

Isomerisation of Epoxides to Carbonyl Compounds induced by Iodides in Dimethyl Sulphoxide

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A NOVEL isomerisation of epoxides to carbonyl compounds by methyl iodide and sodium iodide in dimethylformamide was described earlier.¹ A study of the analogous reaction in dimethyl sulphoxide has now revealed features of its mechanism, linking it with other reactions involving activated dimethyl sulphoxide.²

In a typical experiment an approximately 0.07M-solution of the epoxide in dimethyl sulphoxide containing n-propyl iodide (0.35M) and sodium iodide (0.35M) is kept at 80° for 3 hr. The dark reaction mixture contains iodine (which is not catalytically active) but, after aqueous work-up, it affords the carbonyl compound(s) in high yield; thus octan-4-one (100%) is obtained from *cis*-4,5-epoxyoctane, octanal (80%) and octan-2-one (4%) from 1,2-epoxyoctane, phenylacetaldehyde (85%) and acetophenone (4%) from styrene oxide, and cyclohexanone (90%) from epoxycyclohexane.

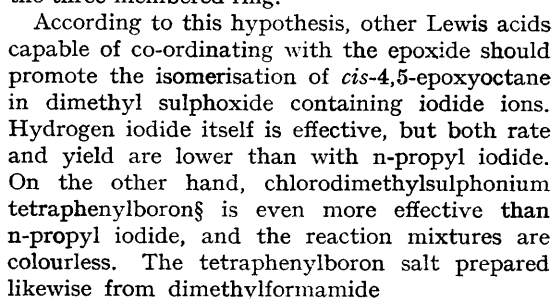
The kinetics of isomerisation of *cis*-4,5-epoxyoctane at 70° have been studied; 2% water was added to eliminate rate variations from imperfect dryness, but it has little effect on the yield of the reaction at this level. Reaction mixtures were partitioned between chloroform and water and the

organic components analysed by gas chromatography using a 100 ft. stainless-steel capillary column with 18% dinonyl phthalate as stationary phase and diglyme as internal standard. The rate of disappearance of the epoxide is substantially faster than the rate of appearance of octan-4-one, indicating the formation of a metastable intermediate which is not detectable by these techniques. Over the first 50% of reaction, the rate of disappearance of the epoxide is proportional to the epoxide concentration,[†] and it increases with increasing concentration of n-propyl iodide, though not proportionately; it is independent of the concentration of sodium iodide. However, sodium iodide has a marked accelerating effect on the production of octan-4-one, and, indeed, terminal yields of ketone fall significantly when no sodium iodide is added to the reaction mixture.

Alkyl iodides are known to alkylate dimethyl sulphoxide either on oxygen to form alkoxydimethylsulphonium iodides or on sulphur to form alkoxydimethylsulphonium iodides.^{3,4} Trimethyloxosulphonium iodide⁵ is ineffective in isomerising epoxides under conditions where methyl iodide gives a smooth conversion. On the other hand,

[†] In the later stages of the disappearance of the epoxide the rate increases, an effect which appears to be related to the formation of appreciable amounts of octan-4-one.

not detectable by our analytical method. Significantly, this iodohydrin, prepared separately, yields octan-4-one when treated with sodium iodide in dimethyl sulphoxide at 70°. The direction of isomerisation of the unsymmetrical epoxides 1,2-epoxyoctane and styrene oxide is explained, since the major products are analogous to those expected from acid-catalysed opening of the three-membered ring.



(i.e., $\text{Cl}\cdot\text{CH}:\overset{+}{\text{N}}\text{Me}_2 \overset{-}{\text{BPh}_4}$) is also a very effective

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$$\begin{array}{c}
 \text{PrI} + \text{Me}_2\text{SO} \rightleftharpoons \text{Me}_2\text{S}^+\text{OPr} + \text{I}^- \\
 \\
 \text{Me}_2\text{S}^+\text{OPr} + \text{PrCH}(\text{O})\text{CHPr} \rightleftharpoons \text{Me}_2\text{SOPr}-\text{O}^+-\text{C}(\text{Pr})(\text{H})-\text{CH}(\text{Pr}) \quad (\text{I}) \\
 \downarrow \text{I}^- \quad \text{slow} \\
 \text{Me}_2\text{SOPr}-\text{O}-\text{CH}(\text{Pr})-\text{CH}(\text{Pr})-\text{I} \quad (\text{II}) \\
 \downarrow \text{I}^- \quad \text{slow} \\
 \text{Me}_2\text{SOPr}-\text{O}-\text{CH}(\text{Pr})-\text{CH}(\text{Pr})^+ \quad (\text{III}) \\
 \downarrow \quad \downarrow \\
 \text{Me}_2\text{S}^+\text{OPr} + \text{PrCO}-\text{CH}_2\text{Pr}
 \end{array}$$

§ New compound, m.p. 108°, prepared from phosgene and dimethyl sulphoxide with subsequent addition of sodium tetraphenylboron in aqueous methanol at 0°.

catalyst, and this supports our initial suggestion that the reactions in dimethylformamide and dimethyl sulphoxide are essentially similar. Alkyl iodides and sodium iodide do *not* significantly

catalyse the isomerisation in sulpholane, acetonitrile, or butan-2-one.

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¹ G. W. Kenner and E. Stenhagen, *Acta Chem. Scand.*, **1964**, **18**, 1551.

² See for example, A. H. Fenselau and J. G. Moffatt, *J. Amer. Chem. Soc.*, **1966**, **88**, 1762; W. W. Epstein and K. W. Sweat, *Chem. Rev.*, **1967**, **67**, 247; J. D. Albright and L. Goldman, *J. Amer. Chem. Soc.*, **1967**, **89**, 2415.

³ S. G. Smith and S. Winstein, *Tetrahedron*, **1958**, **3**, 317.

⁴ R. O. Pick, Thesis, University of California at Los Angeles, 1967.

⁵ R. Kuhn and H. Trischmann, *Annalen*, **1958**, **611**, 117.

⁶ K. Torssell, *Tetrahedron Letters*, **1966**, 4445; *Acta Chem. Scand.*, **1967**, **21**, 1.