Catalysis: Autoxidation in the Presence of Active Cobalt Oxide

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Summary An active oxide of cobalt is an effective autoxidation catalyst towards organic substrates.

HETEROGENEOUS oxidations of benzyl alcohol and related compounds occur with active oxides of manganese¹ and nickel² prepared by ozonisation. We have now found that a black oxide of cobalt prepared in a similar manner is a good autoxidation catalyst.

There are numerous examples³⁻⁵ of cobalt salts acting as homogeneous oxidation catalysts. A recent patent⁶ describes the oxidation of n-butanol to butyric acid with a mixture of sodium hypochlorite, sodium hydroxide, and cobalt(II) chloride. The patent describes the substitution of a cobalt oxide, prepared by the reaction of cobalt(II) chloride and sodium hypochlorite, for cobalt(II) chloride in the procedure, presumably, leading to a heterogeneous reaction. The black cobalt oxide, prepared by ozonisation, when used with oxygen and an aprotic solvent behaves as manganese dioxide and nickel peroxide in the oxidation of benzylic alcohols, active methylene groups, aniline, and benzylamine. All substrates gave the expected products except 2-phenylethanol from which benzaldehyde was obtained. This last reaction involved carbon to carbon bond cleavage which may have proceeded *via* abstraction

TABLE 1

Reactant	Ratio catalyst : substrate	Time (h)	Product	Yield (%)
Benzyl alcohol	7:1	0.2	Benzaldehyde Benzoic acid	81a
Cinnamyl alcohol	2:1	0.2	Cinnamaldehyde	94a
2-Phenylethanol	10:1	6.0	Benzaldehyde	68ª
Diphenylmethanol	7:1	0.2	Benzophenone	91a
Fluorene	1:1	1.5	Fluorenone	99
Diphenylmethane	6:1	6.0	Benzophenone	998
4-Methylcyclohexanol	10:1	6.0	4-Methylcyclohexanone	10b
Aniline	7:1	6.0	Azobenzene	62
Benzylamine	7:1	0.2	Benzonitrile	85

^a Determined as 2,4-dinitrophenylhydrazone. ^bDetermined by g.l.c.

of a benzylic hydrogen or a terminal hydrogen atom followed by rearrangement in a similar manner to that postulated for the oxidation of carboxylic acids with cobalt(III) perchlorate.7

All reactions reported in Table 1 were conducted in dry benzene at 80° and oxygen was passed through at a rate of approximately 45 cm³/min. The concentration of substrate in all cases was about 200 mg in 30 cm³ of dry benzene.

Benzyl alcohol was studied at various catalyst : substrate ratios with and without added oxygen. It is seen in Table 2 that yields were significantly lower when oxygen was not added, and no improvement was observed with increased catalyst ratio.

The cobalt oxide was prepared by addition of 7n-sodium hydroxide (200 cm³) dropwise during 4 h to a stirred solution of cobalt(II) sulphate heptahydrate (281 g) in water $(1 \cdot 2 1)$ while an ozone-oxygen mixture was passed

¹ J. S. Belew and Chwang Tek-Ling, Chem. and Ind., 1967, 1958.

² J. S. Belew and Chwang Tek-Ling, *Chem. Comm.*, 1967, 1100. ³ A. S. Hay and H. S. Blanchard, *Canad. J. Chem.*, 1965, **43**, 1306.

⁴ G. de Vries and A. Schors, *Tetrahedron Letters*, 1968, 5689.
⁵ H. M. Van Dort and H. S. Geursen, *Rec. Trav. chim.*, 1967, 86, 520.

⁶ D. A. Edwards, E. B. Evans, and B. T. Fowler, Chem. Abs., 1967, 66, 65113. (Fr. P. 1,447,697.)

⁷ P. R. Sharan, P. Smith, and W. A. Waters, J. Chem. Soc. (B), 1968, 1322.

through the solution for 20 h at room temperature. The black oxide was washed well with water and dried over anhydrous calcium chloride at reduced pressure.

TABLE 2

Oxidation of benzyl alcohol

Ratio	Yiel	Yield (%) ^a	
catalyst:substrate	Atmosphere	Öxygen added	
1:1	9	34	
3:1	14	50	
5:1	12	76	
7:1	12	85	

^a Determined by g.l.c.

We thank the Robert A. Welch Foundation for financial support.

(Received, April 10th, 1970; Com. 506.)