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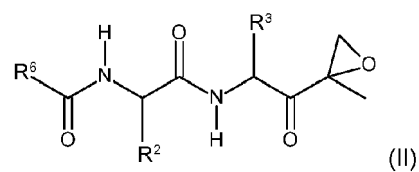
(54)	Title	DIPEPTIDE AND TRIPEPTIDE EPOXY KETONE PROTEASE INHIBITORS
(56)	References Cited:	US-A1- 2007 105 786, EVA HUBER ET AL: "Immuno- and Constitutive Proteasome Crystal Structures Reveal Differences in Substrate and Inhibitor Specificity", CELL, CELL PRESS, US, vol. 148, no. 4, 2 December 2011 (2011-12-02), pages 727-738, XP028458988, ISSN: 0092- 8674, DOI: 10.1016/J.CELL.2011.12.030 [retrieved on 2012-01-12], WO-A2-2007/149512,

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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 (II) eller et farmasøytisk akseptabelt salt av samme,



hvor:

- 5 R^2 og R^3 er hver uavhengig valgt fra C_{1-6} aralkyl og C_{1-6} heteroaralkyl; og R^6 er valgt fra heterosykl og heteroaryl.

2. Forbindelse ifølge krav 1, hvor R^2 er C_{1-6} heteroaralkyl.

- 10 3. Forbindelse ifølge krav 1, hvor R^2 er C_{1-6} aralkyl.

4. Forbindelse ifølge krav 1, hvor R^3 er C_{1-6} heteroaralkyl.

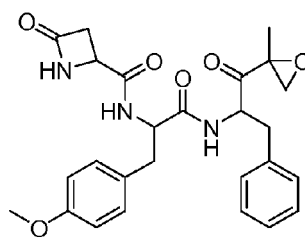
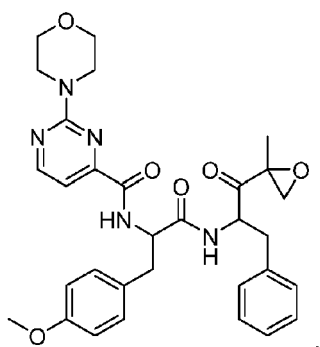
5. Forbindelse ifølge krav 1, hvor R^3 er C_{1-6} aralkyl.

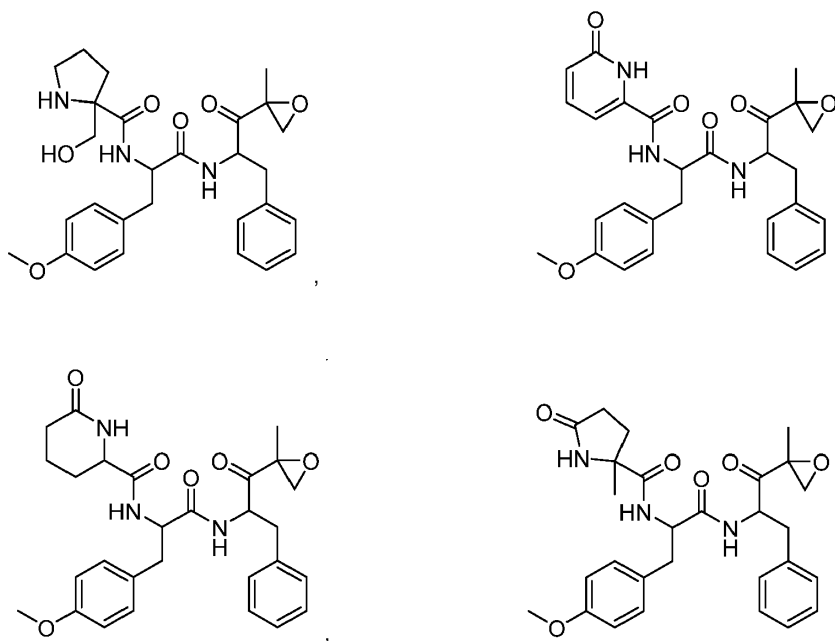
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6. Forbindelse ifølge krav 1, hvor R^6 er heterosykl.

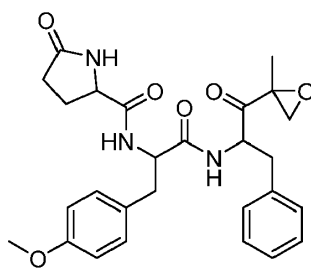
7. Forbindelse ifølge krav 1, hvor R^6 er heteroaryl.

- 20 8. Forbindelse ifølge krav 1 med en struktur valgt fra gruppen bestående av:





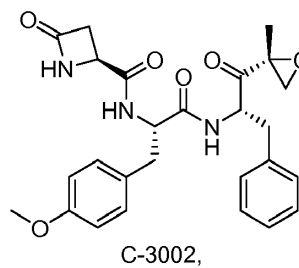
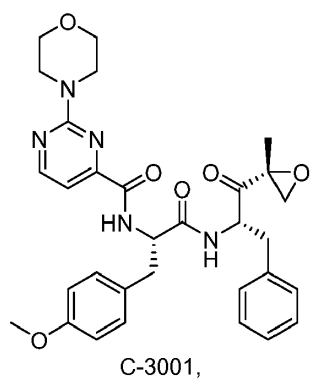
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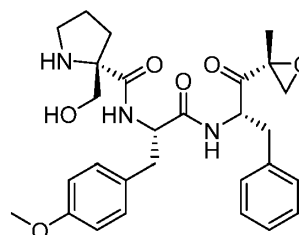
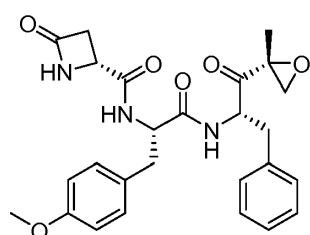
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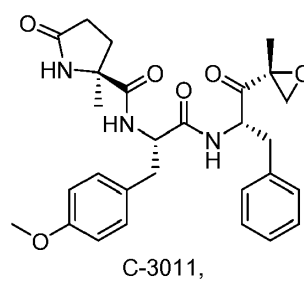
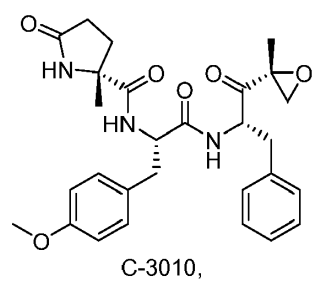
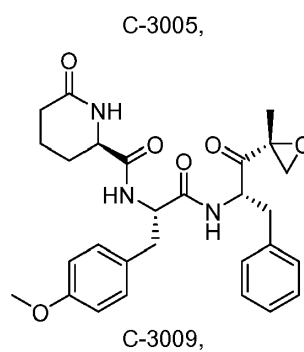
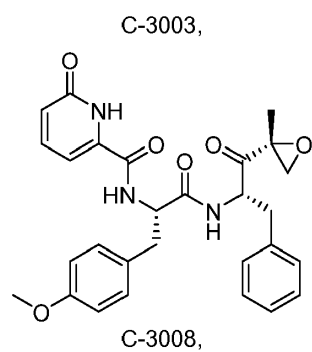
eller et farmasøytisk akseptabelt salt av samme.

9. Forbindelse ifølge krav 1 med en struktur valgt fra gruppen bestående av:

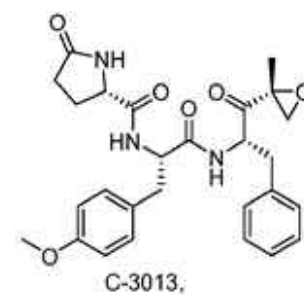
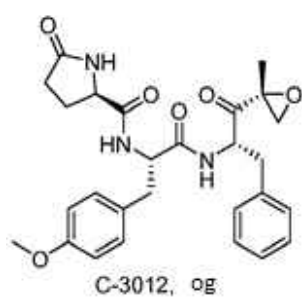


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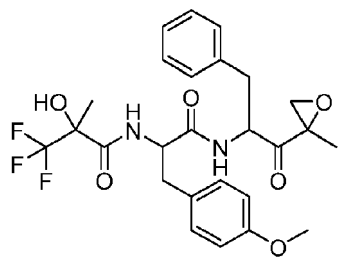


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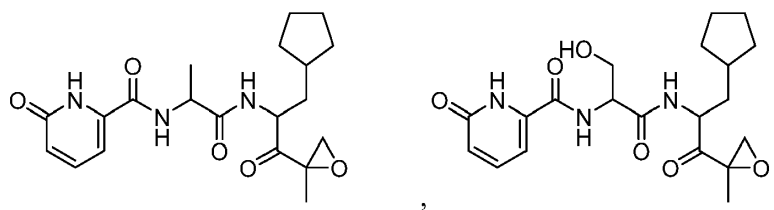
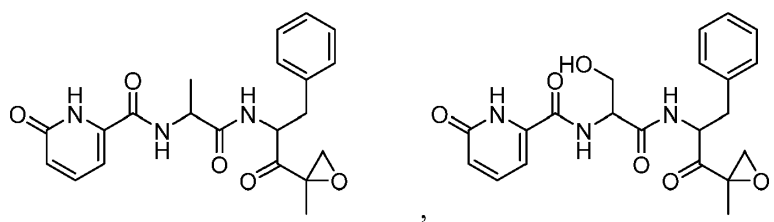
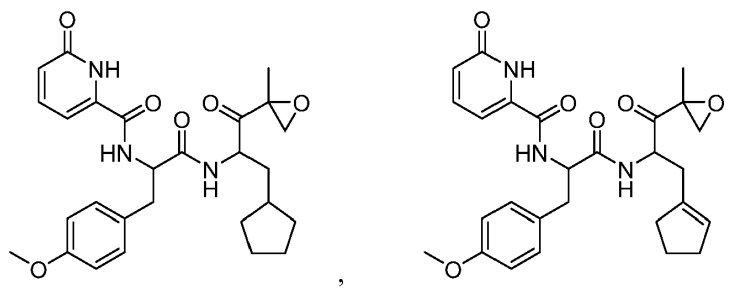
eller et farmasøytisk akseptabelt salt av samme.

10. Forbindelse med en struktur valgt fra gruppen bestående av:

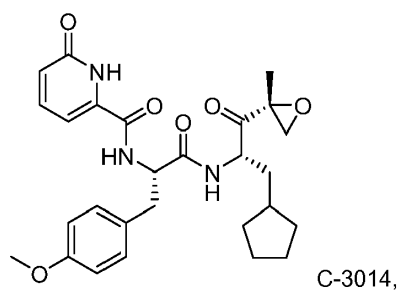
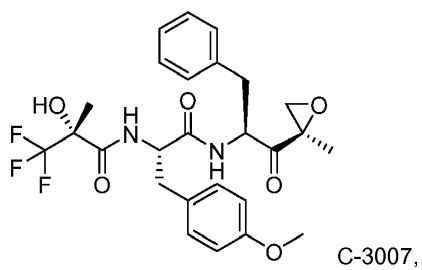


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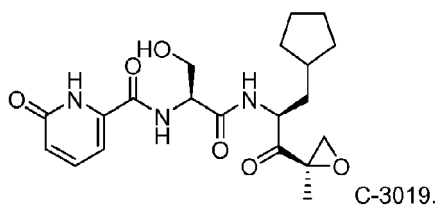
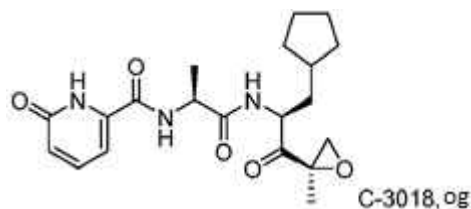
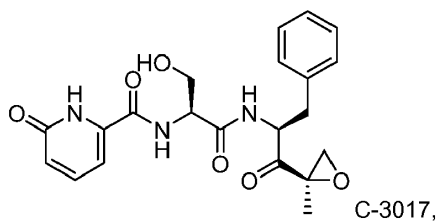
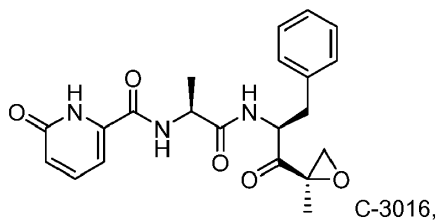
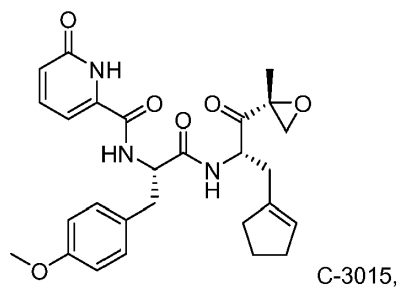
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10 11. Farmasøytisk blanding som omfatter forbindelsen ifølge ett av kravene 1 til 10, eller et
farmasøytisk akseptabelt salt av samme, og en farmasøytisk akseptabel bærer eller
fortynningsmiddel.

12. Forbindelse ifølge ett av kravene 1 til 10, eller blandingen ifølge krav 11, for anvendelse i en
15 framgangsmåte for inhibering av en immunoproteasom av en celle.

13. Forbindelse ifølge ett av kravene 1 to 10 eller blandingen ifølge krav 11 for anvendelse i inhibering av en immunoproteasom i et menneske.

14. Forbindelse for anvendelse ifølge krav 12 eller krav 13, hvori forbindelsen inhiberer LMP2-
5 underenheten av nevnte immunoproteasom.

15. Forbindelse ifølge ett av kravene 1 to 10, eller blandingen ifølge krav 11, for anvendelse i behandlingen av en sykdom valgt fra gruppen bestående av:

- (a) en immun-relatert sykdom; eller
- 10 (b) kreft; eller
- (c) inflammasjon; eller
- (d) infeksjon; eller
- (e) proliferativ sykdom; og
- (f) neurodegenerativ sykdom.

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