

Synthesis of a Regular 24-membered Cyclodepsipeptide by Direct Amide Cyclization

Peter Köttgen*, Anthony Linden, and Heinz Heimgartner

Organisch-chemisches Institut der Universität Zürich, Winterthurerstrasse 190, 8057 Zürich, Switzerland

Reprint requests to Prof. H. Heimgartner. E-mail: heimgart@oci.uzh.ch

Z. Naturforsch. **2009**, *64b*, 689 – 698; received March 30, 2009

Dedicated to Professor Gerhard Maas on the occasion of his 60th birthday

The synthesis of a 24-membered cyclic depsipeptide with an alternating sequence of phenyl-lactic acid and α -aminoisobutyric acid (Aib) is described. The linear precursor was prepared *via* the ‘azirine/oxazolone method’ using 2,2-dimethyl-3-amino-2*H*-azirines as building blocks for the α,α -disubstituted α -amino acid Aib. The macrolactonization leading to the cyclodepsipeptide was achieved by the ‘direct amide cyclization’, *i. e.*, by treatment of a solution of the linear precursor in toluene with HCl gas.

Key words: Cyclodepsipeptides, Direct Amide Cyclization, Azirine/Oxazolone Method, Ainoisobutyric Acid, Crystal Structure