

## Angewandte Corrigendum

Upon further examination of the NMR spectra for the products of the reduction of the Se-phenyl 1-methoxybicyclo[2.2.2]oct-5-ene-2-carboselenoate 5 f the authors of this Communication have found that the reaction did not give the expected 1-methoxybicyclo-[2.2.2]oct-2-ene 6 f in 67% yield as reported in Table 1 (entry f). Instead, three major products were obtained in a combined yield of 67%: the expected product 6f (16%) along with the two products of cyclopropylcarbinyl radical rearrangement, the endo (major, 57%) and the exo (minor, 27%) isomers of 2-methoxybicyclo[3.2.1]oct-6ene, the former being a known compound.[1]

[1] a) M. A. Battiste, J. M. Coxon, A. J. Jones, R. W. King, G. W. Simpson, P. J. Steel, Tetrahedron Lett. 1983, 24, 307-310; b) M. A. Battiste, J. M. Coxon, G. W. Simpson, P. J. Steel, A. J. Jones, Tetrahedron 1984, 40, 3137-44.

Se-Phenyl Prop-2-eneselenoate: An Ethylene Equivalent for Diels-Alder Reactions

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