

Title (en)

METALLOPROTEINASE INHIBITORS, PHARMACEUTICAL COMPOSITIONS CONTAINING THEM AND THEIR PHARMACEUTICAL USES

Title (de)

IHRE PHARMAZEUTISCHE ANWENDUNGENTHALTENDE PHARMAZEUTISCHE ZUSAMMENSETZUNGEN UND IHRE PHARMAZEUTISCHE ANWENDUNG

Title (fr)

INHIBITEURS DE METALLOPROTEINASES, COMPOSITIONS PHARMACEUTIQUES LES CONTENANT ET LEURS UTILISATIONS PHARMACEUTIQUES

Publication

EP 0973748 A1 20000126 (EN)

Application

EP 98914379 A 19980331

Priority

- US 9806365 W 19980331
- US 82531897 A 19970401

Abstract (en)

[origin: WO9843963A1] Compounds of formula (I), wherein Y is O or S, Ar is an aryl group or a heteroaryl group, R is H, an alkyl group, a cycloalkyl group, a heterocycloalkyl group, an aryl group, a heteroaryl group, or -C(O)R₁, wherein R₁ is hydrogen, an alkyl group, a cycloalkyl group, a heterocycloalkyl group, an aryl group, a heteroaryl group, or NR₂R₃ wherein R₂ and R₃ independently are hydrogen, an alkyl group, a cycloalkyl group, a heterocycloalkyl group, an aryl group, or a heteroaryl group, and X is -NH-OH or -OH. Pharmaceutically acceptable prodrugs, salts and solvates of these compounds. Methods of inhibiting the activity of metalloproteinases by administering a compound of the formula (I) or a prodrug, salt or solvate thereof. Pharmaceutical compositions comprising an effective amount of these compounds, prodrugs, salts, and solvates.

IPC 1-7

C07D 233/54; C07D 213/32; C07D 261/10; C07C 323/57; C07C 311/29; C07D 213/68; C07D 213/12; C07D 213/64; C07D 405/12; C07D 213/34; A61K 31/44; A61K 31/195; A61K 31/41

IPC 8 full level

A61K 31/18 (2006.01); A61K 31/198 (2006.01); A61K 31/216 (2006.01); A61K 31/4164 (2006.01); A61K 31/42 (2006.01); A61K 31/4406 (2006.01); A61K 31/4409 (2006.01); A61K 31/443 (2006.01); A61K 31/4439 (2006.01); A61P 1/02 (2006.01); A61P 9/10 (2006.01); A61P 13/00 (2006.01); A61P 17/00 (2006.01); A61P 19/02 (2006.01); A61P 19/10 (2006.01); A61P 25/28 (2006.01); A61P 27/02 (2006.01); A61P 29/00 (2006.01); A61P 35/00 (2006.01); A61P 35/04 (2006.01); A61P 43/00 (2006.01); C07C 311/29 (2006.01); C07C 323/52 (2006.01); C07C 323/59 (2006.01); C07C 323/60 (2006.01); C07D 213/32 (2006.01); C07D 213/34 (2006.01); C07D 213/643 (2006.01); C07D 213/68 (2006.01); C07D 213/70 (2006.01); C07D 233/54 (2006.01); C07D 261/08 (2006.01); C07D 405/12 (2006.01); C07D 413/12 (2006.01); C07D 213/64 (2006.01)

CPC (source: EP KR)

A61P 1/02 (2017.12 - EP); A61P 9/10 (2017.12 - EP); A61P 13/00 (2017.12 - EP); A61P 17/00 (2017.12 - EP); A61P 19/02 (2017.12 - EP); A61P 19/10 (2017.12 - EP); A61P 25/28 (2017.12 - EP); A61P 27/02 (2017.12 - EP); A61P 29/00 (2017.12 - EP); A61P 35/00 (2017.12 - EP); A61P 35/04 (2017.12 - EP); A61P 43/00 (2017.12 - EP); C07C 311/16 (2013.01 - KR); C07C 311/29 (2013.01 - EP); C07C 323/52 (2013.01 - EP); C07C 323/60 (2013.01 - EP); C07D 213/32 (2013.01 - EP); C07D 213/34 (2013.01 - EP); C07D 213/643 (2013.01 - EP); C07D 213/68 (2013.01 - EP); C07D 213/70 (2013.01 - EP); C07D 233/64 (2013.01 - EP); C07D 261/08 (2013.01 - EP); C07D 405/12 (2013.01 - EP); C07D 413/12 (2013.01 - EP)

Citation (search report)

See references of WO 9843963A1

Designated contracting state (EPC)

AT BE CH DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE

DOCDB simple family (publication)

WO 9843963 A1 19981008; AU 6874498 A 19981022; AU 747280 B2 20020509; BR 9809062 A 20000801; CA 2285372 A1 19981008; EP 0973748 A1 20000126; JP 2001521504 A 20011106; KR 20010005940 A 20010115; NZ 338082 A 20000623

DOCDB simple family (application)

US 9806365 W 19980331; AU 6874498 A 19980331; BR 9809062 A 19980331; CA 2285372 A 19980331; EP 98914379 A 19980331; JP 54194598 A 19980331; KR 19997009017 A 19991001; NZ 33808298 A 19980331