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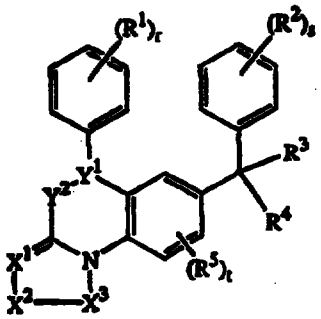
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(57) Abstract

This invention concerns compounds of formula (I), the pharmaceutically acceptable acid addition salts and the stereochemically isomeric forms thereof, wherein $=X^1–X^2–X^3–$ is a trivalent radical; $>Y^1–Y^2–$ is a trivalent radical; r and s are each independently 0, 1, 2, 3, 4 or 5; t is 0, 1, 2 or 3; each R^1 and R^2 are independently hydroxy, halo, cyano, C_{1-6}alkyl, trihalomethyl, trihalomethoxy, C_{2-6}alkenyl, C_{1-6}alkyloxy, hydroxyC_{1-6}alkyloxy, C_{1-6}alkylthio, C_{1-6}alkyloxyC_{1-6}alkyloxy, C_{1-6}alkyloxycarbonyl, aminoC_{1-6}alkyloxy, mono– or di(C_{1-6}alkyl)amino, mono– or di(C_{1-6}alkyl)aminoC_{1-6}alkyloxy, aryl, arylC_{1-6}alkyl, aryloxy or arylC_{1-6}alkyloxy, hydroxycarbonyl, C_{1-6}alkyloxycarbonyl; or two R^1 or R^2 on adjacent positions form together a bivalent radical; R^3 is hydrogen, halo, C_{1-6}alkyl, cyano, haloC_{1-6}alkyl, hydroxyC_{1-6}alkyl, cyanoC_{1-6}alkyl, aminoC_{1-6}alkyl, C_{1-6}alkyloxyC_{1-6}alkyl, C_{1-6}alkylthio–C_{1-6}alkyl, aminocarbonylC_{1-6}alkyl, hydroxycarbonyl, hydroxycarbonylC_{1-6}alkyl, C_{1-6}alkyloxycarbonylC_{1-6}alkyl, C_{1-6}alkylcarbonylC_{1-6}alkyl, C_{1-6}alkyloxycarbonyl, aryl, arylC_{1-6}alkyloxyC_{1-6}alkyl, mono– or di(C_{1-6}alkyl)aminoC_{1-6}alkyl, or a radical of formula $–O–R^{10}$, $–S–R^{10}$ or $–NR^{11}R^{12}$; R^4 is an optionally substituted imidazolyl; aryl is an optionally substituted phenyl or naphthalenyl; having farnesyl transferase and geranylgeranyl transferase inhibiting activity; their preparation, compositions containing them and their use as a medicine.



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