

CORRECTED VERSION

(19) World Intellectual Property Organization
International Bureau(43) International Publication Date
6 June 2002 (06.06.2002)

PCT

(10) International Publication Number
WO 02/043648 A2

(51) International Patent Classification⁷: **A61K**

(21) International Application Number: PCT/IB01/02302

(22) International Filing Date:
28 November 2001 (28.11.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
0029102.1 29 November 2000 (29.11.2000) GB

(81) Designated States (*national*): AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZM, ZW.

(84) Designated States (*regional*): ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

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Published:

— without international search report and to be republished upon receipt of that report

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(48) Date of publication of this corrected version:

6 September 2002

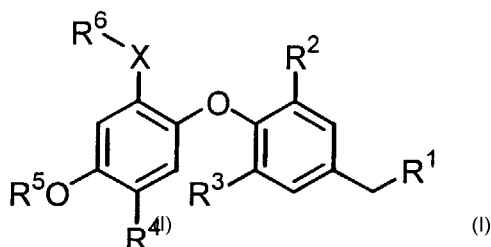
(15) Information about Correction:

see PCT Gazette No. 36/2002 of 6 September 2002, Section II

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For two-letter codes and other abbreviations, refer to the "Guidance Notes on Codes and Abbreviations" appearing at the beginning of each regular issue of the PCT Gazette.

(54) Title: COMPOUNDS ACTIVE AT THE GLUCOCORTICOID RECEPTOR II



hydrogen, halogen, and C₁₋₆ alkyl, provided that one of R² or R³ is other than hydrogen; R⁴ is selected from: C₁₋₆ alkyl, C₂₋₆ alkenyl, C₂₋₆ alkynyl, halogen, (R⁹)(R¹⁰)N, R⁸C(Z)N(R¹¹), R⁸OC(Z)N(R¹¹), (R⁹)(R¹⁰)NC(Z)N(R¹¹), R⁸S(O)₂N(R¹¹), (R⁹)(R¹⁰)NS(O)₂N(R¹¹), and R⁸SC(Z)N(R¹¹); R⁵ is selected from: (i) C₁₋₆-alkyl which is substituted by a group selected from A, provided that A is not halogen; (ii) C₇₋₁₂-alkyl, C₂₋₁₂-alkenyl and C₂₋₁₂-alkynyl; (iii) C₁₋₁₂-alkyl, where one or more carbon atoms are replaced by Y, and where one or more carbons are optionally substituted by a group selected from A, provided that if more than one carbon is replaced by Y, the said Y groups are not directly connected to each other; R⁶ is selected from: C₁₋₁₂-alkyl, C₃₋₈-cycloalkyl, C₃₋₈-heterocycloalkyl, C₂₋₆-alkenyl, and C₂₋₆-alkynyl, aryl, and heteroaryl; R⁷ is optionally selected from hydrogen; R⁸, R⁹, R¹⁰ and R¹¹ are independently of each other selected from: hydrogen, C₁₋₆-alkyl, C₃₋₈-cycloalkyl, C₂₋₆-alkenyl, C₂₋₆-alkynyl, and C₃₋₈-heterocycloalkyl, aryl and heteroaryl; or where any pair of R⁸, R⁹, R¹⁰ and R¹¹ together with the atom or atoms to which they are bound form a ring having 3-7 ring members; or pharmaceutically acceptable salts, stereoisomers or prodrugs thereof.