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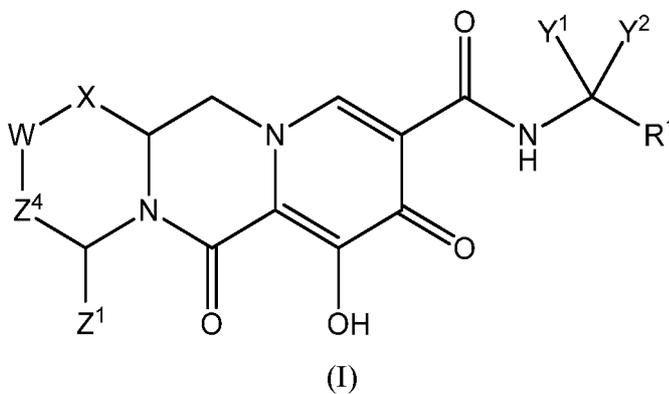
(54) Title **POLYCYCLIC-CARBAMOYLPYRIDONE COMPOUNDS AND THEIR PHARMACEUTICAL
USE**

(56) References
Cited: WO-A1-2006/116764
EP-A1- 2 412 709

Enclosed is a translation of the patent claims in Norwegian. Please note that as per the Norwegian Patents Acts, section 66i the patent will receive protection in Norway only as far as there is agreement between the translation and the language of the application/patent granted at the EPO. In matters concerning the validity of the patent, language of the application/patent granted at the EPO will be used as the basis for the decision. The patent documents published by the EPO are available through Espacenet (<http://worldwide.espacenet.com>) or via the search engine on our website here: <https://search.patentstyret.no/>

Patentkrav

1. Forbindelse med formel (I):



eller en stereoisomer eller et farmasøytisk akseptabelt salt derav, hvor:

5 X er -O-;

W er -CHZ²-;

Z¹ og Z², tatt sammen, danner -L- hvor L er -C(R^a)₂C(R^a)₂-;

Z⁴ er -CH₂-;

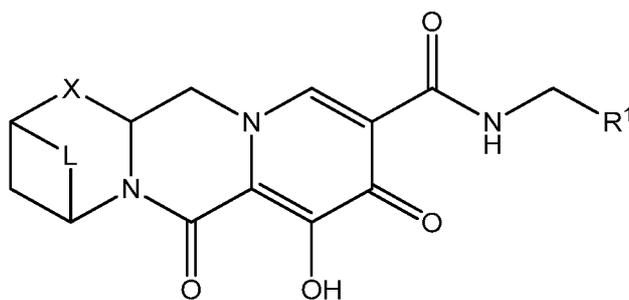
Y¹ og Y² er hver, uavhengig, hydrogen, C₁₋₃-alkyl eller C₁₋₃-halogenalkyl;

10 R¹ er fenyl substituert med ett eller to halogenatomer; og

hver R^a er, uavhengig, hydrogen, halogen, hydroksy eller C₁₋₄-alkyl.

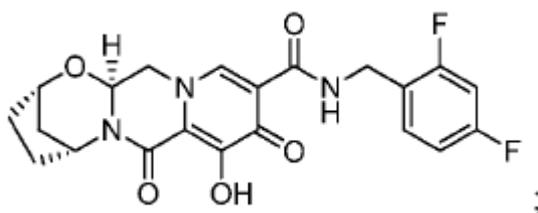
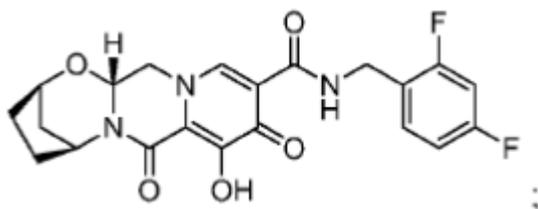
2. Forbindelse ifølge krav 1, med formel (II-A):

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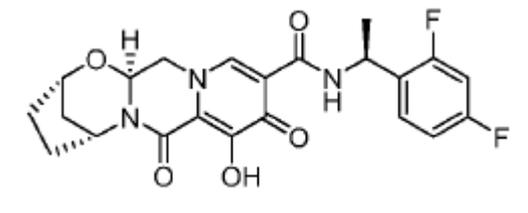
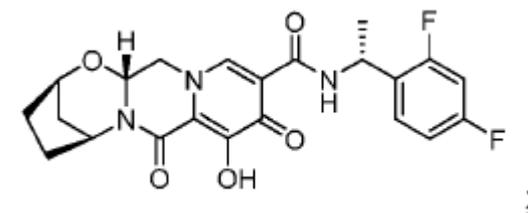
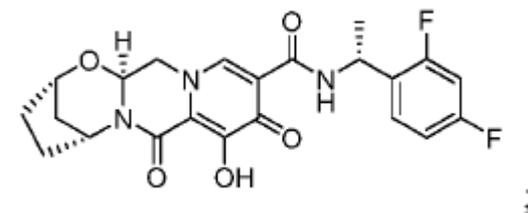
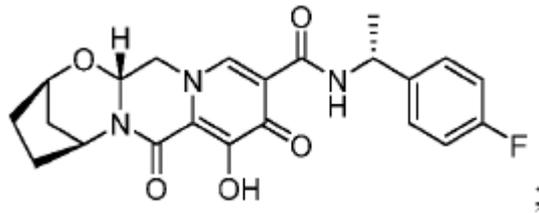
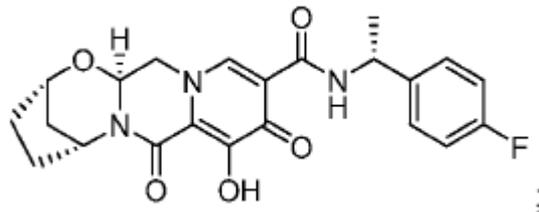


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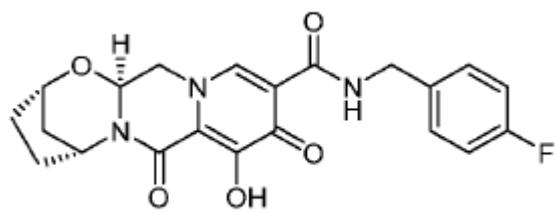
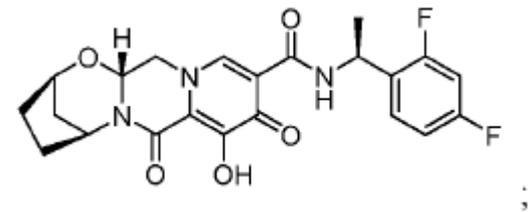
3. Forbindelse ifølge krav 1, hvor Y¹ og Y² er hver uavhengig hydrogen, metyl eller trifluormetyl.
4. Forbindelse ifølge krav 1 eller krav 2, hvor R¹ er substituert med ett
5 halogenatom.
5. Forbindelse ifølge krav 4, hvor R¹ er 4-fluorfenyl eller 2-fluorfenyl.
6. Forbindelse ifølge krav 1 eller krav 2, hvor R¹ er substituert med to halogenatomer.
7. Forbindelse ifølge krav 6, hvor R¹ er 2,4-difluorfenyl, 2,3-difluorfenyl, 2,6-
10 difluorfenyl, 3-fluor-4-klorfenyl, 3,4-difluorfenyl, 2-fluor-4-klorfenyl eller 3,5-difluorfenyl.
8. Forbindelse ifølge krav 7, hvor R¹ er 2,4-difluorfenyl.
9. Forbindelse ifølge krav 1, valgt fra:



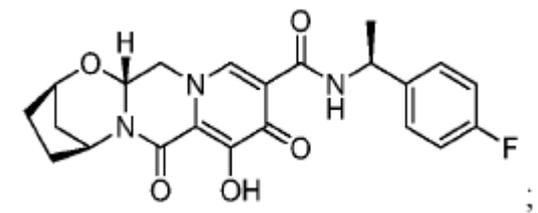
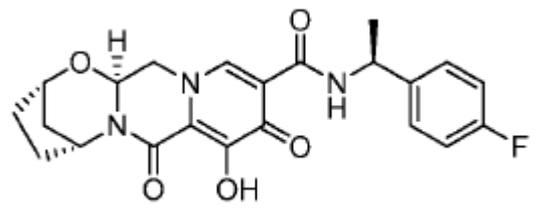
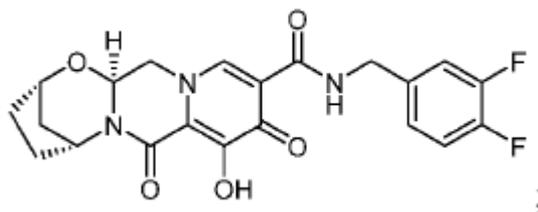
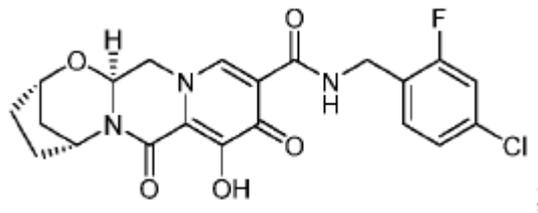
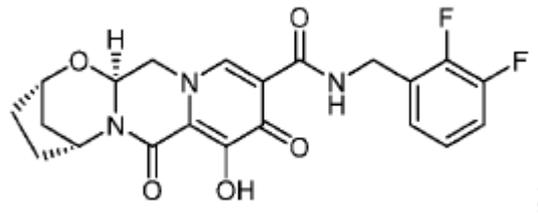
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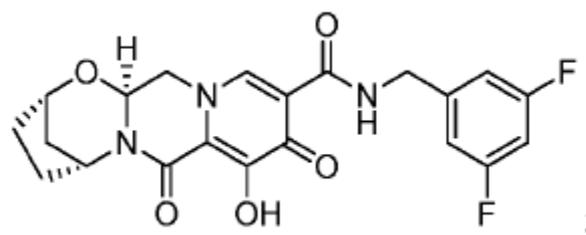
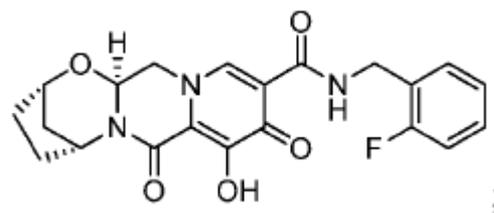
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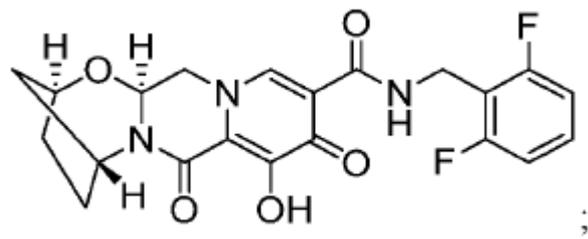
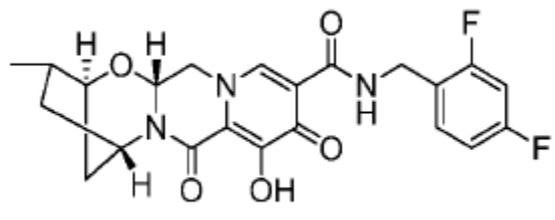
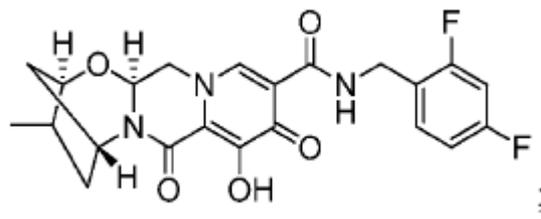
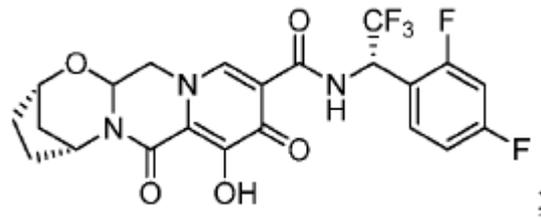
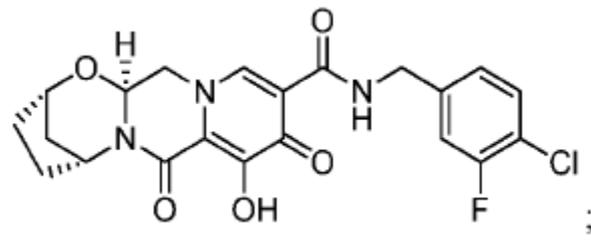
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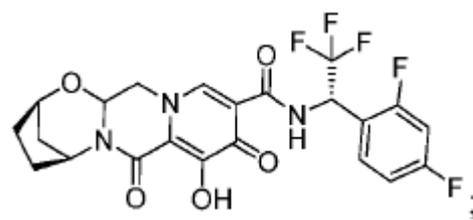
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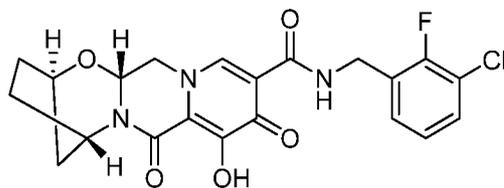
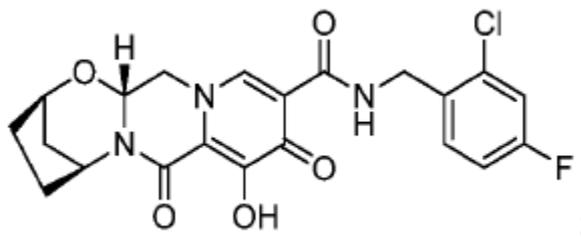


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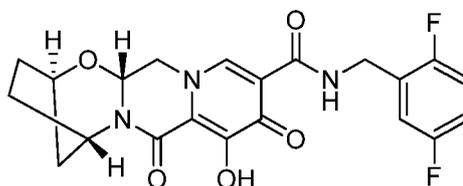


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og



- 5 10. Farmasøytisk sammensetning omfattende en forbindelse ifølge et hvilket som helst av kravene 1-9, eller en stereoisomer eller et farmasøytisk akseptabelt salt derav, og en farmasøytisk akseptabel bærer, et fortynningsmiddel eller en eksipiens.
- 10 11. Farmasøytisk sammensetning ifølge krav 10, ytterligere omfattende ett eller flere ytterligere terapeutiske midler.
12. Farmasøytisk sammensetning ifølge krav 11, hvor det ene eller flere ytterligere terapeutiske midler er et middel mot humant immunsviktivirus.
- 15 13. Farmasøytisk sammensetning ifølge krav 12, hvor det ene eller flere ytterligere terapeutiske midler er valgt fra gruppen bestående av humant immunsviktivirus-proteasehemmere, humant immunsviktivirus-ikke-nukleosidhemmere av revers transkriptase, humant immunsviktivirus-nukleosidhemmere av revers transkriptase, humant immunsviktivirus-nukleotidhemmere av revers transkriptase, og kombinasjoner derav.