

Stereospecific Reaction of Olefins with Sulphur Trioxide—Dioxan

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Summary The reaction of *cis*- and *trans*-but-2-ene with sulphur trioxide–dioxan proceeds stereospecifically yielding *cis*- and *trans*-2,3-butanedisulfone, respectively, as initial sulphonation products.

SULPHONATION of olefins with sulphur trioxide has been extensively investigated,¹ but no stereochemical study on the reaction has been reported. We now report the stereochemistry of the initial products of the sulphonation reaction between *cis*- or *trans*-but-2-ene and sulphur trioxide–dioxan.

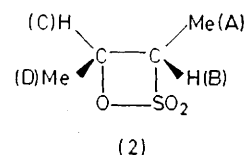
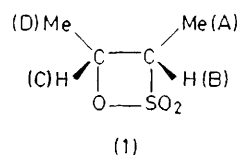
An equimolar amount of sulphur trioxide–dioxan (1:1) was added to *cis*- or *trans*-but-2-ene in chloroform to give the disulfones (1) and (2); (1) δ 1.5(q), 4.7–5.1(m), J_{AB} 7.9, J_{BC} 8.0, J_{CD} 6.0 Hz; (2) δ 1.5(q), 4.2–4.6(m), J_{AB} 6.8, J_{BC} 5.4, J_{CD} 6.8 Hz.

These results indicate that the butane-2,3-disulfones were produced stereospecifically from the corresponding but-2-ene isomers.†

† Satisfactory analytical data were obtained.

¹ For leading references see, e.g. F. G. Bordwell and M. L. Peterson, *J. Amer. Chem. Soc.*, 1959, **81**, 2002; I. L. Knunyants and G. A. Sokolski, *Angew. Chem. Internat. Edn.*, 1972, **11**, 583.

Similarly, sulphonation of *cis*- and *trans*-pent-2-ene with sulphur trioxide–dioxan gave the corresponding 2,3-pentanedisulfones.



Addition of an equimolar amount of sulphur trioxide alone to *cis*- or *trans*-but-2-ene in chloroform at 0° also resulted in stereospecific formation of *cis*- and *trans*-2,3-butanedisulfone.

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