Methylaluminium bis(4-substituted-2,6-di-tert-butylphenoxide): MAD and MABR Lewis acids

Compiled by Cyril Papamicaël

The Dyson Perrins Laboratory, University of Oxford, South Parks Road, Oxford, OX1 3QY, UK

Cyril Papamicaël obtained his PhD under the guidance of Professor G. Quéguiner and G. Dupas at the University of Rouen (France) in 1997. He then spent two years working with Professor A. Gossauer at the University of Fribourg (Switzerland). He is currently engaged in a postdoctoral work with Professor C.J. Schofield at the University of Oxford (United-Kingdom).

Abstracts

(A) In the presence of the MAD Lewis acid, which complexes to the oxygen atom of the sulfoxide group, the radical cyclization/β-elimination tandem reaction leads to the R enantiomer 4. Without the MAD reagent, the S enantiomer is isolated (93% yield, >96% ee).²

(B) A general method of six-membered carbocycles has been demonstrated, which involves the stereo-controlled cyclization of olefinic epoxides with MABR via the epoxide rearrangement and subsequent intramolecular ene reaction with high stereoselectivity.⁷ Treatment of 5 with MABR afforded trans-decalin-1-ol 6 as a sole isolable product with rigorous stereochemistry in 70% yield.

(C) The MAD ¹ or MABR ² can be successfully utilised as a highly efficient nonchelating Lewis acid for achieving high stereoselectivity in 1,n asymmetric induction in cyclic as well as acyclic systems. Thus, Diels-Alder reaction of the acrylate of D-pantolactone and cyclopentadiene exhibits high diastereoselectivity not observable with ordinary Lewis acids.³

References and Notes


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