

Title (en)
INTEGRIN ANTAGONISTS

Title (de)
INTEGRINANTAGONISTEN

Title (fr)
ANTAGONISTES D'INTEGRINE

Publication
EP 1392654 A1 20040303 (DE)

Application
EP 02742933 A 20020503

Priority
• DE 10127041 A 20010602
• EP 0204856 W 20020503

Abstract (en)
[origin: DE10127041A1] N-Substituted glycine N-(1-biphenyl-4-yl-2-(tetrazolyl or sulfonylaminocarbonyl)-ethyl)-amides (I) are new. Biphenyl derivatives of formula (I) and their stereoisomers, salts and solvates are new. A = NH₂, C(=NH)NH₂, NHC(=NH)NH₂, A'C(=NH)NH₂, Het1 or -NH-Het1 (where the primary amino groups are optionally conventionally protected); B' = tetrazolyl or RSO₂NHCO-; R = H, A', 6-14C cycloalkyl, 6-10C aryl or 7-14C aralkyl (optionally substituted by one or more R₃ and optionally having the alkyl chain interrupted by O); R₁, R'₁, R''₁, R₂, R'₂, R''₂ = H, halo, NO₂, NH₂, NHR, N(R)₂, OH, OR, COR, SO₃R, SO₂R or SR; R₃ = halo, NO₂, CF₃, OH, CN, OCF₃, SCF₃, OMe or OEt; Het1 = mono- or bicyclic heterocycle containing 1-4 N, optionally substituted by 1 or 2 of A', NHA', N(A')₂ and/or NH₂; A' = 1-8C alkyl; X = direct bond, O, NH or CH₂; n = 2-4. An Independent claim is also included for the preparation of (I).

IPC 1-7
C07D 213/74; C07D 401/12; A61K 31/41; A61K 31/435; A61P 9/00; A61P 35/00

IPC 8 full level
A61P 9/00 (2006.01); **A61P 35/00** (2006.01); **C07D 213/74** (2006.01); **C07D 401/12** (2006.01)

CPC (source: EP US)
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