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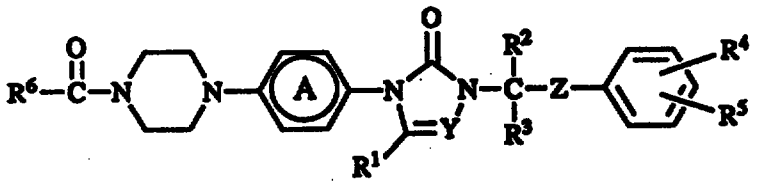
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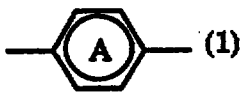
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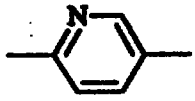
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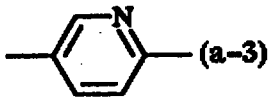
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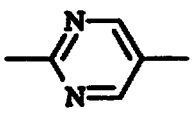
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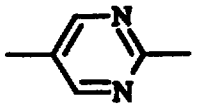
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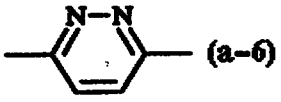
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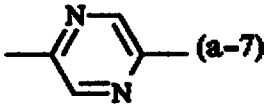
(a-4)



(a-5)



(a-6)



(a-7)

(57) Abstract

The invention is concerned with the compounds having formula (I), the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein Y is CH or N; R^1, R^2 and R^3 each independently are hydrogen or C_{1-4}alkyl; R^4 and R^5 each independently are hydrogen, halo, C_{1-4}alkyl, C_{1-4}alkyloxy, hydroxy, trifluoromethyl, trifluoromethyloxy or difluoromethyloxy; R^6 is hydrogen; or R^6 is phenyl optionally substituted with halo; pyridinyl; furanyl; thienyl; 3-chlorobenzo[b]thien-2-yl; trifluoromethyl; C_{1-4}alkyloxycarbonyl; dihalophenylcyclopropanyl; C_{3-6}cycloalkyl; adamantyl; C_{2-6}alkenyl optionally substituted with halophenyl; or C_{1-4}alkyl optionally substituted with halo, phenyl, halophenyl, dihalophenyl, phenyloxy, piperidinyl, pyrrolidinyl, piperazinyl, C_{1-4}alkylpiperazinyl, C_{1-4}alkyl-carbonylpiperazinyl, C_{1-4}alkyloxycarbonylpiperazinyl, phthalimino, amino, mono- or di(C_{1-20})alkylamino or C_{3-6}cycloalkylamino; Z is C=O or CHOH; and (1) is a radical of formula (a-1), (a-2), (a-3), (a-4), (a-5), (a-6), or (a-7). Compositions comprising said compounds, processes for preparing the same and the use of these compounds as a medicine.

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