UNITED STATES DEPARTMENT OF THE INTERIOR BUREAU OF MINES OFFICE OF SYNTHETIC LIQUID FUELS LOUISIANA, MISSOURI

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W. M. Sternberg Oct. 21, 1947

Holten, 12/14/41

CONNECTING THE PRESSURE AND LOW PRESSURE SYNTHESIS IN SERIES BASED ON 65,000 m³ water Gas.

To Dir. Alberts.

I am presenting my proposal (made in accordance with my report of 4/3/41 given in the appendix) for obtaining the greatest possible proportion of paraffin and the production of a greater liquefaction in the pressure synthesis by connecting in series the pressure synthesis (D) and the low pressure synthesis (N).

Nothing stands in the way in the operations and construction of the system of operation in series, after the assumptions made at one time regarding the reactor load and gas conversion have been confirmed, and after the installation of the large separators behind the condensation stage and heaters and in front of the synthesis stages. The construction of the third stage of condensation is ready since the middle of January, and a thruput of 25,000 m² end gas is possible. The available condensation groups of the 1-st and second stages were then used for the end gas of the 1-st synthesis stage in parallel, and the supplementary condensation used for the end gas of the second synthesis stage.

After the shortly to-be-installed cooler elements, and with the operating level reached with three machines (with one more in reserve), the compressor installation will handle the required 62,000 m³. The CO₂ scrubber must be considered safe for operations after the installation back of it (beginning of March) of a separator the size of the scrubber proper, and a safety ball valve to automatically step the carrying-over of vater; they may accordingly be considered as available for the washing of the converted gas intended for the pressure synthesis. A gas thruput of only 10,000 m³ is possible until March, because at present the 2-nd scrubber is in use as a water separator.

With these connections, modified by to-day's proposal, all of the gas (based on 65,600 m 3 of water gas) is passed thru the pressure synthesis,

except for that portion of the gas which is used after the conversion for producing the required CO: H2 proportion of 1: 2 in the gas used for normal pressure synthesis (cf. the operations flow sheet, app.2).

The composition of the intake and the conversion results are as follows:

	8,8	7.6	77.4	0.4	5.8
Converter gas (75% CO2 removed in	washing	:(:			
Converter gas (80% CO conversion)	28	6,0	61	0.3	4.7
	5.2	39.9	48.2	0.4	6.3
Water gas analysis:	ගෘ	CO	Ha	CHE	N ₃

Input, first step:

44,000 m3 water gas

+ 8.000 washed converter gas 52,000 M³ synthesis gas, 1-st stage (D synthesis)

The composition of this gas: CO₂ CO H₂ CH₄ N₂
5.8 35 52.5 0.4 5.3

Water gas input, 1-st stage

44,000 + 7,600 (8,000 m³ washed conv. gas) = 51,600 m³ washer gas

Efficiency, 1-st stage

Contraction 51%
CO conversion 50%
COs formation 00%
CHo formation 10%

Proportion of CO: He utilized = 1:2.03
and therefore 25,400 m3 of the end see after 1-st stage
has the following composition CO: CO: He CH: No.

11.8 35.6 35.5 4.4 12.7

The proportion CO: H₂ for 2-nd stage synthesis gas must be again raised to 1: 1.5

Addition of washed converted gas - 7,000 m³ = 6.700 m³ washed gas which will increase the input: 51,600 + 6,700 = 58,200 m³ water gas Composition of synthesis gas, 2-nd stage

CO₂ CO H₂ CH₄ N₂
11.2 29.5 44.5 3.6 11.7

2-nd stage gas imput: 32,400 m3

conversion 50%

Contraction 44%

CO2 formation 00%

CHe formation 10%

CO and H2 consumed in proportion 1 : 2.03

which results in an end gas of the following composition:

CO₂ CO H₂ CH₄ N₅

19.6 26.3 25.3 9.1 19.7

Volume of end gas: 18,300 m³

The proportion CO : H_0 in this gas is raised to 1 : 2 by the addition of converted gas. Required amount of unwashed converted gas : 10,000 m^3

= 7.400 m³ water gas

The total water gas addition figures than to

D. synth, 1-st stage D synth., 2-nd stage N. synth.

51,600 + 6,700 + 7,400 = 65,600 m³/h water gas.

Imput of gas into N. synthesis

18,300 + 6,700 = 28,300 m^3/h

of the following composition:

CO_S CO H₂ CH₄ N₂

22,5 19,0 37,8 6,1 14,6

With as

Conversion of CO 65% Formation of CO₂ 3% Formation of CH₃ 15% Contraction 36%

there will remain $18,200 \text{ m}^3/\text{h}$ of end gas of the composition 35.8 10.4 19.2 12.1 22.6

The synthesis and converter load may be calculated from the following considerations:

Pressure Synthesis:

46 reactors, 1-st stage, 1120 m³/h, 35% CO, 52.5% H₂, CO : H₂ = 1:1.5;

CO conversion 50%

2-nd " 1800 m³/h, 29.5% Co, 44.5% H₂, CO : H₂ = 1:1.5

CO conversion 50%

64 # 1+2 st., 925 m5/h (sic) 31.6% CO. 55.8%H2, CO: H2 = 1:1.76; CO conversion 74

Low Pressure Synthesis:

40 reactors, single stage, 700 m³/h, 19,6% CO, 37,6% Hg, CO : Hg = 1:1,99 CO conversion 65%

A balance and comparison of the CO and H₂ imputs and conversion in both synthesis:

The combined imput was:

co: 19,300 m3/h

Ea: 38,700 #

Of this amount, in the 1-st. and 2-nd. stages of the D synthesis

CO: 18,730 m3/h

Ha: 32,600 #

Of this amount, converted:

CO: 13,900 m3/h

Hg: 28,000 *

1.e., with a degree of conversion of

CO: 76%

Ha: 86%

Imput into the N. Synthesis:

00: 5,400 m³/h

Ha: 10,700 0

Of this amount, converted

00: 3,500 m3/h

Ha: 7,200 #

which is represented by a degree of conversion of

CO: 65%

H2: 67%

There remains in the residual gas after the N. conversion

CO: 1,500 m3/h

H2: 3,500 #

These figures give for the total degree of conversion of all CO and $H_{\mathbf{2}}$ sent through the system:

90.3% for co

91.0% for Ha

and, when distributed to the different syntheses:

D. synthesis 72.0% CO 72.5% Ha

N. " 18.3% CO 18.5% H₂

The advantages of the new or old system may be judged as follows:

The total gas is worked up in three stages, with the mild conversion and starting conditions required for the operations. The mutual influences of mass—relationships are here eliminated. According to the subdivision given, 80% of the CO conversion proceeds in the pressure synthesis, and 20% in the low pressure synthesis. Disregarding the increased amount of paraffine, the liquefaction in the pressure synthesis is better than in the low pressure synthesis, and this may be assumed as proven for the proportion of CO: $H_2 = 1$: 1.5.

Operations are in general simplified in this method, which is desirable. The production is measured separately for the pressure and the low pressure syntheses up to the production of gasol.

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i.e., with a degree of conversion of

CO: 76%

Ha: 86%

Input into the N. Synthesis:

60: 5,400 m3/h

Ha: 10,700 4

Of this amount, converted

co: 3,500 m3/h

Ha: 7,200 #

which is represented by a degree of conversion of

CO: 65%

H2: 67%

There remains in the residual gas after the H. conversion

co: 1,900 m3/h

Ha: 3,500 #

These figures give for the total degree of conversion of all CO and H_{2} sent through the system:

90.35 for co

91.0% for Ha

and, when distributed to the different syntheses:

D. synthesis 72.0% CO 72.5% Ha

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The advantages of the new or old system may be judged as follows:

The total gas is worked up in three stages, with the mild conversion and starting conditions required for the operations. The initial influences of mass—relationships are here eliminated. According to the subdivision given, 80% of the CO conversion proceeds in the pressure synthesis, and 20% in the low pressure synthesis. Disregarding the increased amount of paraffine, the liquefaction in the pressure synthesis is better than in the low pressure synthesis, and this may be assumed as proven for the proportion of CO: H2 = 1:1.5.

Operations are in general simplified in this method, which is desirable. The production is measured separately for the pressure and the low pressure syntheses up to the production of gasol.

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APPENDIX I

Holten, May 3, 1941

CONNECTING THE HEDIUM AND THE LOW PRESSURE STREHESIS IN SERIES, ON A BASIS OF 60,000 m³ WATER GAS.

The medium pressure synthesis is conducted in two s'ages, with a CO:H2 ratio of 1:1.5, and followed with a single stage low pressure synthesis with a CO:H2 ratio of 1:2.

50,000 m³ water gas including the converted gas were sent through the system, determined by the capacity of the gas compressor, and taking into consideration the load on the pressure reactors, on the pressure condensation and the A.K. installation II. The balance of 10,000 m³ water gas were mixed with the end gas of the pressure synthesis and worked up at the low pressure synthesis.

The load on the reactors is:

Medium pressure synthesis:

36 reactors, 1-st stage, 1280 m3/h: 35% CO, 50% Ha

24 " 2-nd " 1200 " 25.5% CO, 38% H₂

Low Pressure Synthesis:

45 reactors

720 m³/h: 18% co, 35.5% H₂

The total imput is:

00 17, 310 m3

Hg 34, 300 "

of which the amount sent through the first and second stages, D synthesis:

CO 15,410 m³

Ha 22,860 "

and converted:

00 75%

H2 85%

The amount sent through the N. synthesis is:

CO 5,780 m3

Ha 11,340 "

and converted;

CO 4,330 m³

Ha 8,440 *

i.e. with a degree of conversion of

CO 75%

H2 75%

There remain in residual gas of the N. synthesis

CO 1,940 m3

H₂ 2,900 m3

Conversion, figured to the total input is

CO 89\$

Ha 91%

Were we to compare this process with the one in present day use, we would get the following results. At present we pass through the D. synthesis

23,000 m⁵ synthesis gas composed of 28.6% CO and 51.6% He. and in the N. synthesis:

52.8% H2. 23,000 m3 of synthesis gas, cont aining 26.4% Co and

First case: Second *	D. synthesis " (at present)	00 imput 15,410 m3 10,000 #	CO conversion 11,530 m ³ 8,600 "
First case:	N. synthesis " (at present)	5,780 #	4,330 #
Second #		7,400 #	6,590 #

The compositions of the input and conversion are as follows:

Water gas input

Total gas addition (v. flow sheet, app. I)

29,000 m³ water gas

7.000 converted gas

46,000 synthesis gas

of the following composition:

The 7,000 m³ converted gas have been obtained from 5,400 m³ water gas, i.e. the water gas input into the first stage is

End gas, first stage

The proportion CO: H_2 is again restored to 1:1.5 by the addition of converted gas, for which we shall require 6.200 m³ converted gas = 4,800 m³ water gas.

Total input of water gas 44,400 * 4,800 = 49,200 m³

The composition of synthesis gas II:

Gas input into second stage: 23,000 + 6,200 = 29,200 m³ synthesis cas II. Contraction, 2-nd stage 35% CO conversion # 36% CO, formation # 0% CH, formation # 10%

End gas, second stage: 19,000 m3 of the composition COs CO Ha CHA No.

33.0 20.4 20.8 7.0 18.2

The proportion of CO: He in the gas is raised by the addition of converted gas to 1:2, requiring
7.200 m³ converted gas = 5,500 m³ water gas

Producing

25,400 " synthesis gas of the composition:

31.3 16.2 31.7 6.0 14.8

Total water gas imput so far:

With a required input of 60,000 m³, there remain available which are equal to

of the following composition:

5.800 m³ synthesis gas

13.2 18.0 35.3 5.0 13.4

This gas is used in the one-stage low pressure synthesis. With a contraction of 33%, the 32,200 m3 of synthesis gas will produce 21,400 m3 end

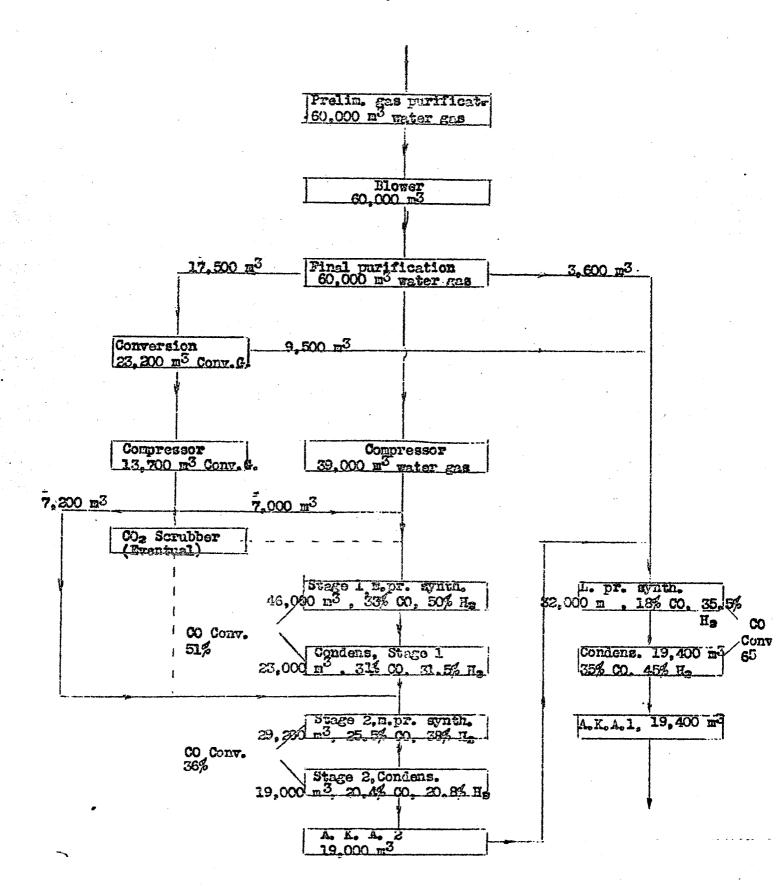
46.0 7.5 15.0 11.5 20.0

No scrubbing cut of CO₂ from the converted gas is foreseen in the above exposition. The scrubbing of the converted gas used in the medium pressure synthesis, 1-st and 2-nd stages, must be taken into consideration. Scrubbing of the converted gas for the low pressure stage must be left out of consideration, because the capacity of the three compressors, 52,000 m³, is practically reached,

When making the decision on whether the proposed method or the cld method are preferable in production, the following must be considered. The gas input and the reactor capacity are left unchanged. In case I, a considerably greater amount of CO is put in and converted in the medium pressure installation under more favorable conditions (CO: $H_2 = 1:1.5$). The low pressure stage with 720 m²/h/ reactor remains a safe stabilizer for the operation of the medium pressure installation.

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Connecting the medium pressure and the low pressure synthesis in series. 60,000 m³ water gas intake



CONNECTING THE MEDIUM PRESSURE AND THE LOW FRESSURE SYNTHESIS IN SERIES. 65,600 m³ HATER GAS INTARE

