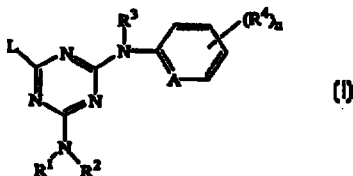




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(54) Title: **TRISUBSTITUTED 1,3,5-TRIAZINE DERIVATIVES FOR TREATMENT OF HIV INFECTIONS**

(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, 1, 2, 3 or 4; R¹ and R² are each independently 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 of the aforementioned C₁₋₁₂alkyl groups may optionally and each individually 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; and each R⁴ independently is hydroxy, halo, C₁₋₆alkyl, C₁₋₆alkyloxy, cyano, aminocarbonyl, nitro, amino, trihalomethyl or trihalomethyloxy; L is -X-R⁵ or -X-Alk-R⁶; wherein R⁵ and R⁶ each independently are indanyl, indolyl or phenyl; each of said indanyl, indolyl or phenyl may be substituted; and X is -NR³-, -NH-NH-, -N=N-, -O-, -S-, -S(=O)- or -S(=O)₂-; 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 pharmaceutical compositions comprising them.