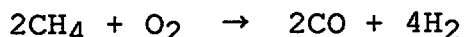


PROCESS FOR THE CATALYTIC PARTIAL OXIDATION OF  
HYDROCARBONS

The present invention relates to a process for the catalytic partial oxidation of hydrocarbons, in particular to a process for the preparation of a mixture of carbon monoxide and hydrogen from methane, natural gas, associated gas or other sources of light hydrocarbons.

The partial oxidation of hydrocarbons, for example methane or natural gas, in the presence of a catalyst is an attractive route for the preparation of mixtures of carbon monoxide and hydrogen, known in the art as synthesis gas. The partial oxidation of a hydrocarbon is a highly exothermic reaction and, in the case in which methane is the hydrocarbon, proceeds by the following reaction:



A number of process regimes have been proposed in the art for carrying out the catalytic partial oxidation reactions. One regime that is most suitable for application on a commercial scale is to contact the feed gases with the catalysts retained in a fixed arrangement, for example as a fixed bed of particles or a monolith structure. The literature contains a number of documents disclosing details of experiments conducted into the catalytic partial oxidation of hydrocarbons, in particular methane, employing a wide range of catalysts in a fixed bed arrangement.

European Patent Application publication No. 0 303 438 (EP-A-0 303 438) discloses a process for the catalytic partial oxidation of a hydrocarbonaceous feedstock in which a gaseous mixture of the hydrocarbonaceous

feedstock, oxygen or an oxygen-containing gas and, optionally, steam, is introduced into a catalytic partial oxidation zone to contact a catalyst retained therein. The catalyst employed in the process may comprise a wide range of catalytically active components, for example palladium, platinum, rhodium, iridium, osmium, ruthenium, nickel, chromium, cobalt, cerium, lanthanum and mixtures thereof. Further, it is stated in EP-A-0 303 438 that materials not normally considered to be catalytically active may also be employed as catalysts, for example refractory oxides such as cordierite, mullite, mullite aluminium titanate, zirconia spinels and alumina. The catalyst may be of a variety of forms, for example sheets of corrugated metal packed to form elongate channels therethrough or wire mesh. However, preference is given in EP-A-0 303 438 to the use of catalysts in the form of extruded honeycomb monoliths. These monoliths comprise a large number of parallel channels extending through the structure in the direction of flow of the feed and product gasses.

European Patent No. 0 262 947 (EP-B-0 262 947) discloses a process for generating hydrogen by the partial oxidation of a hydrocarbon in which a mixture of the hydrocarbon and oxygen is injected into a mass of a catalyst. The catalyst disclosed in EP-B-0 262 947 comprises platinum and chromium oxide supported on a refractory solid. The support structures described in EP-B-0 262 947 are honeycomb monolith supports, of the type used in purifying the exhausts from motor vehicles or from chemical plants, and particulate supports, preferably comprising particles having a maximum dimension of from 1 to 4 mm, for example 1.5 mm.

D.A. Hickman and L.D. Schmidt ("Synthesis Gas Formation by Direct Oxidation of Methane over Pt Monoliths", Journal of Catalysis 138, 267-282, 1992))

have conducted experiments into the partial oxidation of methane in the presence of catalysts comprising either platinum or rhodium. The partial oxidation reactions were conducted at substantially atmospheric pressure and at temperatures in the range of from 600 to 1500 K (337 to 1237 °C). The catalysts employed were in the form of metal gauzes, metal-coated foam monoliths and metal coated extruded monoliths. The metal gauze catalysts comprised 1 to 10 layers of gauzes of either 40 mesh (40 wires per inch) or 80 mesh. The foam monoliths were of alpha-alumina and described as having an open cellular, sponge-like structure. The samples employed had a nominal porosity of 30 to 50 pores per inch (ppi). The extruded monoliths were cordierite extruded monoliths, having 400 square cells/in<sup>2</sup> and consisted of straight parallel channels giving laminar flows of gases through the channels under the conditions of gas flowrate studied.

J.K. Hockmuth ("Catalytic Partial Oxidation of Methane over a monolith Supported Catalyst", Applied Catalysis B: Environmental, 1 (1992) 89-100) reports the catalytic partial oxidation of methane using a catalyst comprising a combination of platinum and palladium supported on a cordierite monolith body.

A number of academic experiments have been reported in the literature in which catalysts have been employed in the form of fixed beds of catalyst particles.

Thus, A.T Ashcroft et al. ("Selective oxidation of methane to synthesis gas using transition metal catalysts", Nature, vol. 344, No. 6264, pages 319 to 321, 22nd March, 1990) disclose the partial oxidation of methane to synthesis gas in the presence of a range of ruthenium-containing catalysts. The objective of the experiments was to establish that the partial oxidation process could be carried out under mild conditions and at

low temperatures. To this end, the experiments were conducted with a low gas hourly space velocity of 40,000/hr, a pressure of 1 atmosphere and a temperature of about 775 °C. The catalyst employed comprised small amounts of a solid, powdered catalyst.

P.D.F. Vernon et al. ("Partial Oxidation of methane to Synthesis Gas", Catalysis Letters 6 (1990) 181-186) disclose a range of experiments in which catalysts comprising nickel, ruthenium, rhodium, palladium, iridium or platinum, either supported on alumina or present in mixed oxide precursors, were applied. Again, the experiments reported are limited to a catalytic partial oxidation process employing only mild operating conditions and using small amounts of catalyst in the form of pellets retained in a fixed bed. The authors report the same experiments in "Partial Oxidation of Methane to Synthesis Gas, and Carbon Dioxide as an Oxidising Agent for Methane Conversion", Catalysis Today, 13 (1992) 417-426.

R.H. Jones et al. ("Catalytic Conversion of Methane to Synthesis Gas over Europium Iridate,  $\text{Eu}_2\text{Ir}_2\text{O}_7$ ", Catalysis Letters 8 (1991) 169-174) report the selective partial oxidation of methane using the europium iridium pyrochlore  $\text{Eu}_2\text{Ir}_2\text{O}_7$ . The reaction was studied under the mild conditions of a pressure of 1 atmosphere and a temperature of 873 K (600 °C). The catalyst was prepared by grinding and subsequent pressing to form pellets. The pelletised catalyst was packed into a porous silica frit and used directly in the experiments.

United States Patent No. 5,149,464 (US-A-5,149,464) is directed to a method for selectively oxygenating methane to carbon monoxide and hydrogen by bringing the reactant gas mixture at a temperature of about 650 °C to 900 °C into contact with a solid catalyst which is generally described as being either:

- a) a catalyst of the formula  $M_xM'_yO_z$ , where:  
M is at least one element selected from Mg, B, Al, Ln, Ga, Si, Ti, Zr and Hf; Ln is at least one member of lanthanum and the lanthanide series of elements;  
M' is a d-block transition metal,  
and each of the ratios  $x/y$  and  $y/z$  and  $(x+y)/z$  is independently from 0.1 to 8; or
- b) an oxide of a d-block transition metal; or
- c) a d-block transition metal on a refractory support; or
- d) a catalyst formed by heating a) or b) under the conditions of the reaction or under non-oxidizing conditions.

The d-block transition metals are said in US-A-5,149,464 to be selected from those having atomic number 21 to 29, 40 to 47 and 72 to 79, the metals scandium, titanium, vanadium, chromium, manganese, iron, cobalt, nickel, copper, zirconium, niobium, molybdenum, technetium, ruthenium, rhodium, palladium, silver, hafnium, tantalum, tungsten, rhenium, osmium, iridium, platinum and gold. It is stated in US-A-5,149,464 that the preferred metals are those in Group VIII of the Periodic Table of the Elements, that is iron, osmium, cobalt, rhenium, iridium, palladium, platinum, nickel and ruthenium.

The process described in US-A-5,149,464 is operated at a temperature in the range of from 650 °C to 900 °C, with a range of from 700 °C to 800 °C being preferred. A range of experiments are described in US-A-5,149,464 in which a variety of catalysts comprising Group VIII metals were tested, including ruthenium oxide, praeisidium/ruthenium oxides, pyrochlores, ruthenium on alumina, rhodium on alumina, palladium on alumina, platinum on alumina, nickel/aluminium oxide, perovskites and nickel oxide.

A similar general disclosure of a catalyst for use in the catalytic partial oxidation process is made in

International Patent Application publication No. WO 92/11199. WO 92/11199 specifically discloses experiments in which catalysts comprising iridium, palladium, ruthenium, rhodium, nickel and platinum supported on alumina were applied. All the experiments were conducted under mild process conditions, with typical conditions being a pressure of 1 atmosphere, a temperature of 1050 K (777 °C) and a gas hourly space velocity of about 20,000/hr.

The experiments described in both US-A-5,149,464 and WO 92/11199 employed catalysts in the form of solid powdered particles retained in a fixed bed arrangement by packing in a reaction tube between two plugs of silica wool.

For successful operation on a commercial scale, the catalytic partial oxidation process must be able to achieve a high conversion of the hydrocarbon feedstock at high gas hourly space velocities. Further, the selectivity of the process to the desired products of carbon monoxide and hydrogen must be high. Both these factors must be met using process equipment which is both economical to construct and economical to operate. In this respect, there exists a significant problem in operating the catalytic partial oxidation process with a fixed bed of catalyst, in that the pressure drop encountered when using the fixed bed prevents the process operating under the high gas space velocities demanded of a commercial operation. European patent application No. 92201879.1 (EP 92201879.1), as yet unpublished, discloses a process for the catalytic partial oxidation of a hydrocarbon in which the catalyst is retained in the form of a fixed bed having a high tortuosity, that is a tortuosity of at least 1.1. In this way, it has been found that only a very shallow fixed bed of catalyst is required in order to achieve a commercially acceptable

rate of conversion and yield. As the fixed bed is only very shallow, the pressure drop through the bed is very low, which in turn allows very high gas space velocities to be applied.

EP 92201879.1 discloses the use of a fixed bed arrangement in which the catalyst is present in the form of particles or as a monolith structure, such as a ceramic foam. Suitable ceramic foams are described as having from 30 to 150 pores per inch (12 to 60 pores per centimetre).

Surprisingly, it has now been found that the performance of the catalytic partial oxidation process disclosed in EP 92201879.1 can be significantly improved if the process is operated with the catalyst retained in a fixed bed arrangement meeting a very specific set of criteria. In particular, it has been found that the selectivity of the process is significantly improved if the fixed bed arrangement combines a high tortuosity with a high number of pores. Specifically, it has been found that the selectivity is significantly improved if the fixed arrangement has a tortuosity of greater than 1.1 and at least 750 pores per square centimetre. Further, it has been found that using a fixed arrangement meeting these two criteria allows the amount of catalytically active metal present in the catalyst to be reduced whilst still maintaining a high level of activity and selectivity.

Accordingly, the present invention provides a process for the catalytic partial oxidation of a hydrocarbon feedstock, which process comprises contacting a feed comprising a hydrocarbon feedstock and an oxygen-containing gas with a catalyst in a reaction zone, which catalyst is retained in the reaction zone in a fixed arrangement having a tortuosity of at least 1.1 and having at least 750 pores per square centimetre.

The process of the present invention may be used to prepare a mixture of carbon monoxide and hydrogen from any gaseous hydrocarbon or hydrocarbon feedstock having a low boiling point. The hydrocarbon feedstock is in the gaseous phase when contacting the catalyst. The process is particularly suitable for the partial oxidation of methane, natural gas, associated gas or other sources of light hydrocarbons. In this respect, the term "light hydrocarbons" is a reference to hydrocarbons having from 1 to 5 carbon atoms. The process may advantageously be applied in the conversion of gas from naturally occurring reserves of methane which contain a substantial amount of carbon dioxide. The feed preferably comprises methane in an amount of at least 50% by volume, more preferably at least 75% by volume, especially at least 80% by volume.

The hydrocarbon feedstock is contacted with the catalyst as a mixture with an oxygen-containing gas. Air is suitable for use as the oxygen-containing gas. However, the use of substantially pure oxygen as the oxygen-containing gas may be preferred. In this way, the need for handling a large volume of inert gas, for example nitrogen when using air as the oxygen-containing gas, is avoided. The feed may optionally comprise steam.

The methane-containing feed and the oxygen-containing gas are mixed in such amounts as to give an oxygen-to-carbon ratio in the range of from 0.3 to 0.8, more preferably, in the range of from 0.45 to 0.75.

References to the oxygen-to-carbon ratio refer to the ratio of oxygen in the form of molecules ( $O_2$ ) to carbon atoms present in the methane-containing feed. Preferably the oxygen-to-carbon ratio is in the range of from 0.45 to 0.7, with oxygen-to-carbon ratios in the region of the stoichiometric ratio of 0.5, that is in a range of from 0.45 to 0.65, being especially preferred. If steam is present in the feed, the steam-to-carbon ratio is

preferably in the range of from above 0.0 to 3.0, more preferably from 0.0 to 2.0. The methane-containing feed, oxygen-containing gas and steam, if present, are preferably well mixed prior to being contacted with the catalyst.

The process of the present invention may be operated at any suitable pressure. For commercial operations, elevated pressures, that is pressures significantly above atmospheric pressure, may be preferred. The process may be operated at pressures in the range of up to 150 bara. Preferably, the process is operated at pressures in the range of from 2 to 125 bara, especially from 2 to 100 bara.

The process may be operated at any suitable temperature. Under conditions of high pressure prevailing in the process, the feed is preferably contacted with the catalyst at high temperatures in order to obtain the desired degree of conversion. Accordingly, the hydrocarbon feedstock and the oxygen-containing gas are preferably contacted with the catalyst at a temperature greater than 950 °C, more preferably a temperature in the range of from 950 to 1300 °C, especially from 1000 to 1200 °C. The hydrocarbon feedstock and the oxygen-containing gas are preferably preheated prior to being contacted with the catalyst.

The hydrocarbon feedstock and the oxygen-containing gas may be provided during the process at any suitable space velocity. It is an advantage of the process of this invention that very high gas space velocities can be achieved. Thus, typical space velocities for the process (expressed as normal litres of gas per kilogram of catalyst per hour) are in the range of from 20,000 to 100,000,000 Nl/kg/hr, more preferably in the range of from 50,000 to 50,000,000 Nl/kg/hr. Space velocities in the range of from 500,000 to 30,000,000 Nl/kg/hr are

particularly suitable.

Catalyst compositions suitable for use in the catalytic partial oxidation of hydrocarbons are known in the art. Preferred catalysts for use in the process of the present invention comprise, as the catalytically active component, a metal selected from Group VIII of the Periodic Table of the Elements. References in this specification to the Periodic Table of the Elements are to the CAS version, as published in the CRC Handbook of Chemistry and Physics, 68th Edition. Preferred catalysts for use in the process comprise a metal selected from ruthenium, rhodium, palladium, osmium, iridium and platinum. Catalysts comprising ruthenium, rhodium or iridium as the catalytically active metal are most suitable for use in the process.

The catalytically active metal is most suitably supported on a carrier. Suitable carrier materials are well known in the art and include the refractory oxides, such as silica, alumina, titania, zirconia and mixtures thereof. Mixed refractory oxides, that is refractory oxides comprising at least two cations, may also be employed as carrier materials for the catalyst.

The catalytically active metal may be deposited on the refractory oxide carrier by techniques well known in the art. A most suitable technique for depositing the metal on the carrier is impregnation, which technique typically comprises contacting the carrier material with a solution of a compound of the catalytically active metal, followed by drying and calcining the resulting material.

In use in the process of the present invention, the catalyst is retained in form of a fixed arrangement. The fixed arrangement may comprise a fixed bed of catalyst particles. Alternatively, the fixed arrangement may comprise the catalyst in the form of a monolith

structure. The fixed arrangement may consist of a single monolith structure or, alternatively, may comprise a number of separate monolith structures combined to form the fixed arrangement. A most preferred monolith structure comprises a ceramic foam. Suitable ceramic foams for use in the process are available commercially.

As described hereinbefore, it is an essential feature of the process of the present invention that the catalyst is retained in a fixed arrangement having a high tortuosity. The term "tortuosity" is a common term in the art which, when referring to a fixed catalyst arrangement, can be defined as the ratio of the length of the path followed by a gas flowing through the arrangement to the length of the shortest straight line path through the arrangement. Thus, the honeycomb monolith structures described in the prior art comprising a number of straight, parallel channels extending through the structure in the direction of the flowing gas, have a tortuosity of 1.0. A typical value for the tortuosity of a fixed bed of catalyst particles is about 1.5. Ceramic foam structures may be prepared having a tortuosity of the order of 1.5 to 4.0, or even higher. In general, the tortuosity of the fixed arrangement of the catalyst for use in the process of the present invention should be in the range of from 1.1 to about 10.0, more preferably to about 5.0. A most suitable range of tortuosity is from 1.3 to 4.0.

Further, it is an essential feature of the process of this invention that the fixed arrangement of the catalyst comprises a large number of pores. In this respect, the term "pore" is a general reference to a space or interstice in the fixed arrangement between two adjacent portions of the catalyst. Thus, in the case of a fixed bed of catalyst particles, the term "pore" refers to the space between two adjacent particles. When referring to

monolith structures, for example ceramic foams, the term "pore" refers to the openings or spaces between adjacent portions or lands of the ceramic structure. Thus, it will be appreciated that the pores referred to in respect of the present invention have a nominal diameter of the order of magnitude of 0.1 mm. These are to be contrasted with pores which may be present in the catalyst support material itself, as is the case, for example, when a ceramic foam or particles prepared from a porous refractory oxide are employed.

The fixed arrangement comprises at least 750 pores per square centimetre. Preferably, the fixed arrangement comprises from about 1000 to about 15000 pores per square centimetre, more preferably from about 1250 to about 10000 pores per square centimetre, most preferably from greater than 3600 to about 10000 pores per square centimetre.

It will be clear that the fixed arrangement of the catalyst in the process of this invention is a porous structure. The fixed arrangement has a typical void fraction in the range of from 0.6 to 0.9.

The gaseous mixture of the hydrocarbon feedstock and the oxygen-containing gas are preferably contacted with the catalyst under adiabatic conditions. For the purposes of this specification, the term "adiabatic" is a reference to reaction conditions in which substantially all heat loss and radiation from the reaction zone is prevented, with the exception of heat leaving in the gaseous effluent stream of the reactor.

In a further aspect, the present invention relates to carbon monoxide or hydrogen whenever prepared by a process as hereinbefore described.

The mixture of carbon monoxide and hydrogen prepared by the process of this invention is particularly suitable for use in the synthesis of hydrocarbons, for example by

means of the Fischer-Tropsch synthesis, or the synthesis of oxygenates, for example methanol. Processes for the conversion of the mixture of carbon monoxide and hydrogen into such products are well known in the art.

The process of the present invention is further described by way of the following illustrative examples, of which Examples 1 and 2 are examples of the process of the present invention and Examples 3 and 4 are for comparison purposes only.

Example 1

A commercially available zirconia ceramic foam ( $\text{ZrO}_2$ , 1550 pores per square centimetre) was impregnated with rhodium using conventional impregnation techniques to give a final rhodium loading of 5% by weight.

A reactor was constructed comprising a transparent sapphire tube mounted concentrically within an outer transparent polycarbonate tube. The rhodium-containing catalyst prepared as hereinbefore described was loaded into the sapphire tube and retained in the tube to form a fixed arrangement having a tortuosity of between 1.5 and 2.5 and 1550 pores per square centimetre.

Methane and oxygen were thoroughly mixed before being introduced into the reactor to contact the fixed bed of catalyst. Methane and oxygen were present in amounts sufficient to give an oxygen-to-carbon ratio of 0.62. The gaseous feed mixture was supplied to the reactor at a gas hourly space velocity (GHSV) of 500,000  $\text{Nl/kg/hr}$  (Normal litres of gas per kilogramme of catalyst per hour) and at a pressure of 3.3 bara.

The operating temperature of the catalyst bed was measured by optical pyrometry. The composition of the gas mixture leaving the reactor was measured by gas chromatography. The conversion and the selectivity of the process to carbon monoxide and hydrogen (on the basis of methane converted) was determined. The operating

conditions of the reactor and the results of the experiment are summarised in the Table hereinbelow.

Example 2

The general procedure described in Example 1 above was followed to prepare and test a rhodium on zirconia ceramic foam catalyst having a rhodium loading of 0.5% by weight.

The operating conditions and results of the experiment are set out in the Table hereinbelow.

Example 3

By way of comparison, the general procedure described in Example 1 hereabove was followed to prepare and test a rhodium on zirconia ceramic foam catalyst having a rhodium loading of 5% by weight. A commercially available zirconia ceramic foam was selected as starting material, which resulted in the fixed catalyst arrangement having a tortuosity of from 1.5 to 2.5 and about 390 pores per square centimetre.

The operating conditions and results of the experiment are set out in the Table hereinbelow.

Example 4

By way of comparison, the general procedure described in Example 1 hereabove was followed to prepare and test a rhodium on zirconia ceramic foam catalyst having a rhodium loading of 1% by weight. A commercially available zirconia ceramic foam was selected as starting material, which resulted in the fixed catalyst arrangement having a tortuosity of from 1.5 to 2.5 and about 390 pores per square centimetre.

The operating conditions and results of the experiment are set out in the Table hereinbelow.

As can be seen from the Table, the process of the present invention, as exemplified in Examples 1 and 2, offers significant advantages in terms of process performance and operating costs. In particular, it can

be seen that the processes of Examples 1 and 2 gave a significantly higher level of methane conversion when compared with the comparative examples, Examples 3 and 4. Further, it is clear that the process of the present invention gives rise to a much improved hydrogen selectivity than the comparative processes. In addition, it can be seen that when the fixed arrangement of the present invention is employed, the amount of the catalytically active component employed in the catalyst, in this case rhodium, may be significantly reduced without adversely affecting the conversion and yields of the process. From the comparative examples, Examples 3 and 4, it can be seen that a reduction in the rhodium loading gives rise to a significant reduction in both the overall methane conversion and the selectivity of the process to hydrogen.

Table

| Example No.                                 | 1       | 2       | 3       | 4       |
|---------------------------------------------|---------|---------|---------|---------|
| <hr/>                                       |         |         |         |         |
| Fixed Catalyst Arrangement                  |         |         |         |         |
| Tortuosity                                  | 1.5-2.5 | 1.5-2.5 | 1.5-2.5 | 1.5-2.5 |
| Pores per square cm                         | 1550    | 1550    | 390     | 390     |
| Rhodium (% wt)                              | 5       | 0.5     | 5       | 1       |
| <hr/>                                       |         |         |         |         |
| Operating Conditions                        |         |         |         |         |
| Temperature (°C)                            | 1155    | 1070    | 1108    | 1150    |
| Pressure (bara)                             | 3.3     | 3.3     | 3.3     | 3.3     |
| GHSV (1000 Nl/kg/hr)                        | 500     | 500     | 400     | 400     |
| oxygen/carbon ratio                         | 0.62    | 0.62    | 0.62    | 0.62    |
| <hr/>                                       |         |         |         |         |
| CH <sub>4</sub> conversion (%)              | 96.7    | 97.5    | 87.3    | 83.6    |
| CO selectivity (%) <sup>1</sup>             | 90.9    | 91.8    | 92.2    | 92.2    |
| H <sub>2</sub> selectivity (%) <sup>2</sup> | 90.0    | 90.1    | 81.8    | 78.3    |

1 selectivity to CO based on CH<sub>4</sub> conversion

2 selectivity to H<sub>2</sub> based on CH<sub>4</sub> conversion



C L A I M S

1. A process for the catalytic partial oxidation of a hydrocarbon feedstock, which process comprises contacting a feed comprising a hydrocarbon feedstock and an oxygen-containing gas with a catalyst in a reaction zone, which catalyst is retained in the reaction zone in a fixed arrangement having a tortuosity of at least 1.1 and having at least 750 pores per square centimetre.
2. A process according to claim 1, characterised in that the hydrocarbon feedstock comprises methane, natural gas, associated gas or a source of light hydrocarbons.
3. A process according to either of claims 1 or 2, characterised in that the oxygen-containing gas is substantially pure oxygen.
4. A process according to any preceding claim, characterised in that the feed comprises the hydrocarbon feedstock and the oxygen-containing gas in amounts giving an oxygen-to-carbon ratio of from 0.3 to 0.8, preferably from 0.45 to 0.75, more preferably from 0.45 to 0.65.
5. A process according to any preceding claim, characterised in that the feed is contacted with the catalyst at a pressure in the range of up to 150 bar, preferably from 2 to 125 bar, more preferably from 2 to 100 bar.
6. A process according to any preceding claim, characterised in that the feed is contacted with the catalyst at a temperature in the range of from 950 to 1300 °C, preferably from 1000 to 1200 °C.
7. A process according to any preceding claim, characterised in that the feed is contacted with the catalyst at a gas hourly space velocity in the range of from 20,000 to 100,000,000 Nl/l/hr, preferably from

50,000 to 50,000,000 Nl/l/hr, more preferably from 500,000 to 30,000,000 Nl/l/hr.

8. A process according to any preceding claim, characterised in that the catalyst comprises a metal selected from rhodium, iridium or ruthenium.

9. A process according to any preceding claim, characterised in that the catalyst is retained in the fixed arrangement in the form of a fixed bed of catalyst particles or in the form of a ceramic foam, more preferably in the form of a ceramic foam.

10. A process according to any preceding claim, characterised in that the fixed arrangement of the catalyst has a tortuosity in the range of from 1.1 to about 10.0, preferably in the range of from 1.1 to about 5.0, more preferably in the range of from 1.3 to about 4.0.

11. A process according to any preceding claim, characterised in that the fixed arrangement of the catalyst has from about 1000 to about 15000 pores per square centimetre, preferably from about 1250 to about 10000 pores per square centimetre, more preferably from greater than 3600 to about 10000 pores per square centimetre.

12. A process according to any preceding claim, characterised in that the fixed arrangement has a void fraction in the range of about 0.6 to about 0.9.

13. A process according to any preceding claim, characterised in that the feed is contacted with the catalyst under substantially adiabatic conditions.

14. Carbon monoxide or hydrogen whenever prepared by a process according to any one of claims 1 to 13.

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