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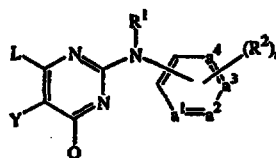
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(54) Title: HIV REPLICATION INHIBITING PYRIMIDINES



(I)

(57) Abstract

This invention concerns the use of compounds of formula (I), the *N*-oxides, the pharmaceutically acceptable addition salts, quaternary amines and the stereochemically isomeric forms thereof, wherein $-a^1=a^2=a^3=a^4-$ forms a phenyl, pyridinyl, pyrimidinyl, pyridazinyl or pyrazinyl with the attached vinyl group; *n* is 0 to 4; and where possible 5; *R*¹ is hydrogen, aryl, formyl, C₁-alkylcarbonyl, C₁-alkyl, C₁-alkyloxycarbonyl, substituted C₁-alkyl, or substituted C₁-alkyloxy; each *R*² independently is hydroxy, halo, optionally substituted C₁-alkyl, C₂-alkenyl or C₂-alkynyl, C₃-cycloalkyl, C₁-alkyloxy, C₁-alkyloxycarbonyl, carboxyl, cyano, nitro, amino, mono- or di(C₁-alkyl)amino, polyhalomethyl, polyhalomethoxy, polyhalomethylthio, $-S(=O)_pR^6$, $-NH-S(=O)_pR^6$, $-C(=O)R^6$, $-NHC(=O)H$, $-C(=O)NHNH_2$, $-NHC(=O)R^6$, $-C(=NH)R^6$ or a 5-membered heterocyclic ring; *p* is 1 or 2; *L* is optionally substituted C₁-10alkyl, C₂-10alkenyl, C₂-10alkynyl or C₃-7cycloalkyl; or *L* is $-X-R^3$ wherein *R*³ is optionally substituted phenyl, pyridinyl, pyrimidinyl, pyrazinyl or pyridazinyl; *X* is $-NR^1$, $-NH-NH$, $-N=N$, $-O$, $-C(=O)$, $-CHOH$, $-S$, $-S(=O)$ or $-S(=O)_2$; *Q* is hydrogen, C₁-alkyl, halo, polyhalo-C₁-alkyl or an optionally substituted amino group; *Y* represents hydroxy, halo, C₃-7cycloalkyl, optionally substituted C₁-alkyl, C₂-alkenyl or C₂-alkynyl, C₁-alkyloxy, C₁-alkyloxycarbonyl, carboxyl, cyano, nitro, amino, mono- or di(C₁-alkyl)amino, polyhalomethyl, polyhalomethoxy, polyhalomethylthio, $-S(=O)_pR^6$, $-NH-S(=O)_pR^6$, $-C(=O)R^6$, $-NHC(=O)H$, $-C(=O)NHNH_2$, $-NHC(=O)R^6$, $-C(=NH)R^6$ or aryl; aryl is optionally substituted phenyl; Het is an optionally substituted heterocyclic radical; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection.