

Europäisches Patentamt

European Patent Office

Office européen des brevets



(11) **EP 1 375 499 A8**

(12)

CORRECTED EUROPEAN PATENT APPLICATION

Note: Bibliography reflects the latest situation

(15) Correction information:

Corrected version no 1 (W1 A1)

INID code(s) 84

(51) Int Cl.⁷: **C07D 413/12**, C07D 263/32, C07C 233/47

(48) Corrigendum issued on: **08.09.2004 Bulletin 2004/37**

(43) Date of publication: **02.01.2004 Bulletin 2004/01**

(21) Application number: 03020537.1

(22) Date of filing: 19.03.1999

(84) Designated Contracting States:

AT BE CH CY DE DK ES FI FR GB GR IE IT LI LU MC NL PT SE

Designated Extension States:

AL LT LV MK RO SI

(30) Priority: 30.03.1998 JP 10409898

(62) Document number(s) of the earlier application(s) in accordance with Art. 76 EPC: 99909293.5 / 0 992 503

(71) Applicant: Japan Tobacco Inc. Tokyo 105-8422 (JP)

(72) Inventors:

 Ando, Koji, 1-1, Murasaki-cho Takatsuki-shi Osaka 569-1125 (JP) Suzuki, Masanobu Tokyo 184-0004 (JP)

(74) Representative:

von Kreisler, Alek, Dipl.-Chem. et al Patentanwälte, von Kreisler-Selting-Werner, Bahnhofsvorplatz 1 (Deichmannhaus) 50667 Köln (DE)

Remarks:

This application was filed on 17-09 - 2003 as a divisional application to the application mentioned under INID code 62.

- (54) Process for the production of 2-(5-methyl-4-oxazolyl)ethanol derivatives
- (57) The present invention relates to a novel method for producing a compound of the formula [11]

wherein R is an optionally substituted aromatic hydrocarbon group, an optionally substituted alicyclic hydrocarbon group, an optionally substituted heterocyclic group or an optionally substituted condensed heterocyclic group, which is useful as a therapeutic agent for diabetes. The method of the present invention is an industrially utilizable method that enables efficient production of the objective compound [11] from β -methyl L-aspartate via an important intermediate compound

wherein R is as defined above, at high yield.