

Isoxazolinium Salts in Asymmetric Synthesis.

1. Stereoselective Reduction Induced by a 3'-Alkoxy Stereocentre. A New Approach to Polyfunctionalized β -Amino Acids* [1, 2]

Marco Henneböhle, Pierre-Yves Le Roy, Matthias Hein, Rudolf Ehrler, and Volker Jäger

Institut für Organische Chemie, Universität Stuttgart, Pfaffenwaldring 55,
D-70569 Stuttgart, Germany

Reprint requests to Prof. Dr. Volker Jäger. Fax: +49(0)711-6854321.
E-mail: jager.ioc@po.uni-stuttgart.de

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Dedicated to Professor Ulrich Schmidt on the occasion of his 80th birthday

A new approach to optically active *N*-methylamino acids is presented, relying on stereoselective reduction of *N*-methylisoxazolinium salts with a dioxyethyl side-chain. The diastereoselectivity of the reduction step is studied systematically, in comparison with that of respective isoxazolines. A two-step transformation of isoxazolinium salts – with NaBH₃(OAc) and subsequent catalytic hydrogenation as well as a one-pot reduction by catalytic hydrogenation led to high (95:5 and 87:13) diastereomeric ratios of protected *erythro-N*-methylaminopentane-1,2,3-triols. The hydroxyethyl side-chain is elaborated by oxidation to afford the β -*N*-methylamino acid **37**, exemplifying the potential of this strategy.

Key words: Isoxazolinium Salts, Methylamino Alcohols, Diastereoselective Reduction,
Homoserine, β -Amino Acids