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Mohamed Osama Ibrahim Mohamed

Graduate School of Pharmaceutical Sciences,
Tohoku University, 6-3 Aza-aoba, Aramaki,
Aoba-ku, Sendai 980-8578, Japan

Total synthesis of the antifungal clinical candidate ASP2397 cyclohexapeptide facilitated by the chemo-selective reduction of secondary *N*-benzyloxyamides

Mohamed Osama Ibrahim Mohamed, Yutong Du, Kosuke Ohsawa, Atsushi Tahara, Yoshitaka Shimotai, Hiroshi Hamamoto, A Ganesan, Takayuki Doi*

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