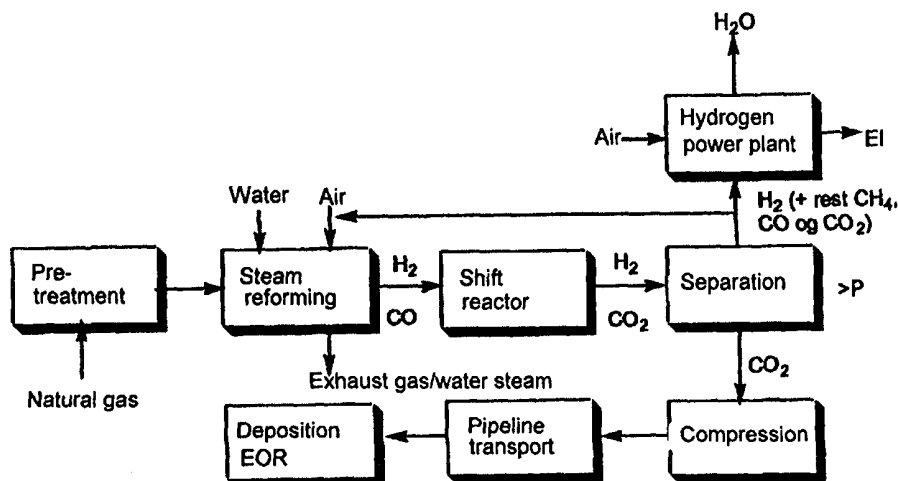




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(21) International Application Number: PCT/NO99/00280 (22) International Filing Date: 14 September 1999 (14.09.99) (30) Priority Data: 19984295 16 September 1998 (16.09.98) NO (71) Applicant (for all designated States except US): DEN NORSKE STATS OLJESELSKAP A.S [NO/NO]; N-4035 Stavanger (NO). (72) Inventors; and (75) Inventors/Applicants (for US only): OLSVIK, Ola [NO/NO]; Haukv. 18, N-7562 Hundhammeren (NO). HANSEN, Roger [NO/NO]; Sildråpev. 70B, N-7048 Trondheim (NO). GRISLINGÅS, Arne [NO/NO]; Ole Nordgaardsvei 36B, N-7049 Trondheim (NO). RYTTER, Erling [NO/NO]; Steinåsen 19, N-7049 Trondheim (NO). (74) Agent: BRYN & AARFLOT AS; P.O. Box 449 Sentrum, N-0104 Oslo (NO).		(81) Designated States: AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CR, CU, CZ, DE, DK, DM, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW, ARIPO patent (GH, GM, KE, LS, MW, SD, SL, SZ, TZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i> <i>Before the expiration of the time limit for amending the</i> <i>claims and to be republished in the event of the receipt of</i> <i>amendments.</i> <i>In English translation (filed in Norwegian).</i>

(54) Title: PROCESS FOR PREPARING A H₂-RICH GAS AND A CO₂-RICH GAS AT HIGH PRESSURE



shows known technique in the form of steam reforming with hydrogen firing

(57) Abstract

The present invention comprises a method for production of a CO₂-rich gas stream and a H₂-rich gas stream, the method comprising the following steps: a) natural gas and water are fed to a reforming reactor and are converted to synthesis gas under supply of a O₂-containing gas; b) the gas stream from a) is shifted, whereby the content of CO is reduced and the amounts of CO₂ and H₂ are increased by reaction of H₂O; c) the gas stream from b) is separated in a separation unit into a CO₂-rich and a H₂-rich gas stream, respectively. The invention also concerns the use of a CO₂-rich gas stream for injection into marine formations, and the use of a H₂-rich gas stream for hydrogenation, as a source of energy/fuel in fuel cells or for production of electricity.

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Method for preparing a H₂-rich gas and a CO₂-rich gas at high pressure.

The present invention comprises a method for production of a CO₂-rich gas for injection purposes or to be deposited, and a hydrogen-rich gas, as well as use thereof.

It is commonly assumed that the greenhouse effect and the climate on earth are closely connected to human made emissions of CO₂. These emissions are primarily formed by combustion of coal and hydrocarbons, i.a. by generation of heat and electric power. A desirable goal is therefore to reduce the emission of CO₂ to the atmosphere.

It is known art to reduce the emission of CO₂ from combustion of natural gas, e.g. by gas reforming and shift technology for preparation of a mixture consisting of hydrogen and carbon dioxide. These components are then separated, whereafter hydrogen is used as fuel in a gas turbine and carbon dioxide is deposited after compression to desired pressure. The deposition can be made on the bottom of the sea or in geological reservoirs. The reservoirs can also contain hydrocarbons. The above mentioned technique is i.a. described in Teknisk Ukeblad No. 16, page 8, 1998.

Known art comprising gas reforming and shift technology as described above is very expensive while providing less energy yield than a conventional, modern gas power plant.

US 3,652,454 describes preparation of CO₂ and H₂ from a gas stream containing CO by an improved continuous catalytical shift reaction at high pressure. The reaction takes place in one or more shift reactors at a superatmospheric pressure of from 35 to 250 atmospheres, and a temperature between 287°C and 537°C. The patent does not describe reforming of natural gas.

From EP 0 000 993-A1 it is known a method for preparation of ammonia by means of a primary and a secondary catalytic reforming of an hydrocarbon stream at superatmospheric pressure. From the primary catalytic reforming the ratio of steam to carbon is from 2.5 to 3.5, the pressure is from 30 to 120 bar and the temperature out of the reactor is from 750 to 850°C. From the secondary catalytic reforming the content of methane is from 0.2 to 10 % by weight on a dry basis and the ratio of hydrogen to nitrogen is from 2.2 to 2.7. To the secondary reforming there is added an excess of air for preparing a gas with a higher content of methane, i.e. at a lower temperature, and/or a lower steam ratio and/or a higher pressure. In the above mentioned EP patent CO₂ is re-

moved at a low pressure by taking out hydrogen at an elevated pressure for further use by the preparation of ammonia.

EP 0 289 419 describes catalytic steam reforming of hydrocarbons for preparing hydrogen in an ammonia process. The catalytic steam reforming takes place at a pressure from 25 to 120 bar, a temperature from 800 to 1000°C and at a ratio steam:carbon of 1.8-2.5. The process is operated in such a way that there are less than 0.3% impurities in the H₂-rich gas which is to be used for production of ammonia. The present invention allows a higher content than 0.3% of CO, CO₂ and CH₄ in the H₂-rich gas stream.

CA 868,821 describes the preparation of synthesis gas by steam reforming of hydrocarbons in a gas and a liquid at 50-250 absolute atmospheres, preferably 160 abs. atm. for production of ammonia and methanol.

Known art does not deal with a simplified plant as comprised by the present invention, where natural gas is reformed at high pressure in an autothermal reformer followed by a shift reactor, whereafter a CO₂-rich gas is provided at a pressure in the interval from 1 to 100 bar, which leads to reduced compression costs by deposition and reinjection. The simplified plant results in that a limited part of the carbon in the raw material is in the form of methane or carbon monoxide when the hydrogen-rich gas mixture is fed as a fuel to a gas power plant. A result of the reduced demand to the purity of the product gas involves that one or more of the process steps can be operated at a higher pressure, a lower temperature and/or a lower ratio steam/carbon.

The present invention comprises a method for production of a CO₂-rich gas stream and a H₂-rich gas stream, whereby the method is characterized by the following steps:

- a) natural gas and water are fed to a reforming reactor and are converted to synthesis gas under supply of a O₂-containing gas;
- b) the gas stream from a) is shifted, whereby the content of CO is reduced and the amounts of CO₂ and H₂ are increased by reaction of H₂O;
- c) the gas stream from b) is separated in a separation unit into a CO₂-rich and a H₂-rich gas-stream, respectively.

It is a further object of the present invention that the shift process in b) is carried out in one step where the ratio H₂O:CO in the shift process can be from 1 to 9, preferably 1.5 to 4. In the present invention the pressure in the CO₂-rich gas stream after the separation unit is from 1 to 100 bar, preferably from 5 to 50 bar. The carbon part in the H₂-rich

gas stream is from 1 to 20 % by volume, preferably from 5 to 15 % by volume. The invention is characterized in that to the natural gas in step a) it is added an oxygen rich gas or air/oxygen enriched air stream. The reforming reactor according to the invention is preferably a partial oxidation reactor, especially an autothermal reformer. The gas stream from a combustion chamber in an autothermal reformer is brought in contact with a catalyst bed. When the reforming takes place in an partial oxidation reactor, the reaction is carried out without a catalyst. Further, the gas stream out of the reformer has a temperature within the interval from 800 to 1200°C. When the natural gas in step a) is supplied with air/oxygen enriched air at least a part of N_2 follows the CO_2 -rich gas stream through the process. The present invention also concerns the use of the CO_2 -rich gas stream prepared according to the previously mentioned method, where the CO_2 -rich gas stream is injected into marine formations. Further, the invention comprises the use of the H_2 -rich gas stream prepared according to the invention, where the H_2 -rich gas stream can be utilized for hydrogenation, in the production of electricity and as a source of energy / fuel in fuel cells.

An important feature of the invention is that the CO_2 -rich stream is present at an elevated pressure, which means that the compression work which is necessary to bring the pressure up to the injection pressure or deposition pressure (50-250 bar) becomes as little as possible. This requires a high pressure through all units (reformer, shift unit and separator unit). As an example the natural gas which is fed into the reformer can have a pressure of 30-200 bar. By a marginal pressure drop in the reformer and the shift unit, the pressure into the separation unit will be within the same interval.

The present invention provides several advantages when producing a H_2 -rich and a CO_2 -rich gas, respectively. One of the advantages is that increased pressure gives increased reaction for each volume unit in reformer and shift reactor, and further a reduced compression work of CO_2 by a possible reinjection or deposition. In addition, increased pressure gives reduced compression work for feed to the gas turbine. In the reformer there is high primary formation of CO_2 as a result of high pressure. Another advantage of the present invention is a simplified design and operation of a possible a gas turbine for combustion of the H_2 -rich gas stream as a result of a certain slip of CH_4 in the reformer, CO in the shift reactor and CO_2 in the separation step. The use of air or oxygen enriched air in the case of an autothermal reformer (ATR) and partial oxidation (POX) give savings in oxygen units and gives a simplified design and operation of ATR and gas turbine. A cer-

tain CO slip, i.e. not reacted gas component, makes low temperature shift and methanation unnecessary. In the present invention it might be necessary to supply steam to the shift reactor. A certain slip of CO₂ in the present invention simplifies the separation process. If the separation is carried out with amine washing, a moderate pressure release might be sufficient.

The present invention comprises a number of embodiments where i.a. natural gas is cleaned and pre-treated in a suitable manner so that the gas feed mainly will consist of methane after treatment. The reforming takes place at a pressure within the interval 30 to 200 bar. It is preferred to use a shift reactor when carrying out the present invention.

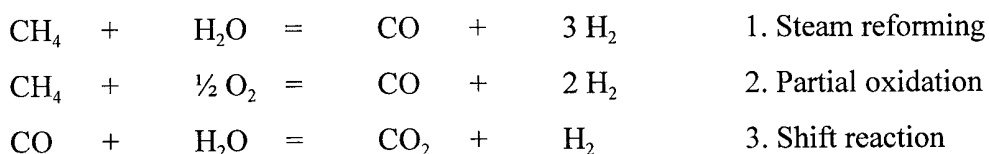
Steam can be supplied to the shift reactor, but it can also be operated without supply of steam. At the outlet of the shift reactor a content on carbon basis of CO up to 2 % by volume, preferably 5 % by volume, and methane with 2 % by volume, preferably 5 % by volume, is acceptable. Further, the H₂-rich gas stream according to the invention can contain at least 1 % by volume and up to maximum 20 % by volume of the original amount of carbon in the natural gas. The steam reformer can be fired with the hydrogen rich fuel gas. If autothermal reforming is used at least a part of the hydrogen rich stream can be used for pre-heating of the feed gas. Pre-heating of the feed gas can also be carried out by heat exchange with warm gas from a gas turbine.

The object of the present invention is that a limited part of the carbon in the feed stream/raw material is in the form of methane, carbon monoxide or carbon dioxide when the hydrogen rich gas mixture is fed as fuel to a gas power plant. In the present invention reforming of natural gas, shift of carbon monoxide to carbon dioxide and separation of the gas mixture into CO₂-rich and H₂-rich gas, respectively, takes place at conditions which have not been described in known technique. The mentioned conditions are also outside of those which is accepted and recommended in the petrochemical industry.

Because the method is characterized by a reduced requirement to methane and carbon monoxide in the feed stream/raw material which i.a. is passed to a gas power plant, this will involve that one or more of the process steps can take place at higher pressure than normal for reforming of natural gas by autothermal reforming (ATR), combined reforming (CR) or partial oxidation (POX). The fact that reforming and shift can take place at a higher pressure leads to a reduction in capital costs and in compression energy for CO₂ before deposition. It is also possible to separate out and compress CO₂ in liquid form,

which can give further savings. In the present invention the reforming and shift reactions can be carried out at lower temperatures than those being necessary/required at a given pressure. Lower temperatures at a given pressure also lead to savings. Further it can be unnecessary with both low temperature shift and methanation of residual carbon monoxide, which is standard technology today in production of hydrogen for synthesis of ammonia. A method to separate CO₂ can be by an amine washing process, which in the present invention can be carried out by a simplified method by using pressure release instead of steam stripping.

The following chemical reactions take place by production of synthesis gas and hydrogen by reforming of natural gas:



The heat of reaction for the strongly endothermic steam reforming can be provided either by external heating, as in a steam reformer, or by combination with the partial oxidation in an autothermal reformer.

In steam reformer (SR) natural gas (methane) is converted in a tube reactor at high temperature and relatively low pressure. A steam reformer consists of many reactor tubes, i.e. 200 – 250 tubes with relative lengths of 12-13 metres, inside diameter of about 10 cm and an outside diameter of about 12 cm. This is a space demanding unit with a length of 30 – 50 metres, width of 10-12 metres and a height of 15 – 20 metres. Conventional steam reformers are operated in the pressure range from 15 to 30 bar. The outlet temperature of the gas from a conventional steam reformer lies in the temperature area 950°C. The energy which is used to carry out the endothermic reactions is supplied by external firing/heating (top-, side-, bottom- or terrace-fired). The ratio between steam and carbon is from 2.5 to 3.5, and the ratio between hydrogen and carbon monoxide in the product stream is from 2.7 to 3.0. A typical synthesis gas produced from a steam reformer contains approximately 3 % by volume of methane.

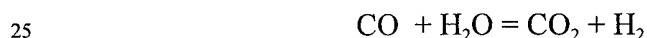
In the present invention the reforming of natural gas (equation 1 and 2 above) can take place in an autothermal reformer (ATR). In an ATR natural gas (methane) is fed together with oxygen/air into a combustion chamber. The energy which is required to operate the endothermic steam reforming reactions is provided by the exothermic reaction

ons between hydrocarbons/hydrogen and oxygen/air. The temperature i the combustion chamber can reach more than 2000°C. After the combustion chamber the reactions are driven to equilibrium over a catalyst bed before the synthesis gas is leavig the reactor at approximately 1000°C. The size of such a unit could be a height of 10-15 metres and a
5 diameter of 5-6 metres. Typical ratio of steam:carbon is from 0.6 to 1.4. The ratio between hydrogen and carbon monoxide is lower than 2. Typical methane slip is 1-2% by volume i the product stream. The ATR can be operated at higher pressure than the SR.

The present invention can also take place in a partial oxidation reactor (POX) which also is an autothermal reformer except that the unit does not comprise a catalyst
10 bed. This unit is often somewhat bigger if the same capacity compared to an ATR shall be maintained. The reason for a somewhat bigger reformer is that the ATR has catalyst which make the endothermic reactions more rapid. The exit temperature for a POX is also somewhat higher than for a typical ATR.

Reforming of natural gas can also be made by combined reforming (CR) which is
15 a combination of a steam reforming (SR) and an autothermal reforming (ATR). A combination of SR and ATR makes it possible to adjust the composition out of the reformer unit by regulating the efforts on the two reformers. SR is in CR operated at «milder» conditions, i.e. lower outlet temperature, which leads to a high methane slip. The residual methane is reacted in the ATR. The ratio steam:carbon is in the area 1.8-2.4, with a ratio between
20 hydrogen and carbon monoxide in the product gas higher than 2.

After reforming of the natural gas the gas mixture according to the present invention is shifted. The gas mixture from the reformer reactor contains mainly the gas components CO, H₂, H₂O, CO₂ and some CH₄. Between these components there is an equilibrium relation given by the stoichiometric equation:



This reaction is called the water shift reaction, and by operating a shift reactor at certain conditions the equilibrium can be forced to the right and a gas mixture is obtained which is rich in hydrogen and carbon dioxide, and where the concentration of carbon monoxide is low. Sufficient reaction velocity is provided by use of suitable catalysts, and in processes where a high degree of reaction of CO is desirable (e.g. ammonia synthesis) two fixed
30 bed reactors are used in series, high temperature shift reactor and low temperature shift reactor, respectively. Two steps are chosen because the equilibrium is favoured by low

temperature, whereas the reaction velocity is favoured by high temperature. By selecting two reactors working in series, a smaller total reactor volume is achieved. The process is nearly pressure independent and normally the same pressure as in the reformer is used. Typical temperature out of the first reactor is 420°C and out of the second reactor 230°C. Catalyst in the first step is normally based on chromium/iron, whereas the catalyst in the second step normally is a copper/zinc catalyst.

In the shift unit CO and H₂O are reacted to CO₂ and H₂, and in known technique it is often a requirement that the mentioned reaction to a highest possible degree is driven to the right, so that as little CO as possible is present in the gas mixture out of the shift unit. A low content of CO in the mentioned gas mixture again gives a high purity of the H₂-rich gas stream out of the separation unit. In known art the ratio H₂O:CO is usually equal to 10:1, which gives a high consumption of H₂O, for thereafter to obtain a high purity of the H₂-rich gas. In the present invention it is, however, acceptable with more impurities in the H₂-rich gas, which implies a lower consumption of H₂O and again a simpler shift process. These are factors which gives total cost savings by the production of a CO₂-rich and H₂-rich gas stream. In the present invention it is acceptable with a ratio H₂O:CO of from 1 to 9, preferably from 1.5 to 4.

If the requirement to reaction of CO₂ is moderate, which will be the case if the gas mixture shall be burned for energy purposes, the shift reaction can be carried out in one step. A simple high pressure shift reactor can then be selected.

Gasses in mixture can be separated more or less completely based on the different properties of the gas molecules. The most common principles are absorption, adsorption, semipermeable membranes and kryogenic distillation.

CO₂ is an acid gas, and the most widely used method to separate the mentioned gas from other non-acid gas molecules is absorption. During absorption the different chemical properties of the gas molecules are utilized. By contacting the gas mixture with a basic liquid the acid gases will to a high degree be dissolved in the liquid. The liquid is separated from the gas and the absorbed gas can then be set free either by altering the composition of the liquid or by altering pressure and temperature. For separation of CO₂ mainly aqueous solutions of alcoholamines are used. The absorption is taking place at a relatively low temperature and a high pressure, while stripping of the gas from the liquid is carried out at a relatively high temperature and low pressure. To liberate CO₂ from the

amine phase in the stripping unit, stripping steam is usually used. If the partial pressure of CO_2 in the gas into the absorber is high, e.g. higher than 15 bar, it is possible to obtain high concentrations in the amine phase, and a large part of absorbed CO_2 can be set free in the stripping unit at elevated pressure, e.g. 5-8 bar.

5 By the use of one or more semipermeable membrane units it is possible to achieve that molecules of different molecular weight and different properties permeate the membrane at different velocities. This principle can be utilized to separate gases. For the gas mixture in question membranes can be selected where H_2 permeates rapidly, whereas CO_2 permeates slowly, whereafter a separation-in-part of the different gas components is
10 achieved. By combining solid membranes and liquid membranes it is also possible to achieve a rapid permeation of CO_2 , while H_2 is kept back. It can be difficult to achieve complete separation of the different gas components by using different separation methods. This is especially the case by the use of membranes. For gas mixtures which are to be burned, a partly separation of hydrogen and CO_2 will be sufficient.

15 In a gas power plant normally natural gas is burned wherein the main component is methane, and air in a combustion chamber at elevated pressure. The combustion gases are driving the turbines which provide necessary shaft moment to the compressor part (which compresses air to the pressure of the combustion chamber) and to operate the generator or other mechanical equipment. Mixing in of natural gas takes place in burners
20 and design/construction of these is important in order to obtain correct flame temperature and to prevent formation of undesired combustion products. If a hydrogen rich gas is used instead of a natural gas, more conditions which are necessary for the construction of burners, combustion chamber and turbine, will be changed. The most important are combustion energy, flame propagation velocity, explosion range and auto ignition temperature.
25 re. These are conditions which lead to the fact that a hydrogen rich gas cannot be used in a gas turbine which is constructed for combustion of methane without modifications. The experience in burning hydrogen rich gases is limited, and the technology is available to a delimited extent.

In the present invention it is desirable to deposit out-separated CO_2 . Large
30 amounts of CO_2 can be deposited according to various methods, of which the three most interesting are deposition at very deep oceans, deposition in deep water reservoirs and deposition in oil reservoirs wherein the gas at the same time functions as drive agent for

enhanced oil recovery. The two last mentioned storage methods are operated commercially. In these storage forms the CO_2 gas has to be brought to high pressure for transport in pipelines to a deposition well and further to injection. The injection pressure will vary, but could be in the range 50 to 300 bar. If the CO_2 gas can be separated from the H_2/CO_2 mixture at an elevated pressure, significant compression work can be avoided, and this is the case in the present invention.

The attached figures show known technique and different embodiments of the present invention.

Figure 1 shows known technique in the form of steam reforming with hydrogen firing.

Figure 2 shows POX or ATR with oxygen where natural gas (1) is pre-treated before this (2) and is passed into an ATR or a POX where water (3) and oxygen gas (4) are added. The reformed gas mixture (5) is thereafter passed to a shift reactor/shift unit whereafter the gas mixture (6) is passed to a separation unit where a H_2 -rich gas (10) and a CO_2 -rich gas (7) are provided. Stream (7) can possibly be compressed further before deposition/reinjection. Stream (10) can be utilized i.a. a hydrogen power plant.

Figure 3 shows POX or ATR with air. This is in principle the same process as described in Figure 2, but in this case air is used as oxidant in the reformer instead of oxygen. This gives somewhat milder conditions than those present by the use of oxygen in the burner. By the use of air relatively big amounts of nitrogen gets into the system and the partial pressure of CO_2 will be considerably lower. This again will render it more difficult to take out CO_2 at an elevated pressure in the separation process. The ratio between H_2 and N_2 will be approximately 1:1.

Example 1

The ATR reactor can be operated under different operating conditions. In the tables below the composition of the gas at the outlet of the reactor is shown for different pressures, temperatures and steam/carbon ratios. In all calculations the ratio oxygen/carbon is set at 0.6. Feed gas has the composition $\text{CO}_2=2.5$; $\text{CH}_4=82$; $\text{C}_2=9$; $\text{C}_3=5$; $\text{C}_{3+}=1.5$.

Table 1 shows composition in mole% in dry gas at a pressure of 80 bar

Steam/carbon	S/C=0.5			S/C=1.8			S/C=2.8		
Temp. °C	800	1000	1200	800	1000	1200	800	1000	1200
H ₂	40.9	59	65	52.9	65.6	66.8	58.8	67.9	69.2
CO	13.4	25.9	36.3	10.7	20.8	27.1	9.4	17.7	23.2
CO ₂	17.4	7.3	3.8	19	10.9	6.1	19.7	13.1	7.6
C ₁	28.2	7.8	1.0	17.5	2.7	0	12.1	1.3	0

Table 2 shows composition in mole% in dry gas at a pressure of 120 bar

Steam/carbon	S/C=0.5			S/C=1.8			S/C=2.8		
Temp. °C	800	1000	1200	800	1000	1200	800	1000	1200
H ₂	36.6	55.7	64	48.8	63.8	67.0	56	-	69.2
CO	11.7	-	30	9.4	19.9	24.6	7.8	19.4	23
CO ₂	19.4	6.4	4	20.1	11.6	8.0	-	9.6	7.8
C ₁	32.8	11.2	1.9	21.6	4.7	0.4	18.4	0	0

5

Example 2

Example 2 comprises a one step shift reactor. The gas mixture into the shift reactor can have a varying composition as is evident from the Tables 1 and 2. If the gas composition in Table 1 is used at 1000°C and S/C of 1.8, this will correspond to a steam/CO ratio out of the ATR reactor of about 3.0. Table 3 below shows a possible composition of feed gas and outcoming gas from a one step shift reactor having an outlet temperature of 400°C.

10

Table 3 shows the gas composition from an one step shift reactor given in moles/100 moles dry gas.

Component	Feed gas	Leaving gas
H ₂	65.6	70.8
CO	20.8	2.4
CO ₂	10.9	24.5
C ₁	2.7	2.3
H ₂ O	62.4	37.3

Example 3

The example comprises a two step shift reactor. If a two step shift reactor is used, higher degree reaction of CO will be obtained. Table 4 shows a possible gas composition for a two step process where the outlet temperature in the first reactor is 400° C and in the second reactor 250°C. Inflowing gas has the same composition as in example 2.

Table 4: Possible gas composition for a two step process given in moles/100 moles dry gas.

Component	Feed gas	Leaving gas
H ₂	65.6	71.5
CO	20.8	0.3
CO ₂	10.9	26.0
C ₁	2.7	2.2
H ₂ O	62.4	35

Example 4

Separation of CO₂ by absorption in ethanolamine solution (amine washing) is described in this example. The gas mixture out from the shift reactor contains significant

amounts of steam. After cooling to e.g. 40°C most of the steam will be condensed out, and below two types of possible gas mixtures from the reforming process are shown:

Table 5: Possible gas compositions from the reforming process

Component	Gas 1	Gas 2
CH ₄	3 mol%	3 mol%
CO	2 mol%	2 mol%
CO ₂	17.7 mol%	29 mol%
H ₂	62.3 mol%	66 mol%
Total pressure	40 bar	80 bar

5 These gas mixtures can be fed to an absorption column where an aqueous methanolamine solution is used as absorption liquid. The amine solution will absorb the main part of CO₂, and the gas out of the absorption column will mainly consist of hydrogen. The CO₂ rich amine solution is fed to a stripping unit where the pressure is e.g. 8 bar and the temperature is e.g. 120°C. A part of the CO₂ gas will be liberated because of the temperature increase and the pressure reduction. Further CO₂ can be set free from the amine solution by stripping with steam.

10 Table 6 below shows a possible composition of the gas phase out from the absorption column and out from the stripping unit when gas of type 2 in Table 5 is used as feed and only pressure reduction and temperature increase is used and no stripping steam.

15

20

Table 6 shows the composition of gas from absorber unit and stripping unit without use of stripping steam.

Component	Out from absorption column	Out from stripping column
CH ₄	4 mol%	-
CO	3 mol%	-
H ₂ O	-	30 mol%
CO ₂	3 mol%	70 mol%
H ₂	90 mol%	-
Total pressure	80 bar	8 bar
Temperature	50°C	120°C

Example 5

Example 5 concerns separation by condensation. By cooling of a CO₂-rich gas CO₂ could be condensed out as a liquid or precipitated as a solid (CO₂ ice). At a temperature of -57°C the freezing point of CO₂ is reached and the vapour pressure of CO₂ is then 5.1 bar. If CO₂ gas is further cooled, solid CO₂ is obtained and at -78°C the vapour pressure over solid CO₂ is 1 bar.

Gas of type 2 in Table 5 is dried completely free of water before it is treated in a condenser unit consisting of more cooling steps, wherein the temperature of the gas in the first cooling step is reduced to about -55° C. At this temperature a big part of the CO₂ will condense out and the gas composition out from the condenser can be as shown in column 2 in the table below:

Table 7 shows the composition of gas phases in a two step cooling process

Component Composition of gas	Composition of gas	
	From cooling step 1	From cooling step 2
CH ₄	3.5 mol%	3.8 mol%
CO	2.3 mol%	2.5 mol%
N ₂	0.2 mol%	0.2 mol%
CO ₂	7.5 mol%	1.0 mol%
H ₂	86.7 mol%	92.5 mol%
Total pressure	80 bar	80 bar

In this example about 70% av CO₂ in the incoming gas will be condensed out in cooling step 1. By lowering the temperature further in a new process unit (cooling step 2) to e.g. -80°C the gas composition could be as shown in column 3 in Table 7 above. By means of the two cooling steps 96% of CO₂ in the incoming gas can be removed. The out-condensed liquid and the precipitated solid will be nearly pure CO₂.

Example 6

This example describes compression of CO₂. The gas phase from the stripping unit in example 4 contains steam and CO₂. By cooling of this gas to e.g. 40°C, most of the water vapour will be condensed out and the nearly pure gas at a pressure of 8 bar can be compressed to a desired shipment pressure for transport of CO₂ in pipeline.

Example 7

The example concerns one or more membrane units where the CO₂-rich gas can have a pressure approximately equal to the partial pressure of CO₂ into the separation unit, as shown in Table 1 below.

Total inlet pressure on the separation unit (bar)	Partial pressure of CO ₂ out from the separation unit (bar)
200	40
250	50
300	60

Claims

1. Method for production of a CO₂-rich gas stream and a H₂-rich gas stream,
characterized in that the method comprises the following steps:
 - 5 a) natural gas and water are fed to a reforming reaktor and converted to synthesis gas under supply of a O₂-containing gass;
 - b) the gas stream from a) is shifted, whereby the content of CO is reduced and the amounts of CO₂ and H₂ are increased by reaction of H₂O;
 - c) the gas stream from b) is separated in a separation unit into a CO₂-rich and a H₂-
10 rich gas stream, respectively.
2. Method according to claim 1,
characterized in that the shift process in b) is carried out in one step.
- 15 3. Method according to claims 1-2,
characterized in that the ratio H₂O:CO in the shift process is from 1 to 9.
4. Method according to claims 1-3,
characterized in that the ratio H₂O:CO in the shift process preferably is from
20 1.5 to 4.
5. Method according to claims 1-4,
characterized in that the pressure in the CO₂-rich gas stream after the separation
unit is from 1 to 100 bar.
25
6. Method according to claims 1-5,
characterized in that the pressure in the CO₂-rich gas stream after the separation
unit is from 5 to 50 bar.
- 30 7. Method according to claims 1-6,
characterized in that the carbon part in the H₂-rich gas stream is from 1 to 20 %
by volume.

8. Method according to claims 1-7,
characterized in that the carbon part in the H₂-rich gas stream is from 5 to 15 %
by volume.
- 5 9. Method according to claims 1-8,
characterized in that the natural gas in step a) is supplied with an oxygen rich
gas.
- 10 10. Method according to claims 1-8,
characterized in that the natural gas in step a) is supplied with air/oxygen en-
riched air.
- 15 11. Method according to claims 1-8,
characterized in that the reformer reactor preferably is a partial oxidation reac-
tor.
- 20 12. Method according to claims 1-11,
characterized in that the reformer reactor particularly is an autothermal refor-
mer.
13. Method according to claim 12,
characterized in that the gas stream from a combustion chamber in an
autothermal reformer is contacted with a catalyst bed.
- 25 14. Method according to claim 11,
characterized in that the reforming is carried out without a catalyst.
- 30 15. Method according to claims 1-14,
characterized in that the gas stream out of the reformer has a temperature wit-
hin the interval from 800 to 1200°C.

16. Method according to claim 10,
c h a r a c t e r i z e d i n that at least a part of N_2 follows the CO_2 -rich gas stream.

17. Use of a CO_2 -rich gas stream produced according to claim 1 for injection into ma-
5 rine formations.

18. Use of a H_2 -rich gas stream produced according to claim 1 for hydrogenation.

19. Use of a H_2 -rich gas stream produced according to claim 1 as a source of energy /
10 fuel in fuel cells.

20. Use of a H_2 -rich gas stream produced according to claim 1 for the production of
electricity.

1/2

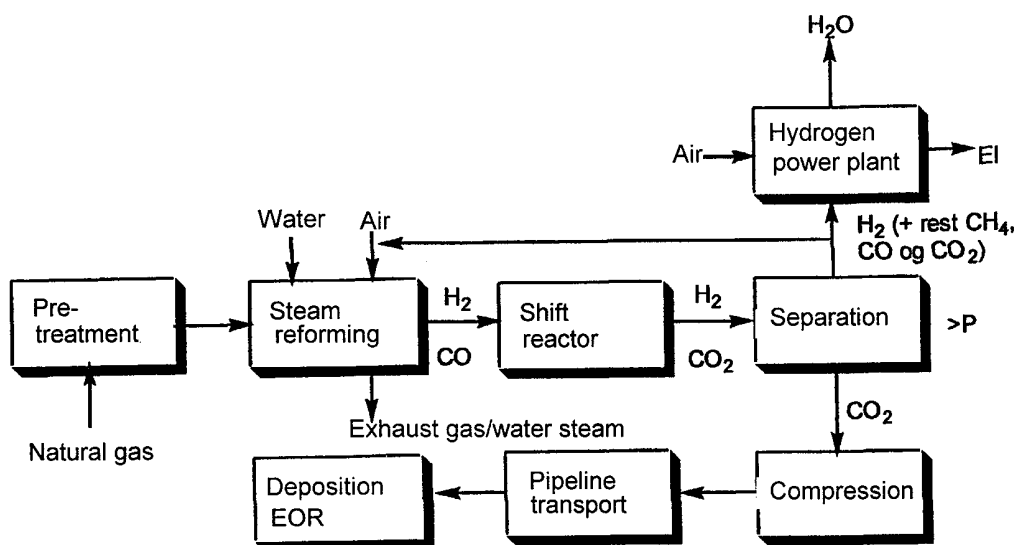


Figure 1 shows known technique in the form of steam reforming with hydrogen firing

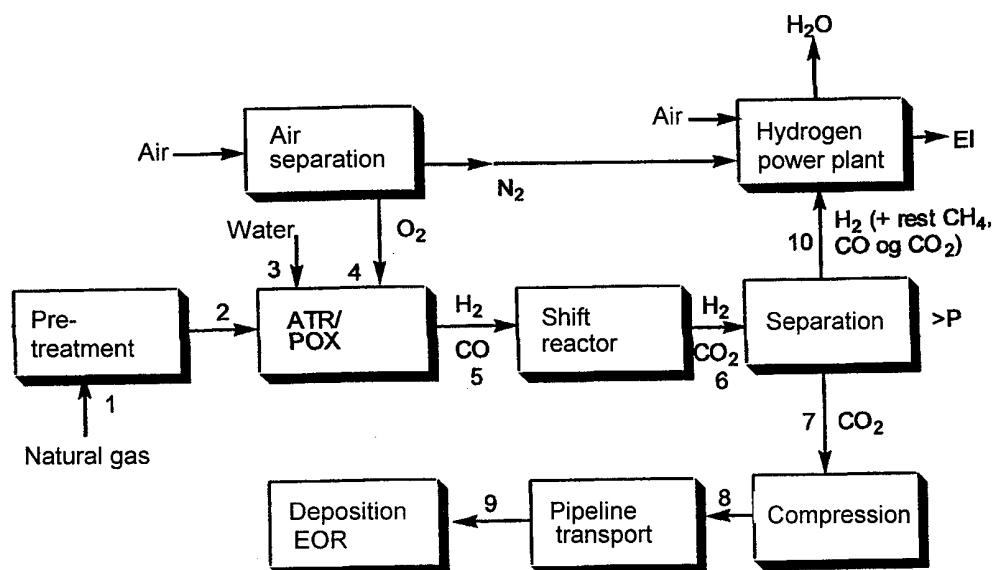


Figure 2 shows POX or ATR with oxygen

2/2

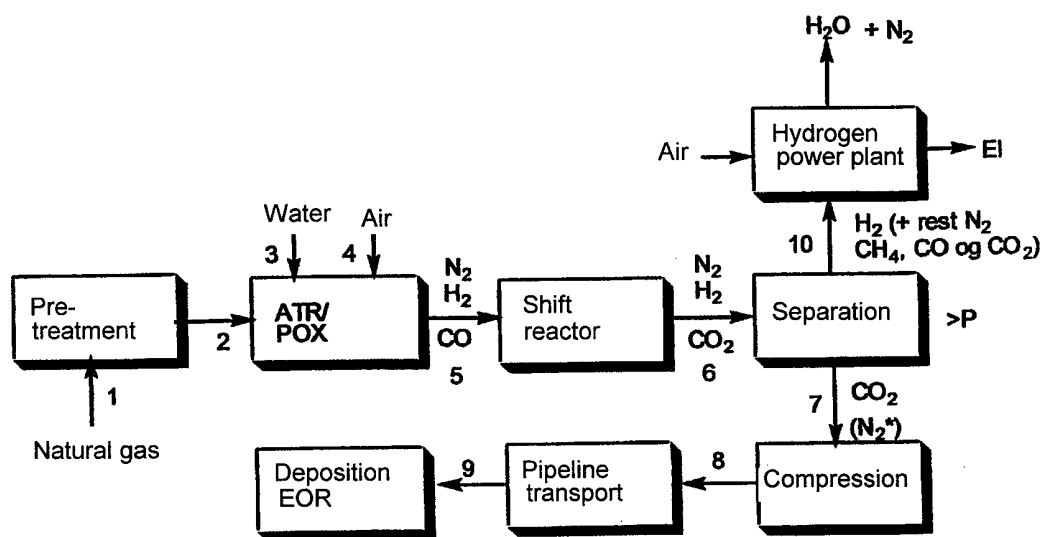


Figure 3 shows POX or ATR with air

(N_2^*): By use of one or more membrane units
 N_2 is run together with CO_2 .

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NO 99/00280

A. CLASSIFICATION OF SUBJECT MATTER

IPC7: C01B 3/32, C01B 31/20

According to International Patent Classification (IPC) or to both national classification and IPC

B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

IPC7: C01B

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

SE,DK,FI,NO classes as above

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

FULLTEXT, EPODOC, WPI

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 0289419 A1 (COMPAGNIE FRANCAISE D'ETUDES ET DE CONSTRUCTION"TECHNIP"), 2 November 1988 (02.11.88), column 1, line 15 - line 40; column 2, line 16 - line 38; column 3, line 4 - line 13, column 4, line 48 - line 51; column 4, line 63 - column 5, line 4; abstract --	1-16
X	US 5714132 A (AKHILESH KAPOOR ET AL), 3 February 1998 (03.02.98), column 2, line 39 - line 45; column 5, line 22 - line 26; column 7, line 50 - column 8, line 18, figures 1,3, example II, Table II --	1-16

☒ Further documents are listed in the continuation of Box C.☒ See patent family annex.

* Special categories of cited documents:

"A" document defining the general state of the art which is not considered to be of particular relevance

"E" earlier document but published on or after the international filing date

"I" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)

"O" document referring to an oral disclosure, use, exhibition or other means

"P" document published prior to the international filing date but later than the priority date claimed

"I" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention

"X" document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone

"Y" document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art

"&" document member of the same patent family

Date of the actual completion of the international search

10 March 2000

Date of mailing of the international search report

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Name and mailing address of the ISA/

Swedish Patent Office

Box 5055, S-102 42 STOCKHOLM

Facsimile No. +46 8 666 02 86

Authorized officer

Helena Hemphälä/ELY

Telephone No. +46 8 782 25 00

INTERNATIONAL SEARCH REPORT

International application No.

PCT/NO 99/00280

C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	US 4699772 A (HANS C. DE LATHOUDER), 13 October 1987 (13.10.87), column 1, line 55 - line 64; column 2, line 6 - line 54; column 3, line 8 - line 9, abstract	1-2,5-6, 9-11,15-16
A	--	3-4,7-8, 12-14
A	US 4681745 A (ALWYN PINTO), 21 July 1987 (21.07.87), column 1, line 11 - line 21, Table 1 & 2	1-16
A	CA 868821 A (GIORGIO PAGANI), 20 April 1971 (20.04.71), column 3, line 20 - line 25; column 4, line 11 - line 22	1-16
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INTERNATIONAL SEARCH REPORT

International application No.
PCT/NO99/00280

Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international search report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:
because they relate to subject matter not required to be searched by this Authority, namely:

2. ☐ Claims Nos.:
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:

Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

See extra sheet

1. ☐ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:

4. ☒ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.: **1-16**

Remark on Protest

- ☐ The additional search fees were accompanied by the applicant's protest.
- ☐ No protest accompanied the payment of additional search fees.

INTERNATIONAL SEARCH REPORT

International application No.
PCT/NO99/00280

- I. Claims 1-16 relate to a process for producing a CO₂- and a H₂-rich gas.
- II. Claim 17 relates to the use of a CO₂-rich gas for injection in marine formations.
- III. Claim 18 relates to the use of a H₂-rich gas for hydrogenation.
- IV. Claim 19 relates to the use of a H₂-rich gas as a source of energy in a fuel cell.
- V. Claim 20 relates to the use of a H₂-rich gas for electricity production.

The special technical feature of Group I is a process for producing the gases H₂ and CO₂. The special technical features of Group II-V involve different applications of the gases H₂ and CO₂. The gases H₂ and CO₂, produced according to claim 1, are well known products. There is a reference in each of the independent claims 17-20 to the process for producing these known products. However, the production process does not give the products any new characteristics, therefore the reference made in claim 17-20 does not involve the features in claim 1. The group of inventions is not so linked as to form a single inventive concept under PCT Rule 13.

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/12/99

International application No.

PCT/NO 99/00280

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0289419 A1	02/11/88	FR 2614612 A,B	04/11/88
US 5714132 A	03/02/98	EP 0643013 A	15/03/95
		ZA 9405891 A	13/06/95
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		FI 865352 A	30/12/86
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		NL 8402091 A	16/01/86
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		AU 1328183 A	20/10/83
		CA 1210222 A	26/08/86
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		JP 4059249 B	21/09/92
		JP 58190821 A	07/11/83
		NZ 203799 A	13/12/85
		ZA 8302524 A	25/01/84
CA 868821 A	20/04/71	NONE	