UNION CARBIDE CORP 27.06.84-US-625371 (16.01.86) B01i-29/14 C07c-1/4 Fischer-Tropsch process giving lower selectivity to methane - and higher selectivity to liq. engine fuels by using catalyst contg. cobalt

E18 H04

with gold copper or silver as inert metal component C86-012042 E(AT BE CH DE FR GB IT LU NL SE) N(AU BR JP)

Synthesis gas is converted to 5C+ hydrocarbon mixts, useful

as liq. motor fuels by contacting at 240-370°C with a Fischer-Tropsch catalyst comprising: (a) Co mixed with (b) an inert metal component comprising Au, Ag, Cu or their mixts., the concn. of (b) being 0.1-50 mole % of (a+b).

ADVANTAGE

RA-028266/04

Component (b) decreases the selectivity of conversion to methane (e.g. from 24.1 to 17.8%), and increases that to liq. fuels (e.g. that of the 420 to 700°F fraction from 15.8 to

PREFERRED CATALYST The content of inert metal is pref. 0.5-5 mole % of that of Co. The Co may be promoted with K. Th or Na. There may also be a co-catalyst/support component, comprising

(c) a steam-stabilised hydrophobic zeolite Y, esp. one in Al-

24.0%). The liq. no longer contains solid prods.

UNIC 27.06.84 E(10-J2D3) H(4-D, 4-E5) N(2-B1, 2-D1, 2-E, 6-A)

extracted form (prepd. e.g. as in US3591488) in which the Co is placed largely within the crystallites; and/or (d) a crystalline microporous SAPO silicoaluminophosphate nonzeolitic mol. sieve (US 4480871).

If a cocatalyst/support is used, the concn. of Co is pref. 1-25 (esp. 5-15) wt. % of the catalyst.

CATALYST PREPARATION

The Co metal componnent (e.g. cobalt carbonate) can be impregnated with a soln, contg. the inert metal component. e.g. as HAuCla, after impregnating with a soln, of the promoter element, e.g. as Th(NO₁)₄, and then mixed with UHP-Y zeolite and SiO, binder, and the mixt. extruded, dried and calcined. Or the Co component can be adsorbed within the zeolite crystals.

EXAMPLE

*WO 8600-295-A

Catalysts comprised: (i) (comparative) 15 wt. % (Co + 15 wt. % Th), 70 wt. % UHP-Y zeolite and 15 wt. % SiO; and (ii) the same, with addn. of 2 wt. & Au relative to cobalt oxide.

The catalysts were used in conversion of a 1: 1 CO/H₂ WO8600295-A+

