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Study on the structures and biosynthetic pathway of indole alkaloids, notoamides

We isolated new alkaloids, notoamides, from a marine-derived *Aspergillus* sp., which was isolated from the

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Diels-Alder (IMDA) reaction between the double bond in the isoprenyl substituent and the diene moiety in the azadiene ring derived from the diketopiperazine ring. However, the feeding experiment of <sup>13</sup>C-labeled 3 did not afford 1 and 2. Further, Williams and Gloer *et al.* reported the isolation of antipodes of 1 and 2 from the terrestrial *Aspergillus versicolor*. Both enantitomers of 1 and 2 were shown to be optically pure. With regard to the generation of Both enantitomers of 1 and 2, the IMDA reaction would occur in a face-selective manner in both *Aspergillus* sp. In

addtion, an enatiomeric pair of versicolamide B (4) was also isolated from both Aspergills. The biosynthetic pathway

common mussel, *Mytilus edulis galloprovincialis*. Among them, notoamide B (1) and stephacidin A (2) contain a bicyclo[2.2.2]diazaoctane ring. They are proposed to be constructed from notoamide E (3) by the intra-molecular

of notoamide congeners is an interesting subject of current investigation.

Angew. Chem. Chem. Int.

Ed. 2007, 46, 2254, 2257,
2262; J. Am. Chem. Soc.

2009, 131, 3834; Org.

Lett. 2009, 11, 1297; Nat.

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