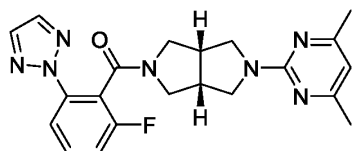


What is claimed is:

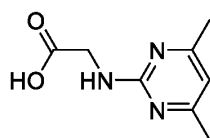
1. A process of preparing (((3a*R*,6a*S*)-5-(4,6-dimethylpyrimidin-2-yl)hexahydropyrrolo[3,4-*c*]pyrrol-2(1*H*)-yl)(2-fluoro-6-(2*H*-1,2,3-triazol-2-yl)phenyl)methanone

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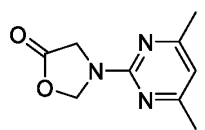
said process comprising step described below:

- a) Oxazolidination of (4,6-dimethylpyrimidin-2-yl)glycine,



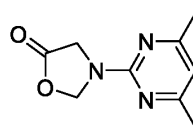
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wherein said oxazolidination is characterized by the use of formaldehyde or paraformaldehyde to obtain 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one



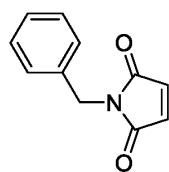
2. The process of claim 1, said process further comprising reaction of 3-(4,6-

dimethylpyrimidin-2-yl)oxazolidin-5-one



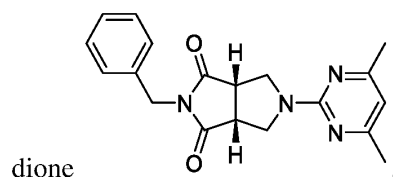
with 1-benzyl-1*H*-pyrrole-

2,5-dione

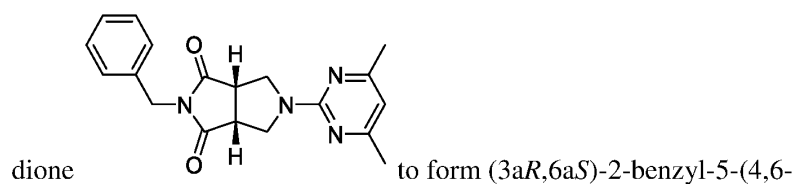


at a temperature greater than 250 °C to form (3a*R*,6a*S*)-2-

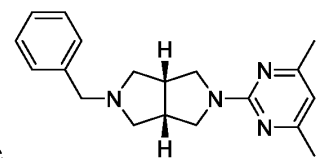
benzyl-5-(4,6-dimethylpyrimidin-2-yl)tetrahydropyrrolo[3,4-c]pyrrole-1,3(2*H*,3*aH*)-



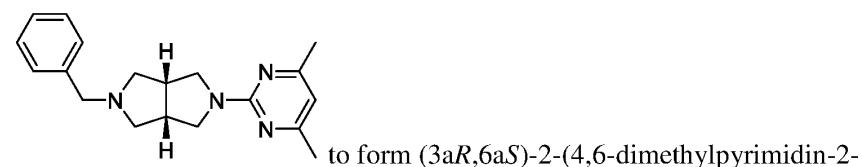
- 5 3. The process of claim 2, said process further comprising reduction of (3*aR*,6*aS*)-2-benzyl-5-(4,6-dimethylpyrimidin-2-yl)tetrahydropyrrolo[3,4-c]pyrrole-1,3(2*H*,3*aH*)-



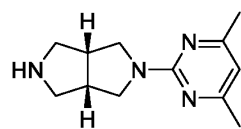
dimethylpyrimidin-2-yl)octahydropyrrolo[3,4-c]pyrrole



- 10 4. The process of claim 3, said process further comprising deprotection of (3*aR*,6*aS*)-2-benzyl-5-(4,6-dimethylpyrimidin-2-yl)octahydropyrrolo[3,4-c]pyrrole



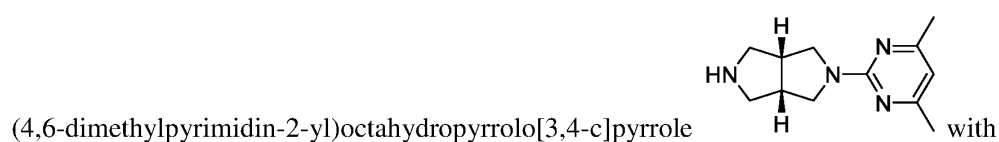
yl)octahydropyrrolo[3,4-c]pyrrole

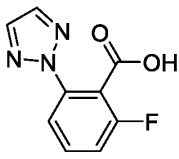


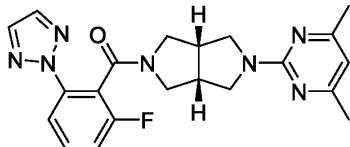
by means of 10% (w/w) Pd/C and ammonium formate.

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5. The process of claim 4, said process further comprising amidation of (3*aR*,6*aS*)-2-

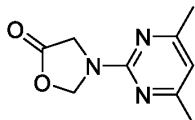


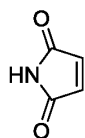
2-fluoro-6-(2*H*-1,2,3-triazol-2-yl)benzoic acid  by means of SOCl₂ to form (((3*aR*,6*aS*)-5-(4,6-dimethylpyrimidin-2-yl)hexahydropyrrolo[3,4-*c*]pyrrol-2(1*H*)-

yl)(2-fluoro-6-(2*H*-1,2,3-triazol-2-yl)phenyl)methanone 

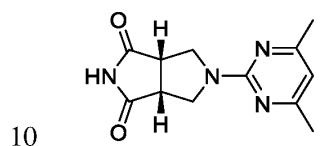
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6. The process of claim 1, said process further comprising reaction of 3-(4,6-

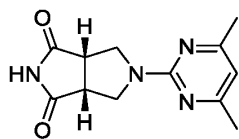
dimethylpyrimidin-2-yl)oxazolidin-5-one  with 1*H*-pyrrole-2,5-dione



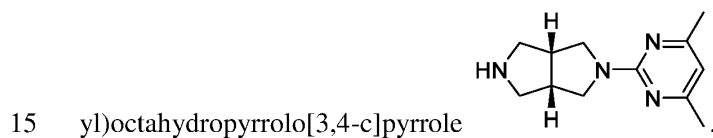
at a temperature greater than 250 °C to form (3*aR*,6*aS*)-5-(4,6-dimethylpyrimidin-2-yl)tetrahydropyrrolo[3,4-*c*]pyrrole-1,3(2*H*,3*aH*)-dione



7. The process of claim 6, said process further comprising reduction of (3*aR*,6*aS*)-5-(4,6-dimethylpyrimidin-2-yl)tetrahydropyrrolo[3,4-*c*]pyrrole-1,3(2*H*,3*aH*)-dione

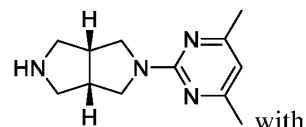


to form (3*aR*,6*aS*)-2-(4,6-dimethylpyrimidin-2-



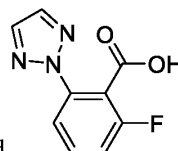
8. The process of claim 7, said process further comprising amidation of (3a*R*,6a*S*)-2-

(4,6-dimethylpyrimidin-2-yl)octahydropyrrolo[3,4-*c*]pyrrole



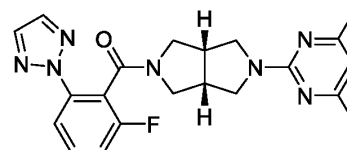
with

2-fluoro-6-(2*H*-1,2,3-triazol-2-yl)benzoic acid



by means of SOCl₂ to

form (((3a*R*,6a*S*)-5-(4,6-dimethylpyrimidin-2-yl)hexahydropyrrolo[3,4-*c*]pyrrol-2(1*H*)-



5 yl)(2-fluoro-6-(2*H*-1,2,3-triazol-2-yl)phenyl)methanone

9. The process of claim 2, wherein said 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one is not isolated prior to reaction with said 1-benzyl-1*H*-pyrrole-2,5-dione.

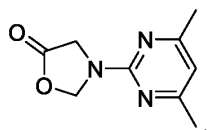
10. The process of claim 2, wherein said 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one is isolated prior to reaction with said 1-benzyl-1*H*-pyrrole-2,5-dione.

11. The process of claim 6, wherein said 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one is not isolated prior to reaction with said 1*H*-pyrrole-2,5-dione.

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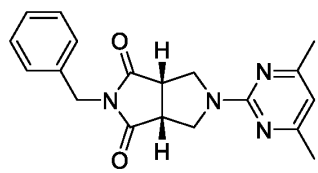
12. The process of claim 6, wherein said 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one is isolated prior to reaction with said 1*H*-pyrrole-2,5-dione.

13. A compound which is 3-(4,6-dimethylpyrimidin-2-yl)oxazolidin-5-one



20 14. A compound which is (3a*R*,6a*S*)-2-benzyl-5-(4,6-dimethylpyrimidin-2-

yl)tetrahydropyrrolo[3,4-*c*]pyrrole-1,3(2*H*,3a*H*)-dione



15. A compound which is (3a*R*,6a*S*)-5-(4,6-dimethylpyrimidin-2-

yl)tetrahydropyrrolo[3,4-*c*]pyrrole-1,3(2*H*,3a*H*)-dione

