R₈ are independently selected from H, -C₁-C₅ alkyl, -C₁-C₅ alkoxy, or -C₃-C₅ cycloalkyl wherein the alkyl groups may be optionally substituted by one or more groups selected from halogen, -OH, or -CN; and

R₇ and R₈ may be optionally joined to form a ring to form a heterocycle or with another heteroatom to form a ring.

2. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -CO₂R₅.

3. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -C₁-C₅-alkyl-CO₂R₅.

4. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -C₃-C₆-cycloalkyl-CO₂R₅.

5. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -NHCO₂R₅.

6. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -N(C₁-C₅ alkyl)-CO₂R₅.

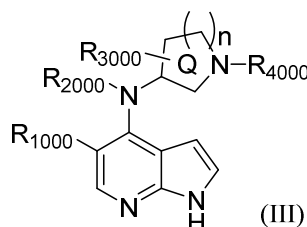
7. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -O-CO₂R₅.

8. The compound of Claim 1, or a pharmaceutically acceptable salt, hydrate or solvate thereof, wherein:

R₁ is -C₁-C₅alkyl-O-CO₂R₅.

9. A compound, or a pharmaceutically acceptable salt, hydrate or solvate thereof, of Formula (III):



wherein:

R₁₀₀₀ is selected from -CO₂R₅₀₀₀, -C₁-C₅alkyl-CO₂R₅₀₀₀, -C₃-C₆-cycloalkyl-CO₂R₅₀₀₀, -NHCO₂R₅₀₀₀, -N(C₁-C₅ alkyl)-CO₂R₅₀₀₀, -O-CO₂R₅₀₀₀, or -C₁-C₅alkyl-O-CO₂R₅₀₀₀;

R₂₀₀₀ is selected from H, -C₁-C₄alkyl, -C₃-C₆cycloalkyl, or -C₁-C₂alkyl-C₃-C₆cycloalkyl, wherein the alkyl or cycloalkyl groups are optionally substituted with one or more groups selected from halogen, -OH, or -O-C₁-C₅alkyl;

n is 0, 1 or 2;

Ring Q is substituted at one or more carbons with one, two, or three R₃₀₀₀ substituents wherein each R₃₀₀₀ group is independently selected from H, halogen, -C₁-C₄alkyl, -C₃-C₆cycloalkyl, -OH, or -O-C₁-C₅alkyl wherein each alkyl or cycloalkyl group is optionally substituted with one or more groups selected from: halogen, -OH, -C₁-C₅alkylalkoxy, or -O-C₁-C₅alkyl;

Two R₃₀₀₀ groups on the same or different carbon atoms of the ring Q may be optionally joined to form a spirocyclic or bicyclic ring system with ring Q;

R₄₀₀₀ is selected from -C(O)-R₆₀₀₀, -CH₂R₆₀₀₀, -C(O)-CH=CH₂, -C(O)-C(CH₂OCH₃)=CH₂, -C(O)-CH=CHCH₃, -C(O)-CH=CHCH₂NR₇₀₀₀R₈₀₀₀, -C(O)-C₁-C₅alkyl, or -C(O)-C₃-C₆cycloalkyl, wherein the alkyl or cycloalkyl groups may be optionally substituted with one or more groups selected from -OH, halogen, alkyne, or -CN;

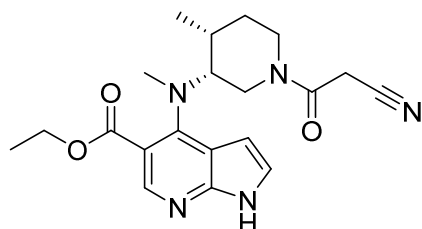
R₅₀₀₀ is selected from H, -C₁-C₅alkyl, and -C₃-C₆cycloalkyl wherein the alkyl or cycloalkyl groups may be optionally substituted by one or more groups selected from halogen, -OH, -C₃-C₅ cycloalkyl, or -O-C₁-C₅alkyl;

R₆₀₀₀ is selected from -C₁-C₅alkyl, -C₃-C₆cycloalkyl, -C₁-C₅alkyl-C₃-C₆cycloalkyl, -NR₇₀₀₀R₈₀₀₀, -O-aryl, -O-heteroaryl, or heteroaryl wherein the alkyl, cycloalkyl, or heteroaryl groups can be optionally substituted by one or more groups selected from halogen, -CN, alkyne, -OH, trifluoromethyl, -O-C₁-C₅alkyl, or -O-C₃-C₆cycloalkyl;

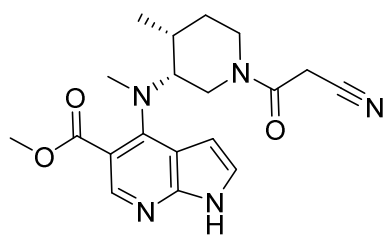
R₇₀₀₀ and R₈₀₀₀ are independently selected from H, -C₁-C₅ alkyl, -C₁-C₅ alkoxy, or -C₃-C₅ cycloalkyl where the alkyl groups may be optionally substituted by one or more groups selected from halogen, -OH, -CN, or -C₃-C₅ cycloalkyl;

R₇₀₀₀ and R₈₀₀₀ may be optionally joined to form a ring to form a heterocycle or with another heteroatom to form a ring.

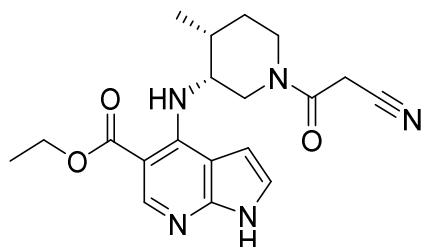
10. The compound of Claim 1, wherein the compound is selected from the group consisting of:



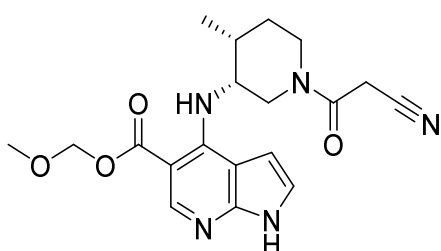
ethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



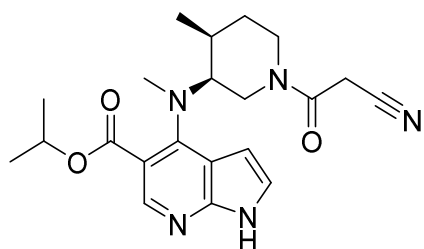
methyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



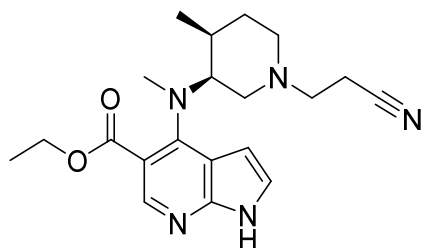
ethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



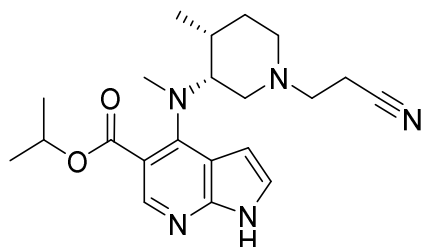
methoxymethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



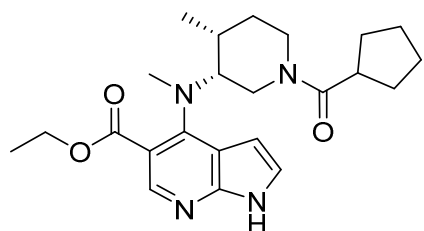
isopropyl 4-(((3*S*,4*S*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



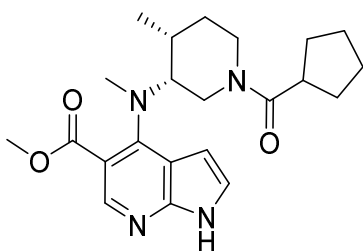
ethyl 4-(((3*S*,4*S*)-1-(2-cyanoethyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



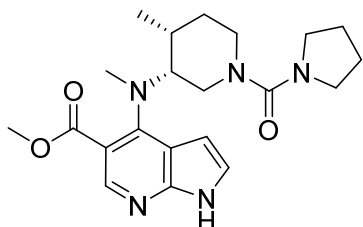
isopropyl 4-(((3*R*,4*R*)-1-(2-cyanoethyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



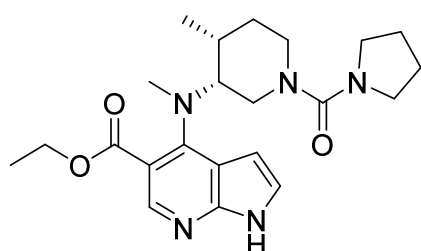
ethyl 4-(((3*R*,4*R*)-1-(cyclopentanecarbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



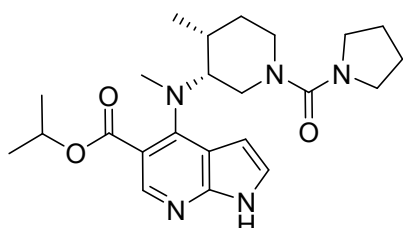
methyl 4-(((3R,4R)-1-(cyclopentanecarbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



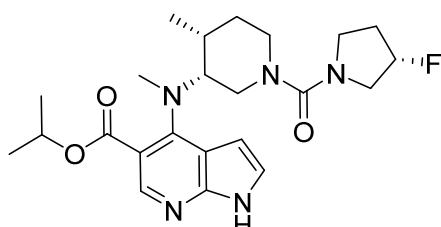
methyl 4-(methyl((3R,4R)-4-methyl-1-(pyrrolidine-1-carbonyl)piperidin-3-yl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



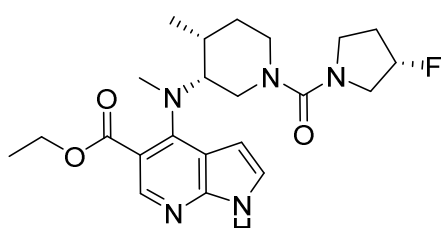
ethyl 4-(methyl((3R,4R)-4-methyl-1-(pyrrolidine-1-carbonyl)piperidin-3-yl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



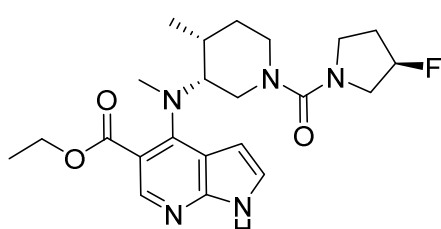
isopropyl 4-(methyl((3R,4R)-4-methyl-1-(pyrrolidine-1-carbonyl)piperidin-3-yl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



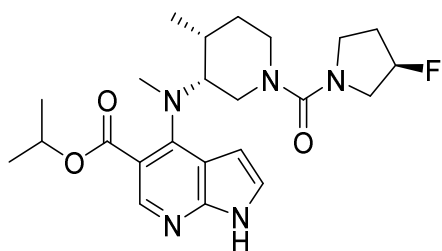
isopropyl 4-(((3R,4R)-1-((S)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



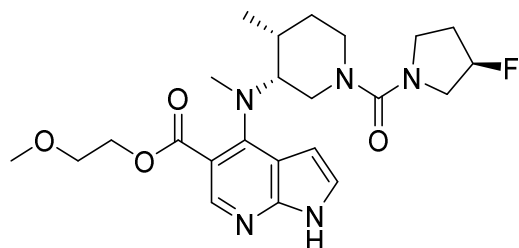
ethyl 4-(((3R,4R)-1-((S)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



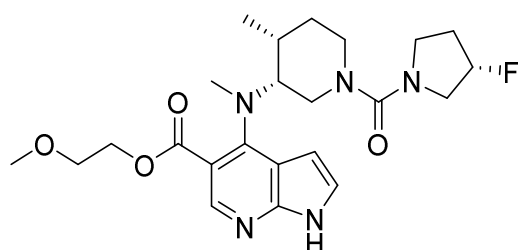
ethyl 4-(((3R,4R)-1-((R)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1H-pyrrolo[2,3-b]pyridine-5-carboxylate,



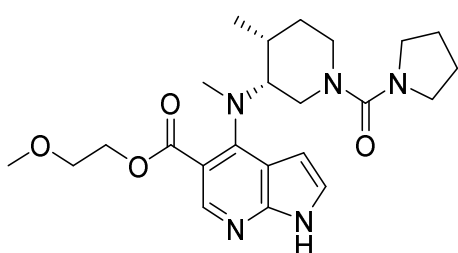
isopropyl 4-(((3*R*,4*R*)-1-((*R*)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



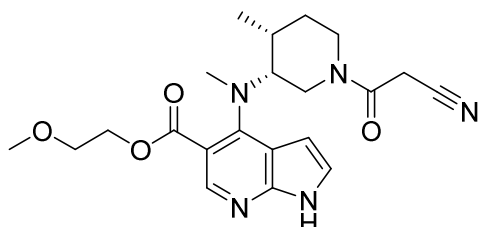
2-methoxyethyl 4-(((3*R*,4*R*)-1-((*R*)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



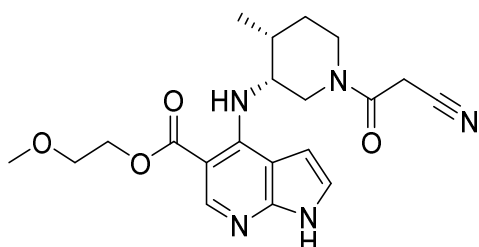
2-methoxyethyl 4-(((3*R*,4*R*)-1-((*S*)-3-fluoropyrrolidine-1-carbonyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



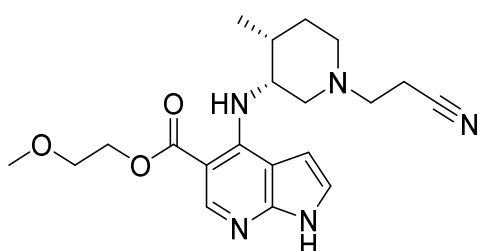
2-methoxyethyl 4-(methyl((3*R*,4*R*)-4-methyl-1-(pyrrolidine-1-carbonyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



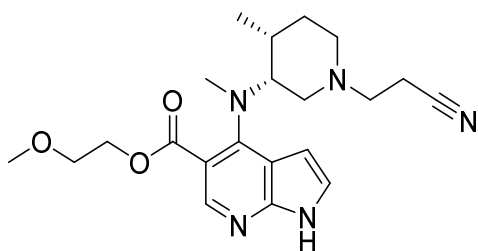
2-methoxyethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



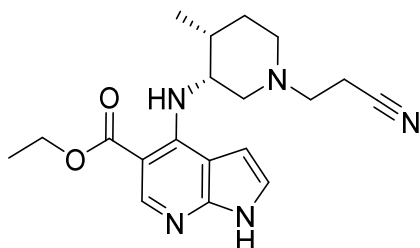
2-methoxyethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



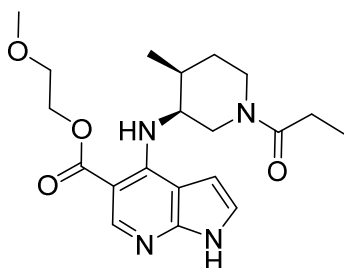
2-methoxyethyl 4-(((3*R*,4*R*)-1-(2-cyanoethyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



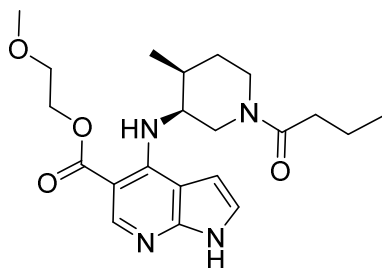
2-methoxyethyl 4-(((3*R*,4*R*)-1-(2-cyanoethyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



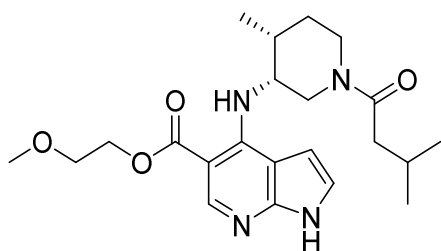
ethyl 4-(((3*R*,4*R*)-1-(2-cyanoethyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



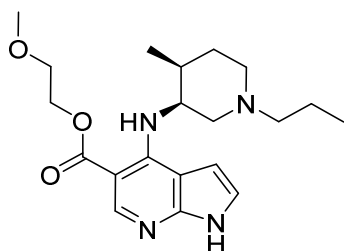
2-methoxyethyl 4-(((3*S*,4*S*)-4-methyl-1-propionylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



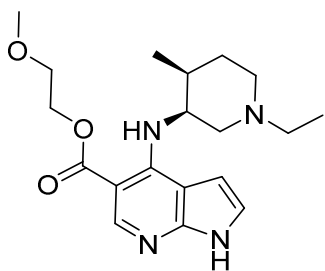
2-methoxyethyl 4-(((3*S*,4*S*)-1-butyryl-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



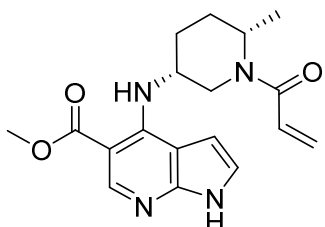
2-methoxyethyl 4-(((3*R*,4*R*)-4-methyl-1-(3-methylbutanoyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



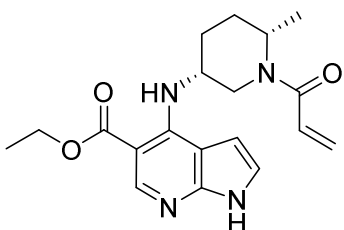
2-methoxyethyl 4-(((3*S*,4*S*)-4-methyl-1-propylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



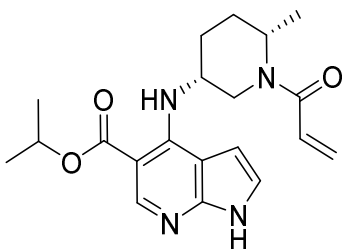
2-methoxyethyl 4-(((3*S*,4*S*)-1-ethyl-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



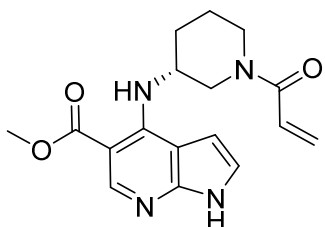
methyl 4-(((3*R*,6*S*)-1-acryloyl-6-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



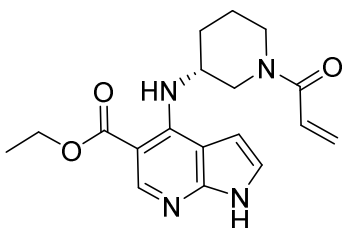
ethyl 4-(((3*R*,6*S*)-1-acryloyl-6-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



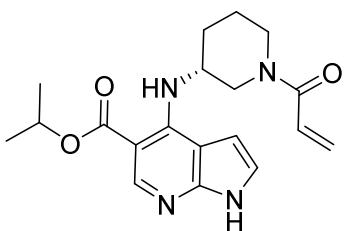
isopropyl 4-(((3*R*,6*S*)-1-acryloyl-6-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



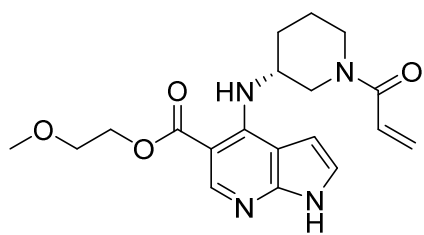
methyl (*R*)-4-((1-acryloylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



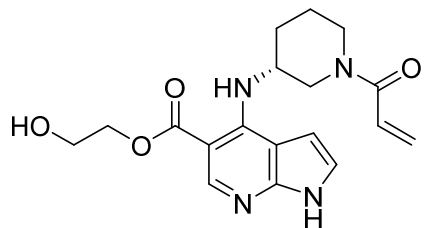
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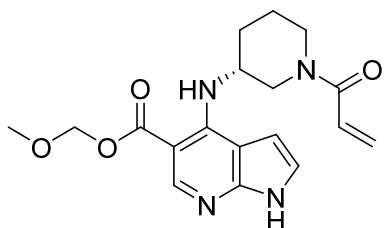
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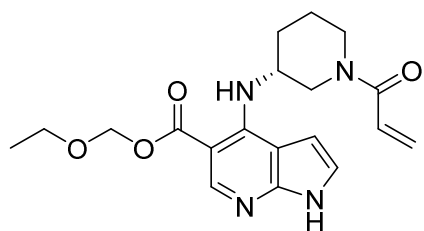
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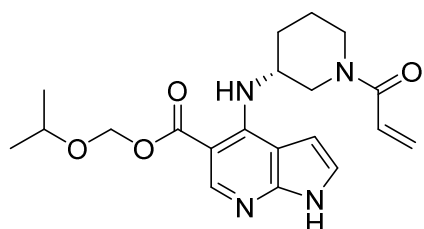
2-hydroxyethyl (*R*)-4-((1-acryloylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



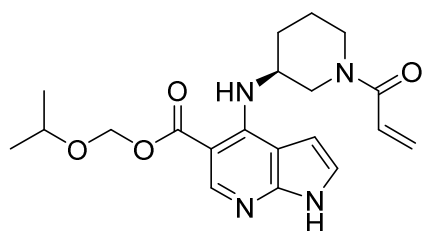
methoxymethyl (*R*)-4-((1-acryloylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



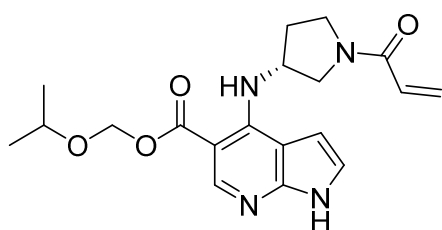
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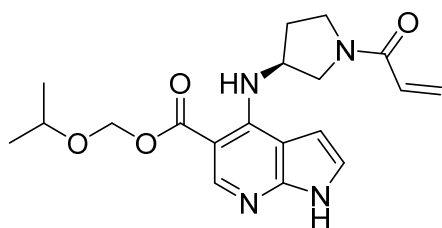
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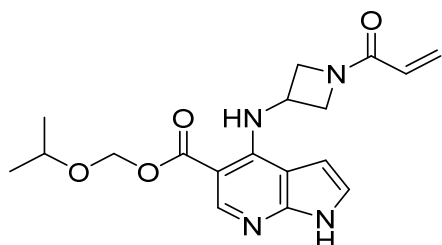
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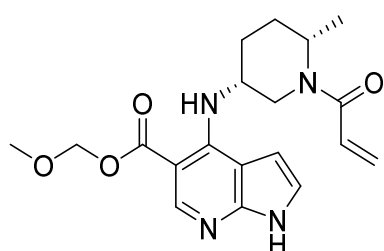
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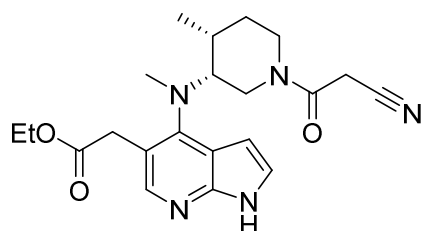
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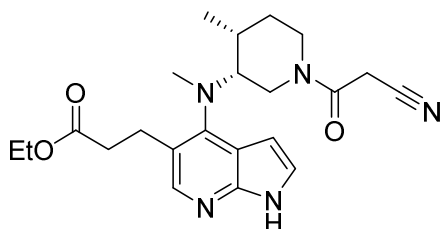
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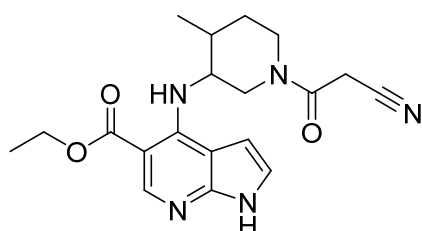
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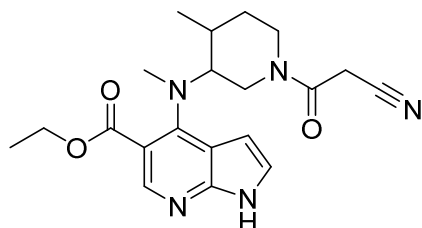
ethyl 2-(4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridin-5-yl)acetate,



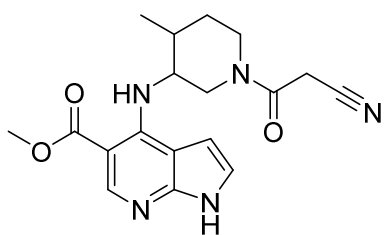
ethyl 3-(4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)(methyl)amino)-1*H*-pyrrolo[2,3-*b*]pyridin-5-yl)propanoate



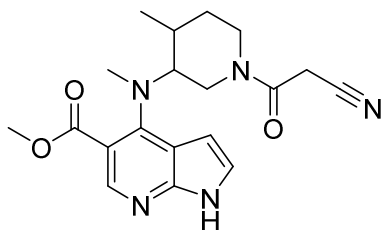
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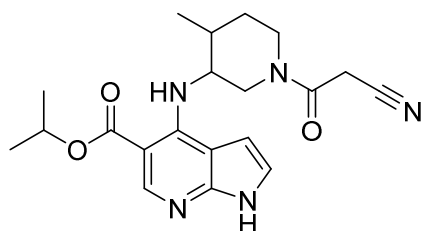
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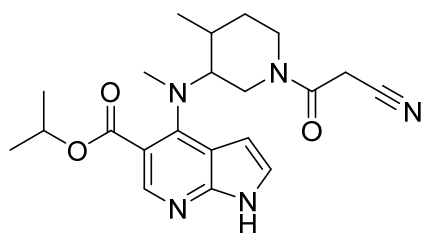
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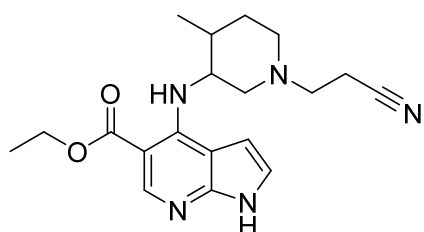
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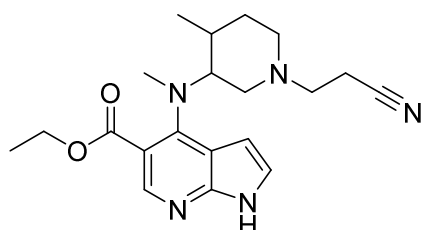
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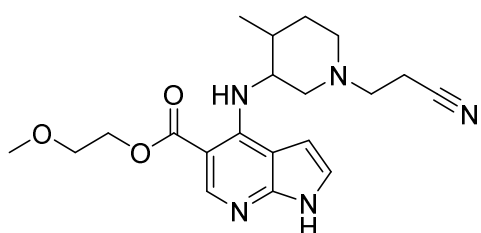
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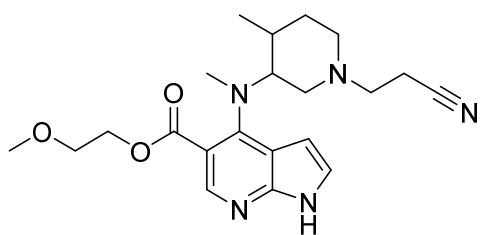
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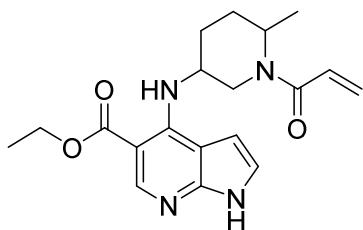
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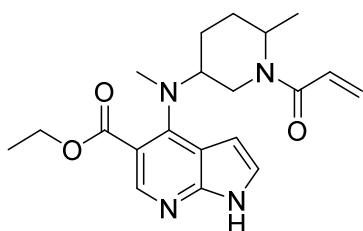
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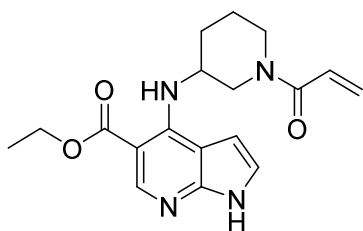
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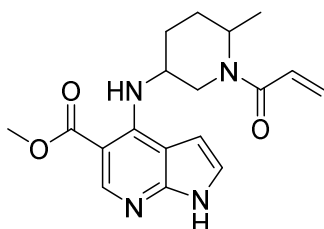
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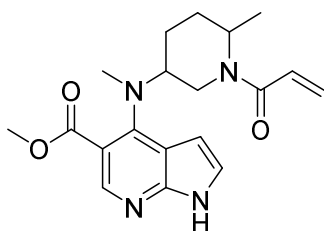
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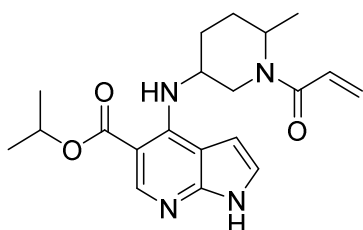
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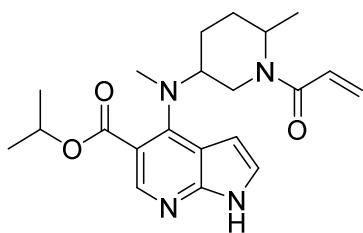
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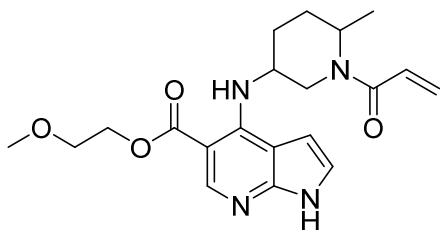
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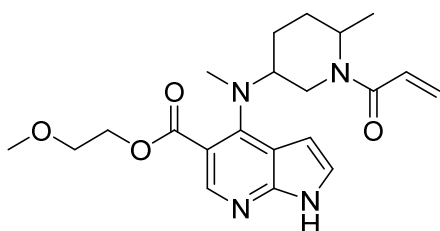
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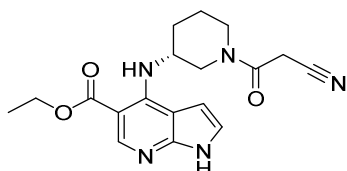
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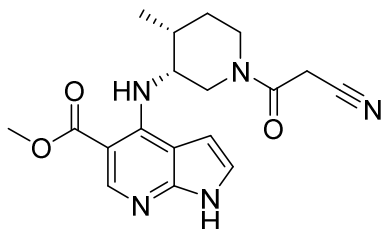
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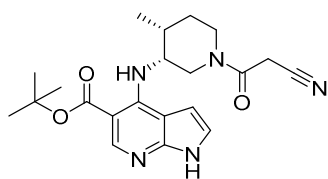
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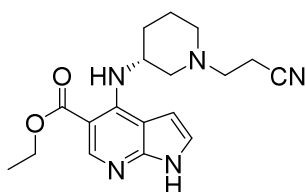
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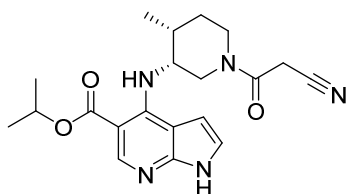
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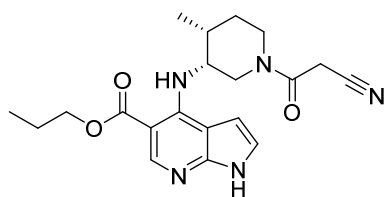
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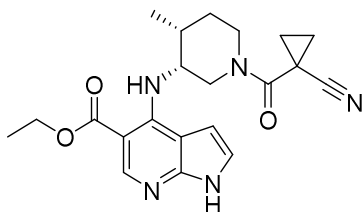
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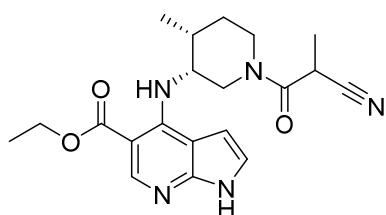
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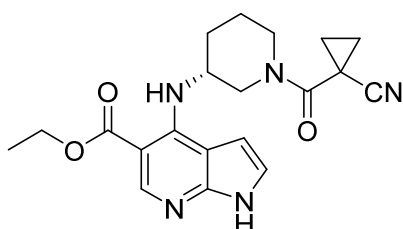
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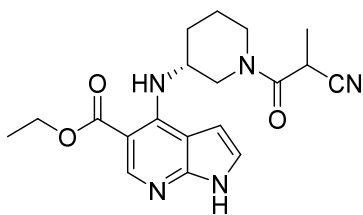
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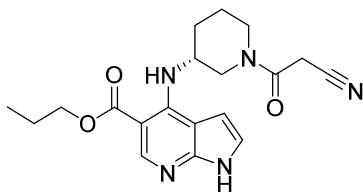
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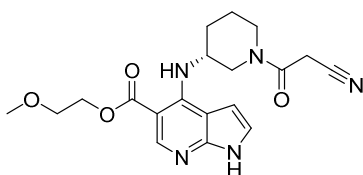
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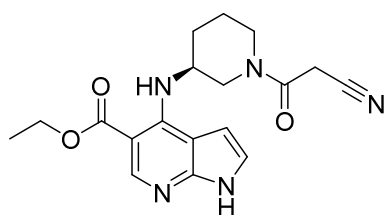
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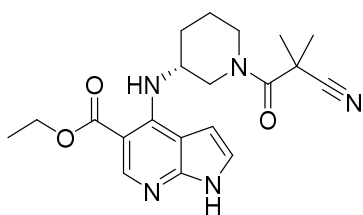
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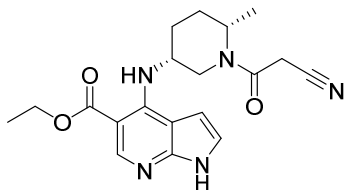
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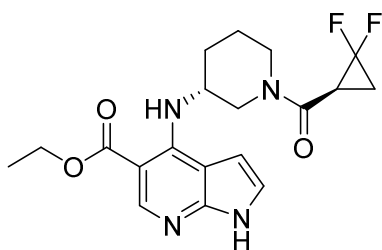
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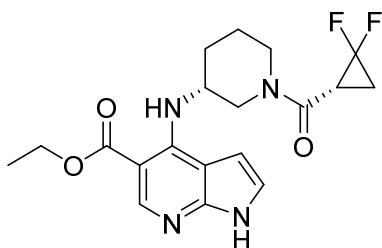
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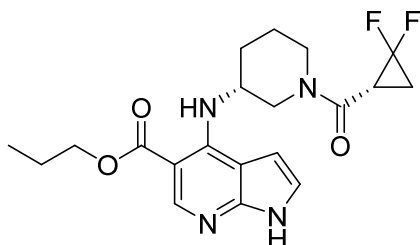
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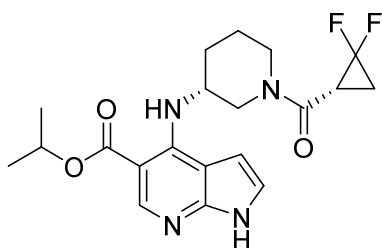
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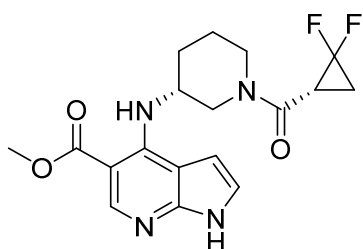
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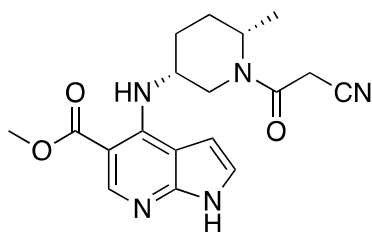
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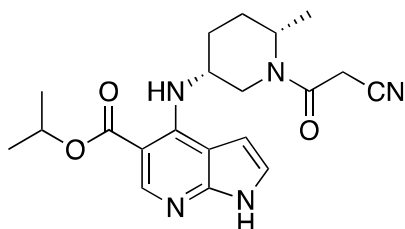
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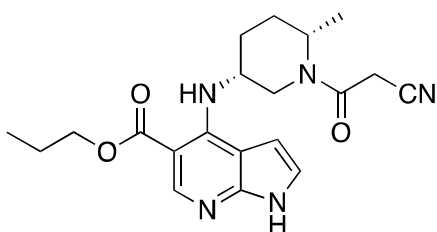
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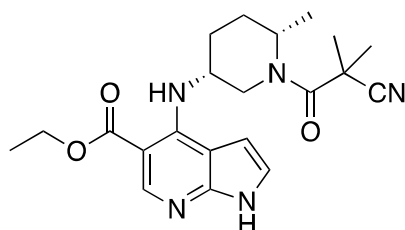
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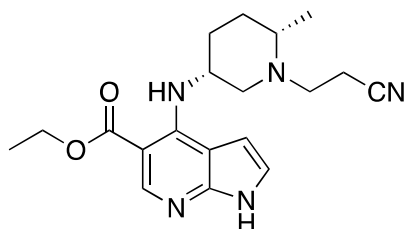
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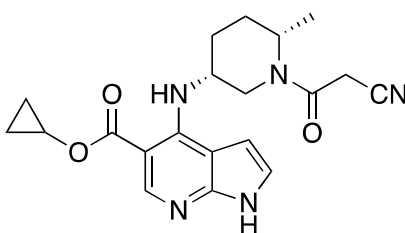
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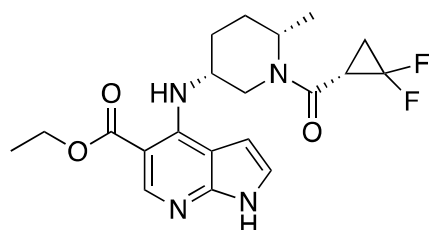
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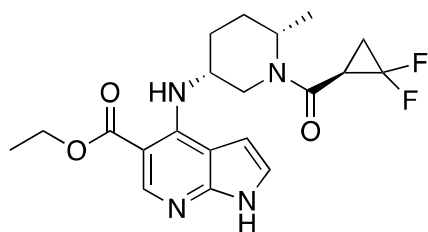
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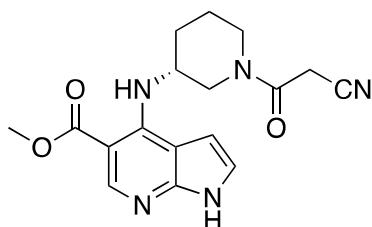
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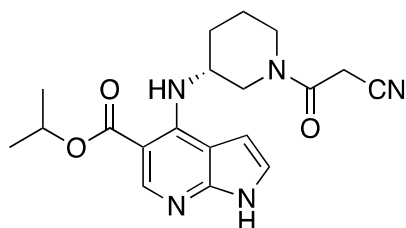
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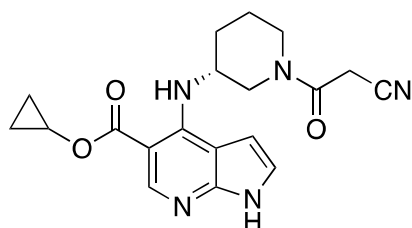
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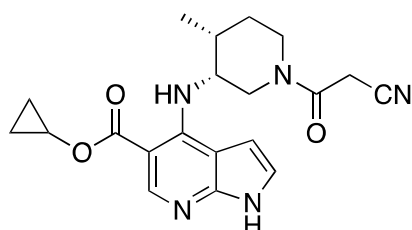
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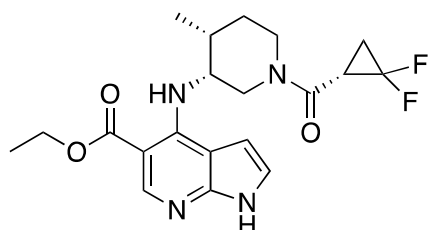
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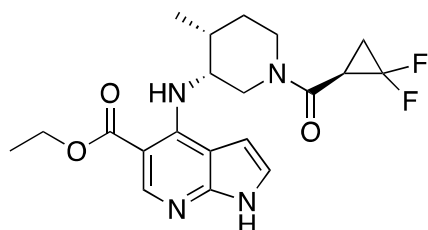
cyclopropyl (*R*)-4-((1-(2-cyanoacetyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



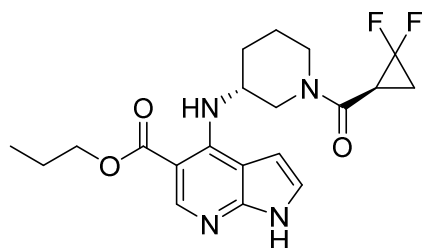
cyclopropyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



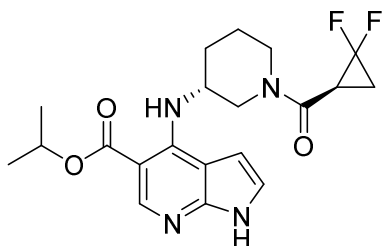
ethyl 4-(((3*R*,4*R*)-1-((*S*)-2,2-difluorocyclopropane-1-carbonyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



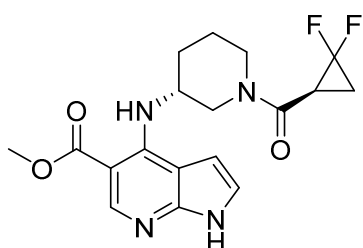
ethyl 4-(((3*R*,4*R*)-1-((*R*)-2,2-difluorocyclopropane-1-carbonyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



propyl 4-(((*R*)-1-((*S*)-2,2-difluorocyclopropane-1-carbonyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,

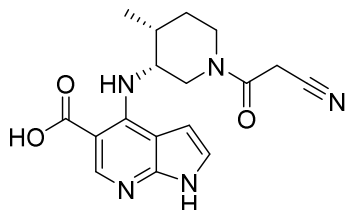


isopropyl 4-(((*R*)-1-((*S*)-2,2-difluorocyclopropane-1-carbonyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate, or

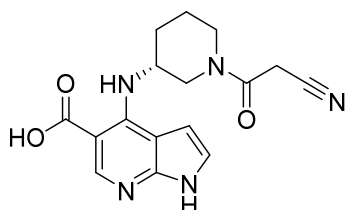


methyl 4-(((*R*)-1-((*S*)-2,2-difluorocyclopropane-1-carbonyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate.

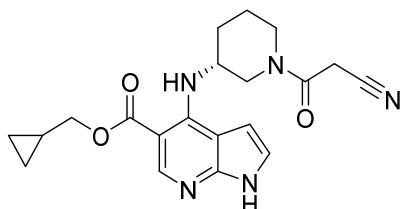
11. The compound of Claim 9, wherein the compound is selected from the group consisting of:



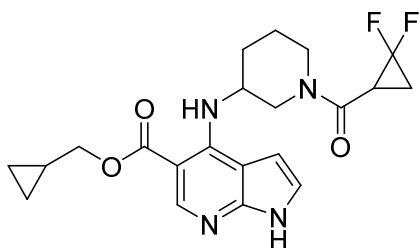
4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylic acid,



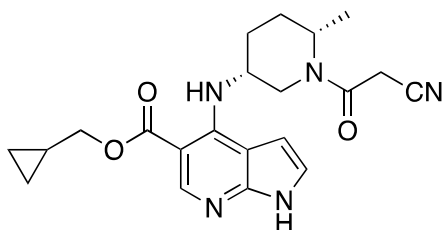
(*R*)-4-((1-(2-cyanoacetyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylic acid,



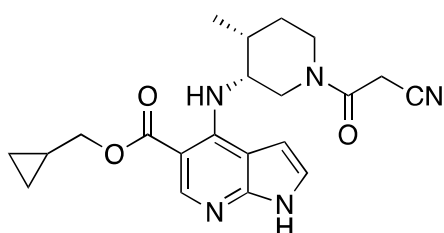
cyclopropylmethyl (*R*)-4-((1-(2-cyanoacetyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,



cyclopropylmethyl 4-((1-(2,2-difluorocyclopropane-1-carbonyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate,

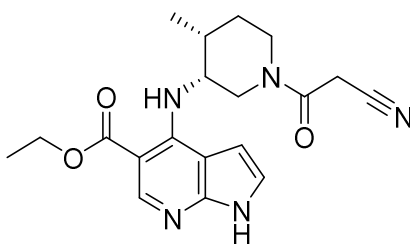


cyclopropylmethyl 4-(((3*R*,6*S*)-1-(2-cyanoacetyl)-6-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate, or



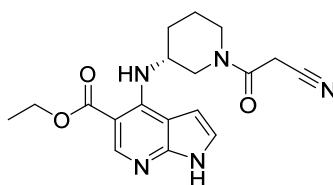
cyclopropylmethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate.

12. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 1, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
13. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 9, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
14. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 10, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
15. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 11, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
16. The compound of Claim 10, wherein the compound is:



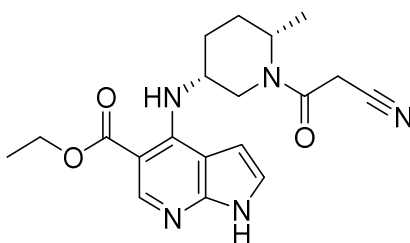
ethyl 4-(((3*R*,4*R*)-1-(2-cyanoacetyl)-4-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate.

17. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 16, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
18. The compound of Claim 10, wherein the compound is:



ethyl (*R*)-4-((1-(2-cyanoacetyl)piperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate.

19. A pharmaceutical composition comprising a therapeutically effective amount of a compound of Claim 18, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.
20. The compound of Claim 10, wherein the compound is:



ethyl 4-(((3*R*,6*S*)-1-(2-cyanoacetyl)-6-methylpiperidin-3-yl)amino)-1*H*-pyrrolo[2,3-*b*]pyridine-5-carboxylate.

21. A pharmaceutical composition comprising a therapeutically effective amount of a

compound of Claim 20, a pharmaceutically acceptable salt thereof, a derivative thereof, or a combination thereof; and a pharmaceutically acceptable carrier.