

(ェ)-Carquinostatin A の主日成こ (ハ)-(-)- 及の (*S*)-(+)-carquinostatin A の不斉全合成研究

稗田雄三、町支臣成、内田佳成、藤岡晴人、藤井さゆり、日比野俐

Chem. Pharm. Bull., 60, 1522-1530 (2012)

Total synthesis of (±)-Carquinostatin A, and Asymmetric Total Synthesis of (R)-(-)-Carquinostatin A and (S)-(+)-Carquinostatin A

Yuhzo Hieda, Tominari Choshi, Yoshinari Uchida, Haruto Fujioka, Sayuri Fujii, and Satoshi Hibino

ABSTRACT: Total syntheses of (\pm)-carquinostatin A, and (R)-(-)-carquinostatin A together with its enanchiomer, (S)-(+)-carquinostatin A, possessing radical scavenging activity, were newly achieved. (\pm)-Carquinostatin A was synthesized from 1-acetonyl-6-bromo-3-ethoxy-2-methylcarbazole. which was derived from the known 1-axetonyl-3-ethoxy-2-methylcarbazole. Introduction of a prenyl groupat the 6-position of caebazole was successful in two steps. For the synthesis of (R)-(-)-carquinostatin A and (S)-(+)-carquinostatin A, (R)-(-)-1-(2-acetoxypropyl)-3-hydroxy-2-methoxycarbazole and (S)-(+)-3-hydroxy-1-(2-hydroxypropyl)-2-methylcarbazole, prepared by Lipase-QLM catalyzed enantioselective transesterification of 3-hydroxy-1-(2-hydroxypropyl)-2-methylcarbazole, were used as chiral starting materials.

抄録 Carquinostatin A は、瀨戸らにより *Streptomyces exfoliates* 2419-SVT2 から単離され、抗酸化作用を有する多置換カルバゾールアルカロイドである。(±)-Carquinostatin A は7工程で全合成を達成した。(*R*)-(¬)-carquinostatin A 及び(*S*)-(+)-carquinostatin A のエナンチオ選択的全合成は、Lipase-QLM による 3-hydroxy-1-(2-hydroxypropyl)-2-methylcarbazole のエナンチオ選択的エステル交換反応により対応する (*R*)-(¬)-1-(2-acetoxypropyl)-3-hydroxy-2-methoxycarbazole 及び(*S*)-(+)-3-hydroxy-1-(2-hydroxypropyl)-2-methylcarbazole を得ることにより達成した。