

Title (en)
THIOUREA INHIBITORS OF HERPES VIRUSES

Title (de)
THIOUREA INHIBITOREN VON HERPESVIREN

Title (fr)
INHIBITEURS DE THIO-UREE DES VIRUS DE L'HERPES

Publication
EP 1140913 A1 20011010 (EN)

Application
EP 99965143 A 19991206

Priority
• US 9928892 W 19991206
• US 20854098 A 19981209

Abstract (en)
[origin: CA2351390A1] Compounds of formula (1) wherein, R1-R5 are independently selected from hydrogen, alkyl of 1 to 6 carbon atoms, alkenyl of 2 to 6 carbon atoms, alkynyl of 2 to 6 carbon atoms, perhaloalkyl of 1 to 6 carbon atoms, cycloalkyl of 3 to 10 carbon atoms, heterocycloalkyl of 3 to 10 carbon members, aryl, heteroaryl, halogen, -CN, -NO₂, -CO₂R₆, -COR₆, -OR₆, -SR₆, -SOR₆, -SO₂R₆, -CONR₇R₈, -NR₆N(R₇R₈), -N(R₇R₈) or W-Y-(CH₂)_n-Z; or R₂ and R₃ or R₃ and R₄, taken together from a 3 to 7 membered heterocycloalkyl or 3 to 7 membered heteroaryl; R₆ and R₇ are independently hydrogen, alkyl of 1 to 6 carbon atoms, perhaloalkyl of 1 to 6 carbon atoms, or aryl; R₈ is hydrogen, alkyl of 1 to 6 carbon atoms, perhaloalkyl of 1 to 6 carbon atoms, cycloalkyl of 3 to 10 carbon atoms, heterocycloalkyl of 3 to 10 members, aryl or heteroaryl, or R₇ and R₈, taken together may form a 3 to 7 membered heterocycloalkyl; A is heteroaryl; W is O, NR₆, or is absent; Y is -(CO)- or -(CO₂)-, or is absent; Z is alkyl of 1 to 4 carbon atoms, -CN, -CO₂R₆, COR₆, -CONR₇R₈, -OCOR₆, -NR₆COR₇, -OCONR₆, -OR₆, -SR₆, -SOR₆, -SO₂R₆, SR₆N(R₇R₈), -N(R₇R₈) or phenyl; G is aryl or heteroaryl; 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; 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; useful in the treatment of diseases associated with herpes viruses including human cytomegalovirus, herpes simplex viruses, Epstein-Barr virus, varicella-zoster virus, human herpes viruses -6 and -7, and Kaposi herpesvirus.

IPC 1-7
C07D 405/12; **C07D 417/12**; **C07D 213/81**; **C07D 213/75**; **A61K 31/33**; **A61P 31/12**

IPC 8 full level
A61K 31/12 (2006.01); **A61K 31/33** (2006.01); **A61K 31/425** (2006.01); **A61K 31/44** (2006.01); **A61K 31/4427** (2006.01); **C07D 213/81** (2006.01); **A61K 31/443** (2006.01); **A61K 31/4436** (2006.01); **A61K 31/444** (2006.01); **A61P 31/12** (2006.01); **A61P 31/22** (2006.01); **C07D 213/75** (2006.01); **C07D 401/12** (2006.01); **C07D 405/12** (2006.01); **C07D 417/12** (2006.01)

IPC 8 main group level
A61K (2006.01); **A61P** (2006.01); **C07D** (2006.01)

CPC (source: EP KR)
C07C 335/20 (2013.01 - KR); **C07C 335/22** (2013.01 - KR); **C07D 213/75** (2013.01 - EP); **C07D 213/81** (2013.01 - EP); **C07D 217/26** (2013.01 - KR); **C07D 307/81** (2013.01 - KR); **C07D 307/85** (2013.01 - KR); **C07D 405/12** (2013.01 - EP); **C07D 417/12** (2013.01 - EP)

Cited by
CN108642926A

Designated contracting state (EPC)
AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE

DOCDB simple family (publication)
AU 3112200 A 20000626; AU 756043 B2 20030102; BG 105580 A 20020131; BR 9916042 A 20011204; CA 2351390 A1 20000615; CN 1335843 A 20020213; CZ 20011958 A3 20011017; EA 200100637 A1 20011224; EP 1140913 A1 20011010; HU P0104758 A2 20020429; HU P0104758 A3 20020528; ID 29064 A 20010726; IL 143203 A0 20020421; JP 2002531558 A 20020924; KR 20010086082 A 20010907; NO 20012836 D0 20010608; NO 20012836 L 20010808; PL 348177 A1 20020506; SK 7692001 A3 20020910; TR 200101598 T2 20011022; ZA 200104373 B 20020918

DOCDB simple family (application)
AU 3112200 A 19991206; BG 10558001 A 20010608; BR 9916042 A 19991206; CA 2351390 A 19991206; CN 99815807 A 19991206; CZ 20011958 A 19991206; EA 200100637 A 19991206; EP 99965143 A 19991206; HU P0104758 A 19991206; ID 20011230 A 19991206; IL 14320399 A 19991206; JP 2000586716 A 19991206; KR 20017007051 A 20010605; NO 20012836 A 20010608; PL 34817799 A 19991206; SK 7692001 A 19991206; TR 200101598 T 19991206; ZA 200104373 A 20010528