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Norwegian Industrial Property Office

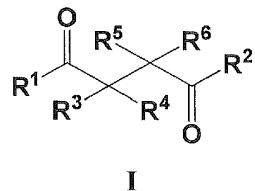
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(54)	Title	DIKETONES AND HYDROXYKETONES AS CATENIN SIGNALING PATHWAY ACTIVATORS
(56)	References Cited:	WO-A1-2010/075551, US-A- 4 014 889, US-A- 4 032 526, US-A1- 2005 267 110, YOSHIMURA HIROYUKI ET AL: "Discovery of Novel and Potent Retinoic Acid Receptor.alpha. Agonists: Syntheses and Evaluation of Benzofuranyl-pyrrole and Benzothiophenyl-pyrrole Derivatives", JOURNAL OF MEDICINAL CHEMISTRY, AMERICAN CHEMICAL SOCIETY, US, vol. 43, 1 January 2000 (2000-01-01), pages 2929-2937, XP002214937, ISSN: 0022-2623, DOI: 10.1021/JM000098S, US-A1- 2009 232 754, US-A1- 2010 152 493, US-B2- 7 041 837, F. V. BRUCHHAUSEN, K. LINGNER: "Eine Synthese von DL-Asarinin und DL-Sesamin", ARCHIV DER PHARMAZIE, vol. 290, 1957, pages 1-16, XP002718639,, US-A1- 2008 146 555

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/>

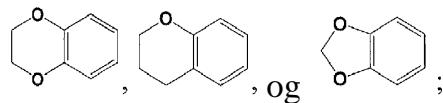
Patentkrav

5 1. Forbindelse eller farmasøytisk akseptabelt salt derav med strukturen i formel I:



hvor R^1 er et substituert eller usubstituert heteroaryl, hvor heteroarylet er valgt

10 fra gruppen bestående av:



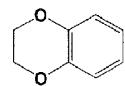
underlagt den betingelse at et karbonatom i R^1 -heteroarylet er knyttet til karbonylet;

15 R^2 er et substituert eller usubstituert aryl, hvor arylet er valgt fra fenyl eller
naftyl; og

R^3 , R^4 , R^5 og R^6 er H.

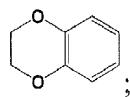
2. Forbindelse ifølge krav 1, hvor R^1 er

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3. Forbindelse ifølge krav 1, hvor R^2 er et substituert eller usubstituert fenyl.

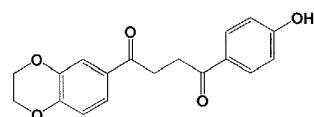
25 4. Forbindelse ifølge krav 1, hvor R^1 er



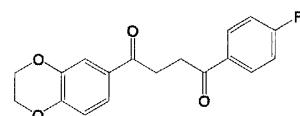
og R² er et substituert eller usubstituert fenyL

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5. Forbindelse ifølge krav 4, hvor forbindelsen med formel I er:

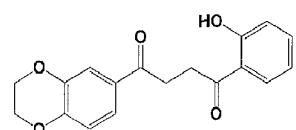


- 10 6. Forbindelse ifølge krav 4, hvor forbindelsen med formel I er:



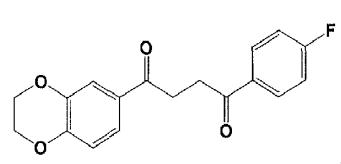
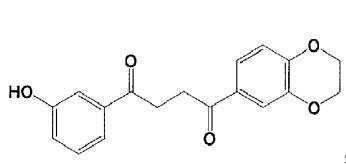
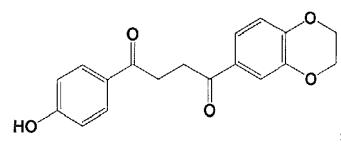
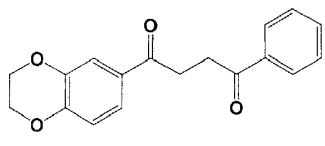
7. Forbindelse ifølge krav 4, hvor forbindelsen med formel I er:

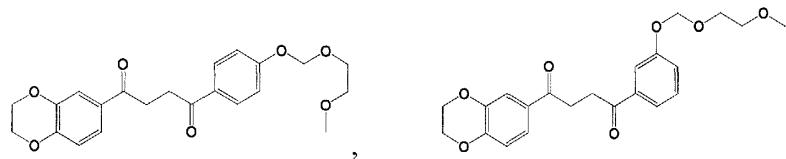
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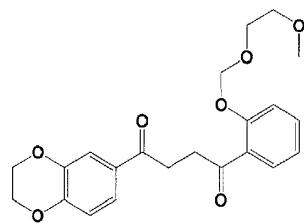
8. Forbindelse ifølge krav 1, med en struktur valgt fra gruppen bestående av:

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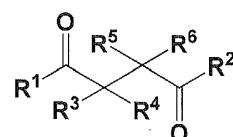




5 og



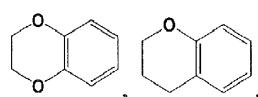
- 10 9. Farmasøytisk sammensetning, omfattende en terapeutisk virksom mengde av en forbindelse ifølge krav 1 eller et farmasøytisk akseptabelt salt derav med strukturen i formel I:



I

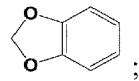
15

hvor R^1 er et substituert eller usubstituert heteroaryl, hvor heteroarylet er valgt fra gruppen bestående av



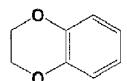
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og



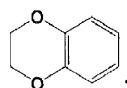
- 5 underlagt den betingelse at et karbonatom i R¹-heteroarylet er knyttet til karbonylet;
- R² er et substituert eller usubstituert aryl, hvor arylet er valgt fra fenyl eller naftyl; og
- R³, R⁴, R⁵ og R⁶ er H;
- 10 og et farmasøytisk akseptabelt tilsetningsmiddel.

10. Farmasøytisk sammensetning ifølge krav 9, hvor R¹ er



15

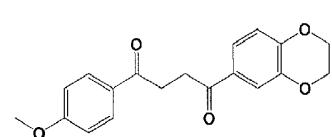
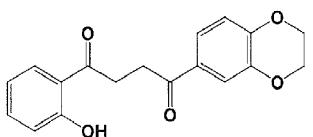
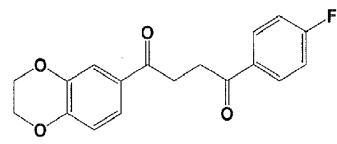
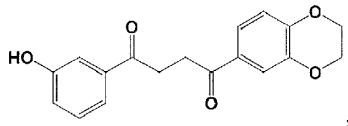
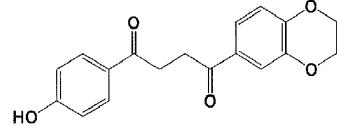
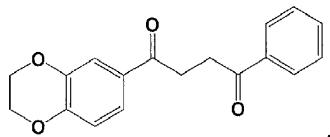
11. Farmasøytisk sammensetning ifølge krav 9, hvor R² er et substituert eller usubstituert fenyl.
- 20 12. Farmasøytisk sammensetning ifølge krav 9, hvor R¹ er



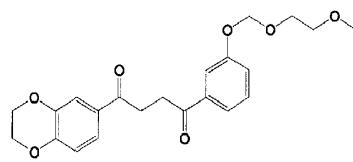
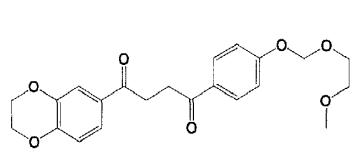
og R² er et substituert eller usubstituert fenyl.

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13. Farmasøytisk sammensetning ifølge krav 9, med en struktur valgt fra gruppen bestående av:

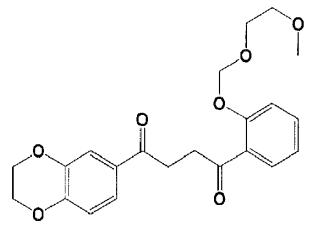


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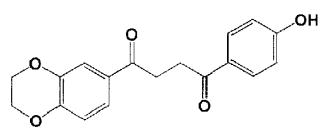
og

10



eller et farmasøytisk akseptabelt salt derav.

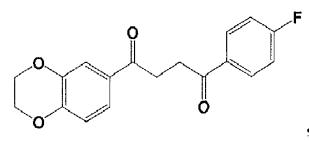
15 14. Farmasøytisk sammensetning ifølge krav 9, hvor forbindelsen med formel I er:



eller et farmasøytisk akseptabelt salt derav.

15. Farmasøytisk sammensetning ifølge krav 9, hvor forbindelsen med formel I er:

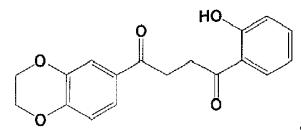
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eller et farmasøytisk akseptabelt salt derav.

16. Farmasøytisk sammensetning ifølge krav 9, hvor forbindelsen med formel I er:

10



eller et farmasøytisk akseptabelt salt derav.