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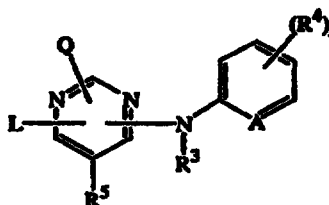
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(54) Title: **HIV INHIBITING PYRIMIDINE DERIVATIVES**

(57) Abstract

This invention concerns the use of the compounds of formula (I), the *N*-oxides, the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein A is CH, CR⁴ or N; n is 0 to 4; Q is hydrogen or -NR¹R²; R¹ and R² are selected from hydrogen, hydroxy, C₁-₁₂alkyl, C₁-₁₂alkyloxy, C₁-₁₂alkylcarbonyl, C₁-₁₂alkyloxycarbonyl, aryl, amino, mono- or di(C₁-₁₂alkyl)- amino, mono- or di(C₁-₁₂alkyl)aminocarbonyl wherein each C₁-₁₂alkyl may optionally be substituted; or R¹ and R² taken together may form pyrrolidinyl, piperidinyl, morpholinyl, azido or mono- or di(C₁-₁₂alkyl)aminoC₁-₄alkylidene; R³ is hydrogen, aryl, C₁-₆alkylcarbonyl, optionally substituted C₁-₆alkyl, C₁-₆alkyloxycarbonyl; and R⁴ is hydroxy, halo, optionally substituted C₁-₆alkyl, C₁-₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl, trihalomethyloxy; R⁵ is hydrogen or C₁-₄alkyl; L is optionally substituted C₁-₁₀alkyl, C₂-₁₀alkenyl, C₃-₁₀alkynyl, C₃-₇cycloalkyl; or L is -X¹R⁶ or -X²-Alk-R⁷ wherein R⁶ and R⁷ are optionally substituted phenyl; X¹ and X² are -NR³-, -NH-NH-, -N=N-, -O-, -S-, -S(=O)- or -S(=O)₂-; Alk is C₁-₄alkanediyl; aryl is optionally substituted phenyl; Het is an optionally substituted aliphatic or aromatic heterocyclic radical; for the manufacture of a medicine for the treatment of subjects suffering from HIV (Human Immunodeficiency Virus) infection. It further relates to new compounds being a subgroup of the compounds of formula (I), their preparation and compositions comprising them.



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