DERWENT PUBLICATIONS LTD.

50251

50251 E/25 E19 H04 SHELL INT RES MIJ BV *BE -891-410 15.12.80-FR-026564 (09.06.82) B01j-23/46 C07b C07c-01/04 C10g Hydrocarbon prodn. from synthesis gas - using metal-contg. catalyst 0.1-10 wt.% Co and impreganted with up to 10 wt.% Cr, and based on laminar support

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Prodn. of hydrocarbons from a mixt. of CO and H₂ is carried out using a catalyst comprising (a) one or more metal components with catalytic activity for conversion cf H₂/CO mixts, to acyclic hydrocarbons, and (b) a support comprising a layered structure (esp. a crystalline silicate) capable of absorbing metal ions or salts by intercalation.

ADVANTAGES

The process gives high conversion at acceptable space velocities, with high selectivity for gasoline-range (5-12C) hydrocarbons.

DETAILS

The support is esp. magadiite (Na₂Si₁₄O₂9.9H₂O), but other silicates of Al, Fe and/or Ga, including other clay minerals of the candite, smectite and/or vermiculite type, can also be used. Component (a) may be Fe, Ni, Co, Cr and/or Ru. The catalysts can be prepd. by ion exchange. Prefd. catalysts comprise (i) magadite ion-exchanged with

SHEL 15.12.80 E(10-J2D) H(4-E5, 4-F2E) N(2, 3-D)

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(ii) magadiite ion-exchanged with 0.1-15 wt.% Ru. Hydrocarbon synthesis is pref. effected at 125-375°C and 1-150

EXAMPLE

A catalyst of compsn. 25 Co/1 Cr/296 SiO₂ was prepd. by converting magadiite to NH4 form by ion exchange with conc. NH4OH, impregnating with aq. Cr(NO₃)₃ soln., ion exchanging with aq. Co(NH₃)₆(NO₃)₂ soln., drying at 110°C, calcining in air at 500°C for 2 hrs., and reducing in H2 at 575°C for 24 hrs. A 1:1 mixt, of H2 and CO was passed over the catalyst at 260°C and 20 bar (GHSV = 1000). The conversion was 71% with 22% selectivity for 1-2C hydrocarbons, 10% for 3-4C, 50% for 5-12C, 12% for 13-19C and 6% for higher hydrocarbons.(12pp367)

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