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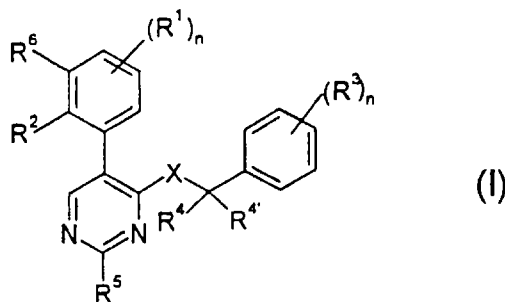
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(54) Title: 5-PHENYL-PYRIMIDINE DERIVATIVES



(57) Abstract: The invention relates to compounds of general formula (I) wherein R¹ is hydrogen or halogen; R² is hydrogen, halogen, lower alkyl or lower alkoxy; R³ is halogen, trifluoromethyl, lower alkoxy or lower alkyl; R⁴/R^{4'} are independently from each other hydrogen or lower alkyl; R⁵ is lower alkyl, lower alkoxy, amino, hydroxy, hydroxy-lower alkyl, -(CH₂)_n-piperazinyl, optionally substituted by lower alkyl, -(CH₂)_n-morpholinyl, -(CH₂)_{n+1}-imidazolyl, -O-(CH₂)_{n+1}-morpholinyl, -O-(CH₂)_{n+1}-piperidinyl, lower alkyl-sulfanyl, lower alkyl-sulfonyl, benzylamino, -NH-(CH₂)_{n+1}N(R^{4'})₂, -(CH₂)_n-NH-(CH₂)_{n+1}N(R^{4'})₂, -(CH₂)_{n+1}N(R^{4'})₂, or -O-(CH₂)_{n+1}N(R^{4'})₂, wherein R^{4'} is hydrogen or lower alkyl; R⁶ is hydrogen; R² and R⁶ or R¹ and R⁶ may be together with the

two carbon ring atoms -CH=CH-CH=CH-, with the proviso that n for R¹ is 1; n is independently 0 - 2; and X is -C(O)N(R^{4'})- or -N(R^{4'})C(O)-, and pharmaceutically acceptable acid addition salts thereof. Compounds of formula (I) have a high affinity to the NK-1 receptor. They are therefore useful for the treatment or diseases which relate to this receptor.



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