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INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification 7 :
C07D 405/12, A61K 31/33, A61P 31/12, C07D 213/40, 417/12, 231/56, 233/44, 285/06, 213/75, 307/52, 207/32, 333/20, 277/28, 209/14, 215/12

A1

(11) International Publication Number: **WO 00/34268**

(43) International Publication Date: 15 June 2000 (15.06.00)

(21) International Application Number: PCT/US99/28838

(22) International Filing Date: 6 December 1999 (06.12.99)

(30) Priority Data:
09/207,961 9 December 1998 (09.12.98) US

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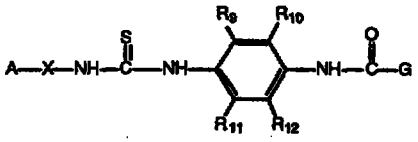
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(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, 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), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

Published
With international search report.
Before the expiration of the time limit for amending the claims and to be republished in the event of the receipt of amendments.

(54) Title: THIOUREA INHIBITORS OF HERPES VIRUSES



(I)

(57) Abstract

Compounds of formula (I) wherein A is heteroaryl; R_9–R_{12} are independently hydrogen, alkyl of 1 to 4 carbon atoms, perhaloalkyl of 1 to 4 carbon atoms, halogen, alkoxy of 1 to 4 carbon atoms, or cyano, or R_9 and R_{10} or R_{11} and R_{12} may be taken together to form aryl of 5 to 7 carbon atoms; W is O, NR_6, or is absent; G is aryl or heteroaryl; and X is a bond, –NH, alkyl of 1 to 6 carbon atoms, alkenyl of 1 to 6 carbon atoms, alkoxy of 1 to 6 carbon atoms, thioalkyl of 1 to 6 carbon atoms, alkylamino of 1 to 6 carbon atoms, or (CH)J; and J is alkyl of 1 to 6 carbon atoms, cycloalkyl of 3 to 7 carbon atoms, phenyl or benzyl; and n is an integer from 1 to 6; or a pharmaceutical salt thereof, useful in the treatment of diseases associated with herpes viruses including human cytomegalovirus, herpes simplex viruses, Epstein–Barr virus, varicella–zoster virus, human herpesviruses–6 and –7, and Kaposi herpesvirus.