ABSTRACTS

Studies on Lactone Formation in Vapor Phase. (II)

Synthesis of *E*-Caprolactone

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Dehydrogenation of 1,6-hexanediol to ε -caprolactone with numerous mixed catalysts has been studied. Attempts to dehydrogenate 1,6-hexanediol with copper-zinc oxide or copper-magnesium oxide catalyst, which gave γ -butyrolactone and δ -valerolactone in good yields, were unsuccessful. The most favorable catalyst giving ε -caprolactone in 68% yield, had the composition of CuO: Cr₂O₃: ZnO= 2: 0.5: 1. The catalysts which contained one of the components excessively lowered both the activity and selectivity. Preparation of 1,6-hexanediol, the starting material for ε -caprolactone, was performed as follows, and the raction conditions of each step have been examined.

- 1) $CH \equiv CH + HCHO \longrightarrow CH \equiv CCH_2OH$
- 2) $2CH \equiv CCH_2OH + 1/2O_2 \longrightarrow HOCH_2C \equiv CC \equiv CCH_2OH + H_2O$
- 3) $HOCH_2C \equiv CC \equiv CCH_2OH + 4H_2 \longrightarrow HO(CH_2)_6OH$

Studies on Lactone Formation in Vapor Phase. (III)

Mechanism of Lactone Formation from Diols

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To clarify the mechanism of lactone formation from diols, the behavior of dialdehydes or oxyaldehydes over the copper catalysts has been studied. With all the catalysts used, it was ascertained that the oxyaldehydes gave the corresponding lactones as easily as the diols did. On the contorary, the dialdehydes examined gave no lactones, and they were recovered up to 70% except in the case of adipaldehyde. From these experimental results, it is concluded that this reaction does not take place by an internal Cannizzaro type reaction of dialdehydes but by an intramolecular acetalization of oxyaldehydes, followed by a dehydrogenation to lactones.