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(54)	Title	EFFECTIVE AMOUNTS OF (3AR)-1,3A,8-TRIMETHYL-1,2,3,3A,8,8A- HEXAHYDROPYRROLO[2,3-B]INDOL-5-YL PHENYLCARBAMATE AND METHODS THEREOF
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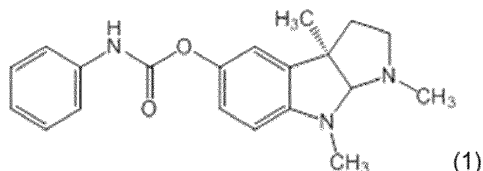
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Enclosed is a translation of the patent claims in Norwegian. Please note that as per the Norwegian Patents Acts, section 66i the patent will receive protection in Norway only as far as there is agreement between the translation and the language of 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. Farmasøytisk preparat som omfatter en farmasøytisk akseptabel bærer og en mengde forbindelse (1) eller et salt derav



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for bruk i behandling av demens hos et menneske, hvor fra 3 µg/kg til 1 mg/kg kroppsvekt av forbindelse (1) administreres en gang om dagen, hvor administrering av nevnte preparat resulterer i et maksimalt plasma-sirkulasjonsnivå av forbindelse (1) lik eller større enn 60 ng/ml i nevnte individ hvorved demens hos nevnte individ blir behandlet.

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2. Farmasøytisk preparat for anvendelse ifølge krav 1, hvor nevnte demens er Alzheimers sykdom

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3. Farmasøytisk preparat for anvendelse ifølge krav 1, hvor det maksimale plasma-sirkulasjonsnivået er oppnådd innen omtrent 6 timer fra administreringstidspunktet.

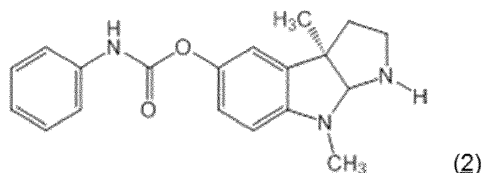
4. Farmasøytisk preparat for anvendelse ifølge krav 1, hvor plasma-sirkulasjonsnivået av forbindelse (1) er lik eller større enn omtrent 20 ng/ml i minst 12 timer etter nevnte administrering.

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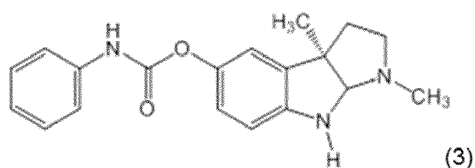
5. Farmasøytisk preparat for anvendelse ifølge krav 1, hvor administreringen resulterer i en maksimal plasmakonsentrasjon i nevnte individ av en eller flere verdier valgt fra gruppen bestående av:

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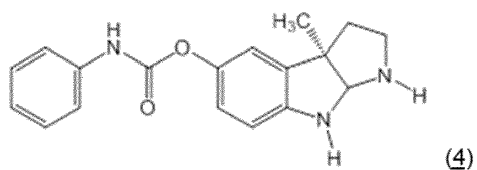
(i) minst 20 ng/ml forbindelse (2);



(ii) minst 20 ng/ml forbindelse (3);



(iii) minst 2 ng/ml forbindelse (4);



og en kombinasjon derav.

- 5 6. Farmasøytisk preparat for anvendelse ifølge krav 1, hvor administreringen resulterer i en plasmakonsentrasjon med jevnt nivå i nevnte individ av en eller flere verdier valgt fra gruppen bestående av: minst omtrent 100 ng/ml forbindelse (1); minst omtrent 10 ng/ml forbindelse (2); minst omtrent. 10 ng/ml forbindelse (3); minst omtrent 3 ng/ml forbindelse (4); og en kombinasjon derav.