12/11/02
2002068625

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization
International Bureau

ACTION

(43) International Publication Date
6 December 2001 (06.12.2001)

PCT

(10) International Publication Number
WO 01/92264 A1

(51) International Patent Classification[7]: C07D 487/14, A61K 31/505, A61P 25/28

(21) International Application Number: PCT/US01/16954

(22) International Filing Date: 24 May 2001 (24.05.2001)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:
60/207,143 26 May 2000 (26.05.2000) US

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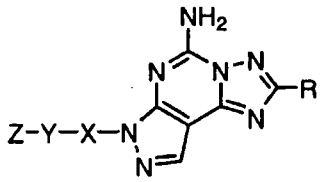
(81) Designated States *(national)*: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CO, CR, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, HR, HU, ID, IL, IN, IS, JP, KG, KR, KZ, LC, LK, LR, LT, LU, LV, MA, MD, MG, MK, MN, MX, MZ, NO, NZ, PL, PT, RO, RU, SE, SG, SI, SK, SL, TJ, TM,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

(84) Designated States *(regional)*: ARIPO patent (GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, 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, GW, ML, MR, NE, SN, TD, TG).

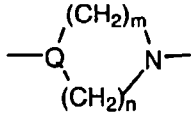
Declaration under Rule 4.17:
— *as to the applicant's entitlement to claim the priority of the earlier application (Rule 4.17(iii)) for all designations*

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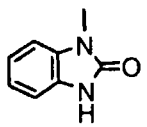
(54) Title: ADENOSINE A2A RECEPTOR ANTAGONISTS



(I)



(Ia)



(Ib)

(57) Abstract: Compounds having the structural formula (I) or a pharmaceutically acceptable salt thereof, wherein R is optionally substituted phenyl, cycloalkenyl, or heteroaryl; X is alkylene or $-C(O)CH_2-$;Y is $-N(R^2)CH_2CH_2N(R^3)-$, $-OCH_2CH_2N(R^2)-$, -O-, -S-, $-CH_2S-$, $-(CH_2)_2-NH-$, or optionally substituted,(Ia), m and n are 2-3, and Q is nitrogen or optionally substituted carbon; and Z is optionally substituted phenyl, phenylalkyl or heteroaryl, diphenylmethyl, $R^6-C(O)-$, R^6-SO_2-, $R^6-OC(O)-$, $R^7-N(R^8)-C(O)-$, $R^7-N(R^8)-C(S)-$, phenyl-CH(OH)-, or phenyl-$C(=NOR^2)-$; or when Q is CH, (Ib),phenylamino or pyridylamino; or Z and Y together are substituted piperidinyl or substituted phenyl; and R^2, R^3, R^6, R^7, and R^8 are as defined in the specification are disclosed, their use in the treatment of Parkinson's disease, alone or in combination with other agents for treating Parkinson's disease, and pharmaceutical compositions comprising them; also disclosed are a processes for preparing intermediates useful for preparing compounds of formula (I).

WO 01/92264 A1

Published:
— *with international search report*
— *before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments*

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