

Title (en)

CONDENSED PYRAZOLE DERIVATIVES AS PPAR AGONISTS II

Title (de)

KONDENSIERTE PYRAZOLDERIVATE ALS PPAR-AGONISTEN II

Title (fr)

DERIVES DE PYRAZOLE CONDENSES UTILISES COMME AGONISTES II DU RECEPTEUR PPAR

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Application

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Abstract (en)

[origin: WO2007036727A1] The invention discloses compounds of formula (I) wherein: R is a carboxylic acid or a derivative thereof; R₁</SUP> and R₂</SUP> are independently H or alkyl, or together R₁</SUP> and R₂</SUP> form an alkylene group; L</SUP> is a single bond, NH, NCH₂, O, S, CH₂ or CH, wherein when L</SUP> is CH the dashed line indicates a double bond and R</SUP> is absent, otherwise the dashed line is a single bond; R</SUP> and R</SUP> are independently H, halo, alkyl or alkoxy; L</SUP> is O or S; L</SUP> is CH₂ or CH₂>CH₂; R</SUP> and R</SUP> are independently H, halo, CF₃, OCF₃, alkyl or alkoxy; Y</SUP> is CH or N; Y</SUP> is CH or N; and R</SUP> and R</SUP> are independently H, halo, CF₃, OCF₃, alkyl or alkoxy; or a pharmaceutically acceptable derivative thereof, useful for treating disorders mediated by peroxisome-proliferator-activated receptor (PPAR) subtype d (PPARD). The compounds of the invention are therefore useful in the treatment of metabolic syndrome, obesity, type-II diabetes, dyslipidemia, wound healing, inflammation, neurodegenerative disorders and multiple sclerosis.

IPC 8 full level

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