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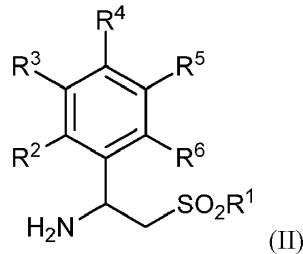
(54)	Title	<b>PROCESSES FOR THE PREPARATION OF (S)-1-(3-ETHOXY-4METHOXYPHENYL)-2-METHANESULFONYLETHYLAMINE</b>
(56)	References Cited:	WO-A2-2010/030345 US-A1- 2010 324 108 FRANCISCO VELÁZQUEZ ET AL: "Steroselective synthesis of beta-substituted beta-amino sulfones and sulfonamides via addition of sulfonyl anions to chiral N-sulfinyl imines", ORGANIC LETTERS , 14(23), 6012-6015 CODEN: ORLEF7; ISSN: 1523-7052, , vol. 8, no. 4 1 January 2006 (2006-01-01), pages 789-792, XP002712270, ISSN: 1523-7060, DOI: 10.1021/OL053132B

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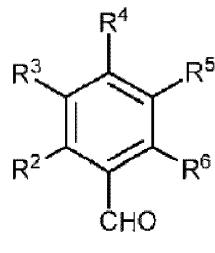
Enclosed is a translation of the patent claims in Norwegian. Please note that as per the Norwegian Patents Acts, section 66i the patent will receive protection in Norway only as far as there is agreement between the translation and the language of the application/patent granted at the EPO. In matters concerning the validity of the patent, language of the application/patent granted at the EPO will be used as the basis for the decision. The patent documents published by the EPO are available through Espacenet (<http://worldwide.espacenet.com>) or via the search engine on our website here: <https://search.patentstyret.no/>

**Patentkrav**

**1.** Fremgangsmåte for å fremstille en forbindelse med formel (II):



- 5 eller et farmasøytisk akseptabelt salt, hydrat, solvat eller polymorf derav, som omfatter:  
 (a) kondensere et kiralt hjelpestoff med et aldehyd som har formelen



- ,  
 hvor det kirale hjelpestoffet er (R)-(+)-tertiært butylsulfonamid eller  
 10 (S)- $\alpha$ -metylbenzylamin;  
 (b) tilsette et nukleofil til det kondenserte produktet, hvori nukleofilet er litiumanionet av dimethylsulfon; og  
 (c) avbeskytte tilleggsproduktet,  
 hvor:  
 15 R<sup>1</sup> er CH<sub>3</sub>; og  
 hver av R<sup>2</sup>, R<sup>3</sup>, R<sup>4</sup>, R<sup>5</sup> og R<sup>6</sup> ved hver forekomst er uavhengig hydrogen, halogen, C<sub>1</sub>-C<sub>6</sub>alkyl, C<sub>1</sub>-C<sub>6</sub>alkoksy, -CF<sub>3</sub>, -CN eller -NO<sub>2</sub>.

- 2.** Fremgangsmåten ifølge krav 1, hvori R<sup>1</sup> er -CH<sub>3</sub>; R<sup>2</sup> er H; R<sup>3</sup> er H; R<sup>4</sup> er -OCH<sub>3</sub>; R<sup>5</sup> er -OCH<sub>2</sub>CH<sub>3</sub>; og R<sup>6</sup> er H.

- 3.** Fremgangsmåten ifølge krav 1, hvori tilsetningen av et nukleofil skjer under basisk tilstand.

- 25 4.** Fremgangsmåten ifølge krav 1, hvori avbeskyttelse av tilsetningsproduktet skjer under sur tilstand.

- 5.** Fremgangsmåten ifølge krav 1, hvori aldehydet er 3-etoksy-4-metoksybenzaldehyd.

**6.** Fremgangsmåten ifølge krav 1, hvori det kirale hjelpestoffet er (R)-(+)-tertiært butylsulfinamid.

**7.** Fremgangsmåten ifølge krav 1, hvori det kirale hjelpestoffet er  
5 (S)- $\alpha$ -metylbenzylamin.

**8.** Fremgangsmåten ifølge krav 1, hvori aldehydet er 3-etoksy-4-metoksybenzaldehyd, det kirale hjelpestoffet er (R)-(+)-tertiært butylsulfinamid eller (S)- $\alpha$ -metylbenzylamin, og nukleofilet er litiumanionet av dimethylsulfon.