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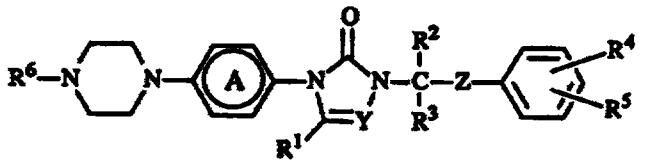
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(54) Title: ANTI-HELICOBACTER HETEROCYCLIC DERIVATIVES OF AZOLONES

(57) Abstract

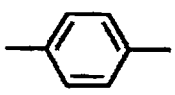
The invention is concerned with the compounds having formula (I), the pharmaceutically acceptable addition salts and the stereochemically isomeric forms thereof, wherein Y is CH or N; R^1, R^2 and R^3 each independently are hydrogen or C_{1-4}alkyl; R^4 and R^5 each independently are hydrogen, halo, C_{1-4}alkyl, C_{1-4}alkyloxy, hydroxy, trifluoromethyl, trifluoromethyloxy or difluoromethyloxy; R^6 is pyridinyl optionally substituted with up to two C_{1-4}alkyl groups; di(C_{1-4}alkyl)hydroxypyridinyl; di(C_{1-4}alkyl)C_{1-4}alkyloxypyridinyl; pyridazinyl optionally substituted with C_{1-4}alkyloxy; pyrimidinyl optionally substituted with hydroxy or C_{1-4}alkyloxy; thiazolyl optionally substituted with C_{1-4}alkyl; thiadiazolyl optionally substituted with C_{1-4}alkyl; benzoxazolyl or benzothiazolyl; or R^6 is pyrazinyl or pyridazinyl substituted with C_{1-4}alkyl; Z is C=O or CHOH; and (a) 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.



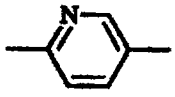
(I)



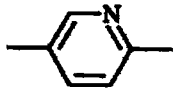
(a)



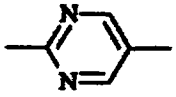
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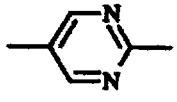
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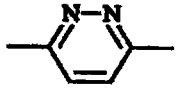
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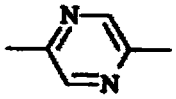
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