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Europäisches Patentamt
European Patent Office
Office européen des brevets

11

Publication number:

**0 321 616
A1**

12

EUROPEAN PATENT APPLICATION

21

Application number: **87311323.7**

51

Int. Cl.⁴ **C07C 1/06**

22

Date of filing: **22.12.87**

43

Date of publication of application:
28.06.89 Bulletin 89/26

84

Designated Contracting States:
BE DE FR GB IT NL

71

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Process for reducing methane production and increasing liquid yields in Fischer-Tropsch reactions.

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A process for reducing methane formation and increasing liquid yields in Fischer-Tropsch processes for synthesizing hydrocarbons from CO and H₂ comprising adding water and/or an olefin to the top of the reactor bed or to the reactor feed at a point below 10% of the distance from the top to the bottom of the reactor bed and above a point 10% above the bottom of the reactor bed to the top of the reactor bed in an amount sufficient to reduce said methane formation, wherein the molar ratio of water (when employed) to CO is from about 0.1 to about 5.

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REDUCING METHANE PRODUCTION AND INCREASING LIQUID YIELDS IN FISCHER-TROPSCH REACTIONSSUMMARY OF THE INVENTION

It has been discovered, in accordance with this invention, that liquid (C₅⁺) yields and methane production in Fischer-Tropsch hydrocarbon synthesis reactions are respectively increased and reduced by adding one or more olefins directly into the reactor bed or by adding water and one or more olefins directly into the reactor bed, above or below the inlet port. Thus, the instant invention relates to a process for reducing methane production in Fischer-Tropsch processes wherein hydrocarbons are synthesized from feeds comprising mixtures of CO and H₂ and by adding at least one olefin to the reactor bed or water and one or more olefins, wherein the olefin is obtained by separation from the product stream and recycled to the reactor bed or from an independent source, such as processing the paraffinic product of the Fischer-Tropsch reaction through a dehydrogenation reactor to convert the paraffins into olefins. In a preferred embodiment, the olefin or olefins will comprise one or more C₂-C₂₀ alpha olefins, wherein the olefin to CO mole ratio in the reactor is from 1/20 to 5/1, more preferably from 1/10 to 2/1, and most preferably from about 1.5 to 1.1. The molar ratio of water to olefin is 0.1 to 50, more preferably 0.2 to 10 and most preferably 0.5 to 10. The catalyst will comprise at least one Group VIII metal supported on an organic refractory oxide support. Additionally, cobalt or iron catalyst can be easily employed.

In a particularly preferred embodiment the catalyst will comprise ruthenium supported on titania.

DETAILED DESCRIPTION OF THE INVENTION

It has now been discovered that methane production in Fischer-Tropsch hydrocarbon synthesis reactions is reduced by adding one or more olefins directly into the reactor bed or by adding water and one or more olefins directly into the reactor bed, above or below the inlet port. Thus, the instant invention relates to a process for reducing methane production in Fischer-Tropsch processes wherein hydrocarbons are synthesized from feeds comprising mixtures of CO and H₂ and by adding at least one olefin and or water and one or more olefin to the reactor bed, wherein the olefin is obtained by separation from the product stream and recycled to the reactor bed or from an independent source. In a preferred embodiment, the olefin or olefins will comprise one or more C₂-C₂₀ alpha olefins, wherein the olefin to CO mole ratio in the reactor is from 1/20 to 5/1, more preferably from 1/10 to 2/1, and most preferably from 1.5 to 1.1. The molar ratio of water to olefin is 0.1 to 50, more preferably 0.2 to 10 and most preferably 0.5 to 10. The catalyst will comprise at least one Group VIII metal supported on an inorganic refractory oxide support. In a particularly preferred embodiment the catalyst will comprise ruthenium supported on titania. Additionally, cobalt or iron catalyst can be readily employed.

Preferred olefins useful in the process of the instant invention include alpha olefins of the type R-CH=CH₂ wherein R is hydrogen or an alkyl group having 1 to 17 carbon atoms, more preferably the alkyl group has 1 to 11 carbon atoms, and most preferably the alkyl group has 1 to 8 carbon atoms. Still more preferably are C₂-C₁₀ alpha olefins. The amount of alpha olefin recycled to the reactor bed will be sufficient to maintain an olefin to CO mole ratio of from 1/100 to 5/1, more preferably 1/20 to 5/1, and most preferably 1/10 to 2/1. Also, internal olefins where unsaturation is away from the terminal carbon can also be added beneficially to the CO/H₂ feed.

Although the process of the instant invention may be practiced in the presence of any suitable Fischer-Tropsch catalyst, in a preferred embodiment it will be practiced in the presence of a catalyst comprising one or more Group VIII metals supported on an inorganic refractory oxide support, preferably ruthenium or cobalt supported on such a support. Thus, suitable supports include oxides of titania, niobia, vanadium, tantalum, silica, alumina, manganese and mixtures thereof. Preferably the catalyst support will be selected from the group consisting of titania, zirconium titanate, mixtures of titania and alumina, mixtures of titania and silica, alkaline earth titanates, alkali titanates, rare earth titanates and mixtures of any one of the foregoing with supports selected from the group consisting of vanadia, niobia, tantalum, alumina, silica and mixtures thereof. Thus, in a particularly preferred embodiment of this invention the process will be carried out in the presence of a catalyst comprising ruthenium supported on a titania support. Alternatively, the catalyst can comprise cobalt or iron supported on one of the aforementioned inorganic refractory oxides.

In general, the amount of cobalt catalytic metal present is 1 to 50 weight percent of the total catalyst

composition, more preferably from 10.0 to 25 weight percent.

Iron catalysts containing 10 to 60 weight percent iron, more preferably 20 to 60 weight percent, and most preferably 30 to 50 weight percent, are unsupported, but promoted with refractory metal oxide (SiO_2 , Al_2O_3 , etc.), alkali (K, Na, Rb) and Group IB metal (Cu, Ag). These catalysts are usually calcined, but usually not reduced, rather they are brought up to reaction temperature directly in the CO/H_2 feed.

In general, the amount of ruthenium catalytic metal present on the catalyst will generally range from 0.01 to 50 weight percent of the total catalyst composition, more preferably from 0.1 to 5.0 weight percent and most preferably from about 0.5 to about 5 weight percent.

In general, in the process of this invention the Fischer-Tropsch reactor temperature will broadly range from 100°C to 500°C , more preferably from 150°C to 300°C and most preferably 200°C to 270°C , at a pressure of 100 to 10,000 kPa, more preferably 300 to 5,000 kPa and most preferably 500 to 3,000 kPa. The space velocity (V_1) of the feed gas ($\text{H}_2 + \text{CO}$) will range from 10 to 10,000 standard $\text{cm}^3/\text{hr}\cdot(\text{cm}^3$ of catalyst), more preferably 100-4,000 and most preferably 200 to 2,000. The space velocity (V_2) of the alpha olefin will range from 0.1 to about 20,000 standard $\text{cm}^3/\text{hr}\cdot(\text{cm}^3$ of catalyst), more preferably 3-4,000 and most preferably 10 to 1,000. Thus, the volume ratio of V_1/V_2 is 0.005 to 10,000, more preferably 0.02 to 1,000 and most preferably 0.5 to 200. The hydrogen to carbon monoxide mole ratio of the feed gas, H_2/CO , will range from 0.5 to 10 and preferably from 1 to 3.

If the alpha olefin is used without water addition, the alpha olefin is injected into the reactor through an inlet port in the sidewall or center of the reactor, wherein the part is positioned below a point which is 10% of the distance from the top of the reactor bed to the bottom of the reactor bed and above a point which is 10% of the distance from the bottom of the reactor bed to the top of the reactor bed.

The water and alpha olefin are injected into the top of the reactor through inlet ports in the sidewall of the reactor or through inlet ports which are positioned anywhere in the reactor, but preferably below a point which is 10% of the distance from the top of the reactor bed to the bottom of the reactor bed and above a point which is 10% of the distance from the bottom of the reactor bed to the top of the reactor bed. The ports for the water and alpha olefin can be located at different points within the reactor.

The reactor bed can comprise a single reactor bed of one catalyst system or it can comprise two catalyst beds with different Fischer-Tropsch synthesis catalysts separated by an inert inter-stage knock-out zone. The water and/or alpha olefin can also be injected through an inlet port located at the interstage knock-out zone. The alpha olefin can also be injected through an inlet port located at the inter-stage knock-out zone. Alternatively, the system can comprise a plurality of tubular reactors joined in a serial fluid communication wherein the alpha olefin is injected into the system either within the first reactor bed or subsequent thereto with or without inter-stage knock-out zone.

DESCRIPTION OF THE PREFERRED EMBODIMENTS

The invention will be more readily understood by reference to the following Examples.

Fischer-Tropsch Hydrocarbon Synthesis

Experimental Procedure

Anywhere from about 5 to 40 g of catalyst was loaded into a stainless steel reactor with an ID of 0.78 cm, having a thermocouple placed within the catalyst bed to measure temperature. The reactor was flushed with He at a flow rate of 500 cc/min. and then pressurized (in He) at 30 atmospheres to confirm the integrity of the unit. The He in the reactor was then brought back to 1 atmosphere and replaced with H_2 at a space velocity between 200-600 v/v/hr at room temperature. The temperature was raised in approximately 2 to 3 hours to a final reduction temperature of between 400°C to 500°C and held at this temperature overnight.

After reduction, each catalyst was brought on stream in either one of two ways, which were found to be substantially equivalent. In one method, the reduced catalyst was treated for 15 to 50 hours at 1 atmosphere $2\text{H}_2/\text{CO}$ at 200°C (the space velocity of the H_2/CO used was varied from 200-1,000 $\text{cm}^3\text{H}_2 + \text{CO}$ h. (cc catalyst)). The conditions in the reactor were then adjusted to the desired temperature, pressure and space velocity. In the other method, the atmospheric treatment in H_2/CO was not done and the catalysts were brought on-stream at the desired experimental pressure in H_2/CO , but an initial low

temperature of 150-170 °C, which was then slowly raised (in 1 to 5 hours) to the desired experimental temperature.

5 EXAMPLES

In all of the Examples, the determination of the Fischer-Tropsch synthesis activity towards the various products was measured by employing an on-line gas chromatograph using an isothermal, fixed-bed, stainless steel reactor (0.78 cm ID) into which from about 5 to 40 g of catalyst was loaded. A thermocouple
10 was placed within the catalyst bed to measure the reaction temperature.

Example 1

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Preparation of Ru Catalyst

20 Titanium dioxide supports prepared by pressing Degussa P-25 powder (20-70% rutile) into wafers using a hydraulic press at a pressure of about 20,000 lb/in². The wafers were then crushed in a mortar to a coarse powder and sieved to retain the 80-140 mesh (U.S. standard) fraction for all the TiO₂ supported catalysts. The so-formed titanium dioxide support samples were calcined in flowing air overnight (about 1 liter per minute) at a temperature of 550 °C to 600 °C. The calcined samples were then reduced in hydrogen flowing
25 at a rate of 1 liter per minute by heating from room temperature to 450 °C for 4 hours, after which the sample was cooled in flowing H₂, flushed with He and discharged. These reduced samples possessed the characteristic blue-gray color of surface-reduced TiO₂.

Catalysts were produced by impregnating the support material with a ruthenium nitrate/acetone mixture. The nitrate was obtained from Engelhardt as 10% weight ruthenium metal in nitric acid.

30 The nitrate-impregnated TiO₂ was reduced in flowing hydrogen (1 l/min.) as follows: (a) heated from room temperature to 100 °C and held for 0.5 hours; (b) heated from 100 °C to 450 °C at 3 °C/min. and held for 4 hours; (c) cooled in H₂ to room temperature overnight; and (d) flushed in the reduction cell with He or Ar (1 l/min.) for one hour. Each sample of reduced catalyst was then passivated by introducing 1% O₂ into the He stream and then increasing the O₂ content to 100% over a period of from about 2-3 hours.

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Example 2

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Preparation of Cobalt Catalyst

45 Degussa P-25 TiO₂ was treated with acetone, dried at 90 °C and milled with 9.6 weight percent Sterotex (vegetable stearine). This material was tableted to 3/8 inch size, using a pilling pressure which yielded a material of 0.35-0.40 ml/g pore volume. The 3/8 inch pills were crushed and screened to 42-80 mesh size and calcined in flowing air at 500 °C for 4 hours to burn out the sterotex. Portions were then re-calcined in flowing air at 650 °C for 16 hours to obtain a high rutile content and screened to 80-150 mesh size. The support composite used to prepare this catalyst was 97% rutile, 13.6 m²/g, 0.173 ml/g.

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Impregnation Procedure

55 108.8 g cobaltous nitrate (Co(NO₂)₂ · 6H₂O) and 16 ml aqueous perchloric acid solution (52 mg Re/ml) were then dissolved in about 300 ml acetone in a 1 liter round bottom flask. 160 g TiO₂ (YAX-0487) was added and the mixture placed on a rotary evaporator until dry. The catalyst was dried in a vacuum oven at 140 °C for 18 hours and calcined in flowing air at 250 °C for 3 hours. The calcined catalyst was re-screened to remove the small amount of fines generated during preparation.

Cobalt Catalyst Preparation

Analytical inspections for the catalyst are given below:

	Cobalt, Wt. %	11.3
5	Rhenium, Wt.%	0.42
	Sodium ppm	369
	% Rutile (ASTM D 3720-78)	97
	Surface Area, m ² /g	14.6
	Pore Volume, ml/g	0.122
10	Bulk Density, g/ml	1.41

Besides quantifying rutile content, the x-ray diffraction spectrum indicates the presence of CO₃O₄.

Example 3

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Effect of Ethylene Feed-Cobalt Catalysts

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The general procedure described for Fischer-Tropsch hydrocarbon synthesis was modified as follows in Examples 3 and 4.

The catalyst was started in H₂/CO at a space velocity required to achieve 25-45% CO conversion and at 1,700-2,500 Kpa. An ethylene/argon mixture was mixed with H₂/CO feed and then introduced to reactor at inlet at an amount of ethylene to give 2.5% and 6.2% ethylene in feed on Co and Ru catalysts, respectively. The total reactor pressure was then increased to the level necessary to maintain the H₂ + CO partial pressure in the reactor at the level found before ethylene addition. The reactor was operated at these conditions for 4 to 24 hours. Ethylene/argon feed was then directed so that it entered the reactor, undiluted by H₂ + CO, at a point in the reactor 1/3 x L (L reactor length) down from inlet, while maintaining H₂ + CO space velocity and partial pressure at its previous level. The reactor was operated at these conditions for 4 to 24 hours. The ethylene/argon feed was discontinued and reactor conditions returned to those of beginning in order to rule out irreversible deleterious effects of ethylene addition on synthesis activity and selectivity.

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	Run	119-342	119-339
	% Ethylene in Feed	0	2.5
5	Location Added	None	Inlet
	<u>Ethylene</u>		
10	% Ethylene Converted	-	99.5
	Selectivity (%) to		
	C ₂ H ₆	-	86.5
15	C ₃ ⁺	-	13.5
	C ₃ ⁺ Yield (%)	-	13.5
20	Selectivity (%)*		
	CH ₄	9.4	8.3
	C ₂	1.3	-
25	C ₃	2.7	3.3
	C ₄	2.4	2.9
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CO ₂	0.4	0.5
C ₅ ⁺	83.8	92.9

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* % of ethylene converted which appeared as these product.

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** % of ethylene added which appeared as these products.

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*** Selectivity is defined as percent of the CO converted which appears as a given product (it is almost identical to weight percent). However, when ethylene is added some of the C-atoms come from it and the sum of selectivities is greater than 100%.

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11.7% Co/0.5 Re/TiO₂, 200°C, 2,050 kPa, H₂ + CO,
H₂/CO = 2.1/1
25-27% CO conversion

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The addition of ethylene (C₂⁺) at reactor inlet lowers the CH₄ selectivity and increases C₅⁺ selectivity. Most of the ethylene (at 2.5% concentration) is converted, but only 13.5% of the converted ethylene appears as desirable C₃⁺ products; the rest (86.5%) is hydrogenated to ethane, which is unreactive towards further chain growth to C₃⁺ at Fischer-Tropsch synthesis conditions and, thus, cannot be recycled without converting it to ethylene in a dehydrogenation reactor.

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Example 4

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Effect of Ethylene Feed-Ruthenium Catalysts

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Run	121-455	121-454
% Ethylene in Feed	0	6.2
Location Added	None	Inlet

Ethylene

% Ethylene Converted	-	97
Selectivity to (%)*		
Ethane	-	82
C ₃ ⁺	-	18
Yield (%) C ₃ ⁺ **	-	17
Selectivity (%)***		
CH ₄	5.5	4.3
C ₂	0.7	-
C ₃	2.1	2.6
C ₄	3.5	4.0
CO ₂	0.7	0.7
C ₅ ⁺	87.5	105.6

* % of ethylene converted which appeared as these products.

** % of ethylene added which appeared as these products.

*** Selectivity is defined as percent of the CO converted which appears as a given product (it

is almost identical to weight percent). However, when ethylene is added some of the C-atoms come from it and the sum of selectivities is greater than 100%.

1.2% Ru/TiO₂, 200°C, 1,700 kPa, H₂/CO = 2.1/1, 35-45% CO conversion.

Addition of ethylene to H₂/CO feed at reactor inlet (at 6.2% level) lowers the CH₄ selectivity from 5.5 to 4.3, while increasing the C₅⁺ selectivity from 87.5 to 105.6%. Most of the ethylene added in (97%) is converted, of which only 18% appears as desirable C₃⁺ products; the rest (82%) is hydrogenated to ethane, which is unreactive towards further chain growth to C₃⁺ at Fischer-Tropsch synthesis conditions and, thus, cannot be recycled without converting it to ethylene in a dehydrogenation reactor.

Example 5

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Top versus Below Inlet Feed Ethylene-Ru

There is a clear advantage to bypassing the reactor inlet by introducing the C₂⁼ below the inlet at the same concentration. The CH₄ selectivity decreases further (from 4.3% to 3.9%), the C₅⁺ selectivity increases (from 105.6% to 117.0%) and the activity of the catalyst is unaffected by the presence of ethylene, either at inlet or below the inlet of the reactor. The apparent reason for the improved product selectivity is an increase in the yield of C₃⁺ from ethylene (the percentage of the ethylene feed converted to C₃⁺) from 17% to 26% in spite of the lower ethylene conversion when added below the inlet. Below inlet addition dramatically decreases the hydrogenation selectivity (from 82% to 44%), while increasing the C₃⁺ selectivity (from 18% to 56%). Therefore, it dramatically increases the efficiency of ethylene utilization while avoiding the formation of species such as ethane which cannot be polymerized further under the Fischer-Tropsch conditions.

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Run	121-453	121-454
% Ethylene in Feed	6.2	6.2
Location Added	Below Top	Top
	Third of	
	Reactor	

Ethylene*

% Ethylene Converted	45.3	97
Selectivity to (%)		
Ethane	44	82
C ₃ ⁺	56	18

Yield (%) C ₃ ⁺ **	26	17
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Selectivity (%)***

CH ₄	3.9	4.3
C ₂	-	-
C ₃	3.4	2.6
C ₄	5.1	4.0
CO ₂	0.7	0.7
C ₅ ⁺	117.0	105.6

* % of ethylene converted which appeared as these products.

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** % of ethylene added which appeared as these products.

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*** Selectivity is defined as percent of the CO converted which appears as a given product (it is almost identical to weight percent).

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However, when ethylene is added some of the C-atoms come from it and the sum of selectivities is greater than 100%.

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1.2% Ru/TiO₂, 200°C, 1,700 kPa, H₂/CO = 2.1/1, 35-45% CO conversion.

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Example 6

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Top versus Below Inlet Feed Ethylene-Co

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There is a clear advantage to bypassing the reactor inlet by introducing the C₂ = below the inlet at the same concentration. The CH₄ selectivity decreases further (from 8.3% to 7.2%), the C₅ ↑ selectivity increases (from 92.9% to 99.3%) and the activity of the catalyst is unaffected by the presence of ethylene, either at inlet or below the inlet of the reactor. The apparent reason is an increase in the yield of C₃ ↑ from ethylene (the percentage of the ethylene feed converted to C₃ ↑) (from 13.% to 27.3%), in spite of the lower ethylene conversion when added below the inlet. Below inlet addition dramatically decreases the hydrogenation selectivity (from 86.5% to 69.5%), while increasing the C₃ selectivity (from 13.5% to 30.5%). Therefore, it dramatically increases the efficiency of ethylene utilization, while avoiding the formation of species such as ethane which cannot be polymerized under Fischer-Tropsch conditions.

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Run	119-336	119-339
% Ethylene in Feed	2.5	2.5
Location Added	Below Top Third	Inlet

Ethylene

% Ethylene Converted	89.5	99.5
Selectivity (%) to		
C ₂ H ₆	69.5	86.5
C ₃ ⁺	30.5	13.5
C ₃ ⁺ Yield (%)	27.3	13.5

Selectivity (%) *

CH ₄	7.2	8.3
C ₂	-	-
C ₃	3.7	3.3
C ₄	3.7	2.9
CO ₂	0.4	0.5
C ₅ ⁺	99.3	92.9

* % of ethylene converted which appeared as these products.

** % of ethylene added which appeared as these products.

*** Selectivity is defined as percent of the CO converted which appears as a given product (it is almost identical to weight percent). However, when ethylene is added some of the C-atoms come from it and the sum of selectivities is greater than 100%.

11.7% Co/0.5 Re/TiO₂, 200°C, 2,050 kPa, H₂ + Co,
H₂/CO - 2.1/1
25-27% CO conversion.

Example 7

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Experimental Procedure for Ethylene Addition

The catalyst was started in H₂/CO at a space velocity required to achieve 15-30% CO conversion at 2,000-2,100 kPa. Conditions were maintained for at least 48 hours. An ethylene/argon mixture was then mixed with the CO/H₂ feed and introduced to the reactor at the inlet at an ethylene concentration of 2-7% by volume. The total reactor pressure was then increased so as to maintain H₂ + CO partial pressure equal to that prior to ethylene cofeed. The reactor was operated at these conditions for at least 24 hours.

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Effect of Ethylene Addition - Ruthenium Catalyst

20	Run	123-608	123-609
	% Ethylene in Feed	0	7
	% Ethylene Converted	-	73.4
25	Selectivity to (%)*	-	

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	Ethane	-	50.5
	C ₃ ⁺	-	49.5
5	Yield to (%)**		
	C ₃ ⁺	-	36.3
10	Rate of Ethylene Conversion		
	(mol C ₂ =/g. atom Ru.h) C ₃ ⁺	-	6.1
	C ₂ H ₆	-	6.3
15	Rate of CO conversion		
	(mol CO/g.atom Ru.H)	12.8	12.5
20	CO Selectivity (%)***		
	CH ₄	3.7	3.0
	C ₂	0.4	-
25	C ₃	1.75	2.26
	C ₄	2.89	3.4
	CO ₂	0.3	0.4
30	C ₅ ⁺	91.0	188.5

1.2% Ru/TiO₂, 191-192°C, 2,100 kPa, H₂+CO, H₂/CO =
2.1/1, 18-22% CO conversion.

* % of ethylene converted which appeared as these products.

** % of ethylene added which appeared as these products.

** Selectivity is defined here as the percent of CO converted which appears as a given product. However, when ethylene is added some of the C-atoms come from it and the value of this "selectivities" can exceed 100%.

The addition of ethylene (C₂=) to the H₂/CO feed at reactor inlet lowers the CH₄ selectivity and increases the C₅⁺ selectivity on the Ru/TiO₂ catalyst. A large fraction of the ethylene (7% concentration to

the feed) added is converted (73.4%), but only 49.5% of the converted $C_2^=$ appears as desired C_3^+ products; the rest (50.5%) is hydrogenated to ethane, which is unreactive towards further chain growth at Fischer-Tropsch synthesis conditions and, thus, cannot be recycled without converting it to ethylene in a dehydrogenation reactor.

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Example 8

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Experimental Procedure for Ethylene Addition and Water Cofeed

The catalyst was started in H_2/CO at a space velocity required to achieve 15-30% CO conversion at 15 2,000-2,100 kPa. Conditions were maintained for at least 48 hours. An ethylene/argon mixture was then mixed with the CO/H_2 feed and introduced to the reactor at the inlet at an ethylene concentration of 2-7% by volume. The total reactor pressure was then increased so as to maintain $H_2 + CO$ partial pressure equal to that prior to ethylene cofeed. The reactor was operated at these conditions for at least 24 hours. When 20 water vapor was introduced at a concentration of 10-20 volume percent by bubbling the $H_2 + CO$ feed through a saturator maintained at 120-145 °C, while keeping the ethylene/argon flow constant. The total reactor pressure was increased so as to maintain the $H_2 + CO$ and the ethylene partial pressures at their level prior to water cofeed. Also, the flow rates of H_2/CO and of ethylene/argon were increased so as to maintain the CO conversion at their level prior to water addition. The reactor was operated at these conditions for at least 24 hours.

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Effect of Ethylene/Water Addition - Ruthenium Catalyst

Run	123-609	123-614
% Ethylene in Feed	7	7
% Water in Feed	0	18
% Ethylene Converted	73.4	38.2
Selectivity to (%) *		
Ethane	50.5	11.5
C_3^+	49.5	88.5
Yield (%) **		
C_3^+	36.3	33.8
Rate of Ethylene Conversion to		
(mole $C_2^=/g.atom Ru.h$)		
C_3^+	6.1	17.3
C_2H_6	6.3	2.2

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Rate of CO Conversion

(mol CO/g.atom Ru.h)	12.5	34.0
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CO Selectivity (%)***

CH ₄	3.0	0.9
C ₂	-	-
C ₃	2.26	0.97
C ₄	3.40	1.46
CO ₂	0.4	0.4
C ₅ ⁺	188.5	198.5

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1.2% Ru/TiO₂, 191-192°C, 2,100 kPa, H₂+CO, H₂/CO = 2.1/1, 18-22% CO conversion.

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* % of ethylene converted which appeared as these products.

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** % of ethylene added which appeared as these products.

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*** Selectivity is defined here as the percent of CO converted which appears as a given product. However, when ethylene is added some of the C-atoms come from it and the value of this "selectivities" can exceed 100%.

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The Table shows that there is a clear advantage to the cofeeding of water vapor with the ethylene and with the H₂ CO feed. While the ethylene conversion decreases from 73.5% to 38.2% with the addition of 18% H₂O, the yield decreases only slightly (36.3% to 33.8%). However, most importantly:

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o the selectivity to desired C₃⁺ products increases from 49.5% to 88.5%;

o the rate of ethylene conversion to C₃⁺ products and of CO conversion increase approximately threefold; in other words, the activity of the catalyst for both reactions, C₂H₄ incorporation and Fischer-Tropsch synthesis, increases threefold.

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o the rate of ethylene conversion to ethane (i.e., hydrogenation rate), actually decreases approximately threefold with the addition of 18% water vapor to the H₂/CO/C₂⁼ feed.

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Accompanying beneficial effects are a decrease in the CH₄ selectivity (3.0% to 0.9%) and an increase in the effective C₅⁺ selectivity from 138.5% to 199.5%.

These results suggest that the effect of water is to enhance the reincorporation and chain growth of added ethylene while dramatically inhibiting its hydrogenation to undesired ethane product.

Notes

- o Mass expressed in pounds (lbs) is converted to kg equivalent by multiplying by 0.4536.
- o Length expressed in inch is converted to cm by multiplying by 2.54.

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Claims

1. A process for reducing methane and formation and increasing liquid (C₅⁺) yields in Fischer-Tropsch
10 process for synthesizing hydrocarbons from CO and H₂, comprising adding one or more olefins to the
reactor bed at a point below 10% of the distance from the top to the bottom of the reactor bed and above a
point 10% above the bottom of the reactor bed to the top of the reactor bed in an amount sufficient to
reduce said methane formation.

2. The process of claim 1 wherein the amount of said one or more olefins added to said reactor bed is
15 based on an olefin to CO mole ratio in the range of from 1/100 to 5/1.

3. The process of claim 1 or claim 2 wherein said olefin(s) is or are alpha olefin of the type R-CH=CH₂,
wherein R is hydrogen or an alkyl group having 1 to 17 carbon atoms.

4. The process of any one of claims 1 to 3 wherein said olefin comprises a C₂-C₁₀ alpha olefin.

5. The process of any one of claims 1 to 4 conducted in the presence of a catalyst.

20 6. The process of claim 5 wherein said catalyst comprises one or more Group VIII catalytic metals.

7. The process of claim 6 wherein said catalytic metal is selected from iron and ruthenium.

8. The process of any one of claims 1 to 7 performed at a temperature of at least 100 °C, preferably in
the range of from 100 °C to 500 °C.

9. The process of any one of claims 1 to 8 wherein water and an olefin are present in the reactor or
25 reactor bed in an amount sufficient to reduce said methane formation and increase liquid yield.

10. The process of any one of claims 1 to 9 comprising adding water and an olefin to the inlet of the
reactor bed or to the reactor at any point in said reactor bed in an amount sufficient to reduce said methane
formation.

11. The process of claim 9 or claim 10 wherein the molar ratio of water to CO is in the range of from
30 0.1 to 5.

12. The process of any one of claims 9 to 11 comprising contacting H₂/CO/H₂O with an olefin.

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DOCUMENTS CONSIDERED TO BE RELEVANT			
Category	Citation of document with indication, where appropriate, of relevant passages	Relevant to claim	CLASSIFICATION OF THE APPLICATION (Int. Cl.4)
X	US-A-4 547 525 (KIM) * Claims * ----- -	1-8	C 07 C 1/06
			TECHNICAL FIELDS SEARCHED (Int. Cl.4)
			C 07 C 1/00
The present search report has been drawn up for all claims			
Place of search THE HAGUE		Date of completion of the search 22-08-1988	Examiner VAN GEYT J.J.A.
<p>CATEGORY OF CITED DOCUMENTS</p> <p>X : particularly relevant if taken alone Y : particularly relevant if combined with another document of the same category A : technological background O : non-written disclosure P : intermediate document</p> <p>T : theory or principle underlying the invention E : earlier patent document, but published on, or after the filing date D : document cited in the application L : document cited for other reasons & : member of the same patent family, corresponding document</p>			