

Title (en)

SEMI-SYNTHETIC ROUTE FOR THE PREPARATION OF PACLITAXEL, DOCETAXEL AND 10-DEACETYLBACCATIN III FROM 9-DIHYDRO-13-ACETYLBACCATIN III

Title (de)

HALBSYNTHETISCHE ROUTE ZUR HERSTELLUNG VON PACLITAXEL, DOCETAXEL UND 10-DEACETYLBACCATIN III AUS 9-DIHYDRO-13-ACETYLBACCATIN III

Title (fr)

VOIE SEMI-SYNTHÉTIQUE POUR LA PRÉPARATION DE PACLITAXEL, DOCÉTAXEL ET 10-DÉSACÉTYLBACCATINE III À PARTIR DE 9-DIHYDRO-13-ACÉTYLBACCATINE III

Publication

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Application

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Abstract (en)

[origin: WO2007143839A1] A novel semi-synthetic route has been provided in the preparation of docetaxel and paclitaxel. This new process involves the conversion of 9-dihydro-13-acetylbaaccatinIII to docetaxel and paclitaxel by the step of converting 9-dihydro-13-acetylbaaccatin III into 7-O-triethylsilyl-9,10-diketobaccatin III, and adding docetaxel and paclitaxel side chain precursors, respectively, to form a new class of taxane intermediates, such as 7-O-triethylsilyl-9,10-diketodocetaxel and 7-O-triethylsilyl-9,10-diketopaclitaxel. These new intermediates then by a series reduction, acetylation of the 10-hydroxyl position for paclitaxel and finally deprotection to yield docetaxel and paclitaxel, the most important anti-cancer drugs.

IPC 8 full level

C07D 305/14 (2006.01); **C07F 7/18** (2006.01)

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Citation (search report)

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- [X] APPENDINO G ET AL: "The Reductive Fragmentation of 7-Hydroxy-9,10-dioxotaxoids", EUROPEAN JOURNAL OF ORGANIC CHEMISTRY, WILEY-VCH VERLAG, WEINHEIM, DE, no. 22, 1 January 2003 (2003-01-01), pages 4422 - 4431, XP002486097, ISSN: 1434-193X
- See references of WO 2007143839A1

Citation (examination)

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