## DERWENT PUBLICATIONS LTD

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74529 E/35 UNION CARBIDE CORP \*US 4345-103

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Hydroxy-methylation of carbon residue of organo silicon cods. - with carbon mon: oxide and hydrogen over ruthenium carbonyl catalyst

Prepn. of alcohols (I) comprises reacting, in the homogeneous liq. phase, an organosilicon cpd. (II) with carbon monoxide and hydrogen in the presence of a ruthenium carbonyl catalyst. (II) has a carbon residue (R), and a gp. X, which is such that a transition metal can add across its bond to Si, bonded to the Si, and (I) has I C-atom and

1 O-atom more than the R of (II). Pref. X is H and (II) is esp. a mono-, di- or trialkylsilane. Reaction is pref. at 50-400°C and 100-15000 psia in a solvent and unreacted CO and H2 are recycled to the liq. phase.

## ADVANTAGE

Process is a relatively low pressure conversion of synthesis gas to alcohols. The Ru cpd. and (II) show a synergistic effect in the prepn. of MeOH, ethylene glycol and (I).

## DETAILS

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Suitable Ru catalysts are described in US2535060; and suitable solvents are ketones, esters, lactones, amides, lactams, sulphones, sulphoxides, aromatic hydrocarbons and ethers. A promotor may also be present. Ratio of CO to H<sub>2</sub> is pref. 40: 1 - 1: 40, esp. 10: 1 - 1: 10. Arnt. of Ru catalyst is pref.  $10^{-3}$  - 20 wt. % and amt. of (II)  $10^{-3}$  -100 wt. %.

## **EXAMPLE**

80 ml Trihexylsilane were reacted at 270°C and 8000 psig for 4 hr. at CO: H<sub>2</sub> ratio 1: 1 using 1 mmole Ru<sub>3</sub>(CO)<sub>12</sub> catalyst. After 1 hr. the prod. contained 1.4g MeOH, 0.41 g ethylene glycol and 5. 6g n-heptanol. After 4 hrs. the prod. contained 3. 9g MeOH, 0. 64g ethylene glycol and 17g n-heptanol. A similar experiment using no Ru3(CO)12 gave, after 4 hr., 0.23g MeOH and no ethylene glycol or n-heptanol (9pp 985).

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