PCT

WORLD INTELLECTUAL PROPERTY ORGANIZATION International Bureau



INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶: C01B 13/02, B01D 53/22, 53/32

A1

(11) International Publication Number:

WO 98/55393

(43) International Publication Date:

10 December 1998 (10.12.98)

(21) International Application Number:

PCT/NO97/00170

(22) International Filing Date:

2 July 1997 (02.07.97)

(30) Priority Data:

972630

6 June 1997 (06.06.97)

NO

(71) Applicant (for all designated States except US): NORSK HYDRO ASA [NO/NO]; N-0240 Oslo (NO).

(72) Inventors; and

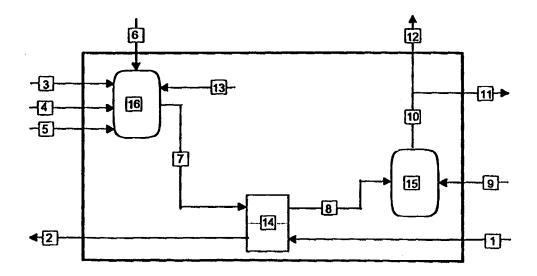
- (75) Inventors/Applicants (for US only): ÅSEN, Knut, Ingvar [NO/NO]; Tiurv. 22, N-3940 Heistad (NO). JULSRUD, Stein [NO/NO]; Gregorius Dagssonsg. 190, N-3713 Skien (NO).
- (74) Agent: SUNDNES, Arne; Norsk Hydro ASA, N-0240 Oslo (NO).

(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, HU, IL, 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, TJ, TM, TR, TT, UA, UG, US, UZ, VN, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG).

Published

With international search report.

(54) Title: METHOD FOR PERFORMING CATALYTIC OR NON-CATALYTIC PROCESSES, WITH ENRICHED OXYGEN AS ONE OF THE REACTANTS



(57) Abstract

The present invention relates to a method for performing a catalytic or non-catalytic process wherein oxygen is one of the reactants and being supplied from an oxygen containing gas mixture, and where oxygen is picked up from the permeate side of a mixed oxygen ion and electron conducting membrane by means of a sweep gas at elevated temperature. The thus formed oxygen containing sweep gas is applied as a reactant for a catalytic or non-catalytic oxidation process. The sweep gas can be preheated in a catalytic or non-catalytic reactor defined as a sweep gas preheater by burning of fuel. The sweep gas containing oxygen picked up from the membrane unit can be used in production of syngas for ammonia formation by being reacted with a hydrocarbon fuel in a catalytic reactor. The process gas used as sweep gas can also be NO-containing gas leaving the ammonia burner in a nitric acid process. The formed oxygen containing gas can then be used in the production of nitric acid.

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	rmenia FI Finland LT Lithuania		Lithuania	SK	Slovakia	
AT	Austria	FR	France	LU Luxembourg		SN	Senegal
AU	Australia	tralia GA Gabon LV Latvia		Latvia	SZ	Swaziland	
AZ			Monaco	TD	Chad		
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav	TM	Turkmenistan
BF	Burkina Faso	GR	Greece		Republic of Macedonia	TR	Turkey
BG	Bulgaria	HU	Hungary	ML	Mali	TT	Trinidad and Tobago
ВJ	Benin	IE	Ireland	MN	Mongolia	UA	Ukraine
BR	Brazil	IL	Israel	MR	Mauritania	UG	Uganda
BY	Belarus	IS	Iceland	MW	Malawi	US	United States of America
CA	Canada	IT	Italy	MX	Mexico	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NE	Niger	VN	Viet Nam
CG	Congo	KE	Kenya	NL	Netherlands	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NO	Norway	$\mathbf{z}\mathbf{w}$	Zimbabwe
CI	Côte d'Ivoire	KP	Democratic People's	NZ	New Zealand		
CM	Cameroon		Republic of Korea	PL	Poland		
CN	China	KR	Republic of Korea	PT	Portugal		
CU	Cuba	KZ	Kazakstan	RO	Romania		
CZ	Czech Republic	LC	Saint Lucia	RU	Russian Federation		
DE	Germany	LI	Liechtenstein	SD	Sudan		
DK	Denmark	LK	Sri Lanka	SE	Sweden		
EE	Estonia	LR	Liberia	\mathbf{SG}	Singapore		

1

METHOD FOR PERFORMING CATALYTIC OR NON–CATALYTIC PROCESSES, WITH ENRICHED OXYGEN AS ONE OF THE REACTANTS

The present invention relates to a method for performing catalytic or non-catalytic processes, wherein oxygen is one of the reactants, comprising supply of oxygen from an oxygen containing gas mixture.

In conventional catalytic or non-catalytic processes where oxygen is one of the reactants, the supply of oxygen is usually performed by compressing air which then is treated in an air liquefaction or cryogenic unit. Recovered oxygen is usually compressed and to some extent reheated. In such processes the energy requirement related to the supply of oxygen comprises a substantial part of the total energy consumption of the process. Thus air liquefaction and compression and reheating of the oxygen stream will require large amounts of energy. Further, the equipments required for these process steps are expensive.

In the field of gas separation, several methods are known. A rather new technique is application of membranes made from materials with both ionic and electronic conductivity. Such a membrane can be a mixed oxygen ion and electron conducting membrane, for instance capable of separating oxygen from oxygen containing gaseous mixtures at 400-1300°C. An oxygen partial pressure difference causes oxygen to be transported through the membrane by reduction of oxygen on the high oxygen partial pressure side (the feed side) and oxidation of the oxygen ions to oxygen gas on the low oxygen partial pressure side (the permeate side). In the bulk of the membrane, oxygen ions are transported by a diffusive process. Simultaneously the electrons flow from the permeate side back to the feed side of the membrane

2

Application of mixed conducting membranes in separation of gas mixtures is generally known from the patent application EP 0 658 367 A2. This application shows separation of oxygen from air by means of a mixed conducting membrane which is integrated with a gas turbine system. Pure oxygen is recovered from the permeate side of the membrane unit at near atmospheric pressure or below. The oxygen thus has to be cooled to below approx. 50°C and recompressed to required process pressure before being supplied to the oxidation reactor.

The main objective of the invention was to arrive at improved methods for performing catalytic and non-catalytic processes in which oxygen is one of the reactants.

Another objective was to supply oxygen to said processes in a way that implied reduced energy consumption and investment costs.

A further objective was to utilise existing process streams in obtaining a cheaper oxygen supply to the process.

It was also an objective to apply the basic concept for production of nitric acid.

One problem the inventors faced in their search for a cheaper oxygen supply, was that the basic processes should not be substantially changed. Further, it would be an advantage to be able to utilise existing hot process streams. The inventors therefore started to look for solutions that might meet both these requirements.

The fact that catalytic or non-catalytic processes where oxygen is one of the reactants usually comprise at least one process stream being at elevated temperature would meet one of the requirements for utilising mixed conductive membranes. However, in order to obtain a high separation efficiency and to avoid cooling and recompression of oxygen an applicable sweep gas for the permeate side of the membrane should be available. One requirement was that the

3

application of a sweep gas in said processes should not require supply of any additional reactant or gases and that it should be possible to make the sweep gas without installation of expensive new process equipment.

After having evaluated various ways of supplying oxygen to said catalytic or non-catalytic processes, the inventors decided to further investigate the possibility of applying mixed conductive membranes in spite of the teachings of the above EP-application. It was then found that in the processes in question there would be available various gas streams at elevated temperature making them applicable as sweep gas. Such sweep gas could be other reactants, steam, the products from a catalytic or non-catalytic chemical reaction, or recycled inert gases. Hot compressed air could be fed to a mixed conducting membrane which on the permeate side was exposed to a sweep gas of the above stated type and then picking up oxygen from the membrane unit. The sweep gas thus enriched with oxygen could then be transferred to a reactor for the final production of the desired products.

Application of a sweep gas in combination with a solid electrolyte membrane to lower the oxygen partial pressure to increase the degree of oxygen removal or oxygen recovery is known from the US Patent No. 5,035,726. In this patent a method for purifying crude argon is disclosed by selective permeation of oxygen through the membrane. Crude argon is compressed and heated and fed to a membrane unit to produce an O_2 depleted argon stream. In order to improve the efficiency of gas separation by the membrane, the permeate side of the membrane is swept by available nitrogen supplied from a cryogenic unit.

In the said argon purification process the use of a high temperature heat exchanger is required to produce a sweep gas having a temperature above at least 500°C and preferably above 600 to 700°C to not cool the membrane and thus reduce the rate of oxygen transport through the membrane. In rather small process plants, or in the case of low oxygen flows preheating of the sweep gas in

4

a high temperature heat exchanger could be economically feasible, but for recovery of several ton of oxygen per day the use of high temperature heat exchangers to preheat the sweep gas could be very expensive and probably not economical. By applying an available hot process stream as sweep gas the problems associated with the said heating of the sweep gas in the above US patent is avoided.

Application of pure oxygen as an oxidant in the ammonia oxidation process is generally known from US Patent No. 3,927,182. According to this patent oxygen is supplied from an oxygen plant, e.g. a cryogenic plant, and said oxygen is mixed with recycled tail gas and the thus formed oxygen containing tail gas is compressed in a recycle compressor and is then mixed with evaporated ammonia and the formed gas mixture is fed to an ammonia burner. The main disadvantage of this process is the high energy requirement and the high cost of oxygen.

The present invention will thus in its widest scope comprise a method for performing a catalytic or non-catalytic process wherein oxygen is one of the reactants being supplied from an oxygen containing gas mixture, and where oxygen is picked up from the permeate side of an oxygen ion and electron conducting membrane by means of a sweep gas at elevated temperature and where the thus formed sweep gas is a reactant for the catalytic or non-catalytic oxidation process downstream the membrane unit from here defined as the main reactor.

In said method the sweep gas can be preheated in a catalytic or non-catalytic reactor located upstream the membrane unit by burning of fuel from here defined as a sweep gas preheater.

A further embodiment of the invention comprises that the process gas used as sweep gas is the gas phase product or part of the gas phase product from the main reactor.

5

Another embodiment of the invention comprises that process gas containing oxygen not utilised in the main reactor is applied as oxidant in the sweep gas preheater.

The process gas used as sweep gas can be steam containing a gas mixture produced by mixing the steam with about stoichiometric amounts of fuel and air. The thus formed mixture can be fed to a sweep gas preheater to increase the temperature of the steam containing sweep gas to between 400 and 1300°C.

The composition and oxygen content of the sweep gas leaving the membrane unit can be regulated to give the desired temperature rise and gas composition of the product leaving the main reactor by regulation of the feed streams to the sweep gas preheater and the amount of oxygen transferred to the permeate side of said membrane unit.

The temperature in the sweep gas preheater can be controlled by addition of a coolant such as steam, water, inert gas or gas mixtures not taking part in the reaction in the main reactor.

The sweep gas having picked up oxygen from the membrane unit can be used in production of syngas for ammonia production by being reacted with a hydrocarbon fuel.

The process gas used as sweep gas is the NO-containing gas leaving the ammonia burner in a nitric acid process and the thus formed oxygen containing gas can be used in the production of nitric acid.

The main product is formed in the sweep gas preheater, and the gas leaving the sweep gas preheater is applied as sweep gas in the membrane unit. The sweep gas containing oxygen is cooled and separated in one product stream and one

stream containing oxygen is recycled to the sweep gas preheater as oxidant in said preheater.

The invention will be further explained and envisaged in the examples and corresponding figures.

Figure 1 shows a process comprising supply of oxygen using a mixed conducting membrane instead of conventional oxygen supply units.

Figure 2 shows a process comprising supply of oxygen according to the invention.

Figure 1 shows a chemical process, which can be catalytic or non-catalytic, or a combination of both, where oxygen is separated from an air stream by means of a mixed conducting membrane 16 at 400 to 1300°C and where the stream of oxygen is separated from air by means of a method as described in the above referred EP 0 658 367 A2. Thus obtained oxygen is then supplied in a conventional way to the chemical process.

Compressed air stream 1 is fed to a mixed conducting membrane unit 16. Pure oxygen is recovered from the permeate side of the membrane unit 16 at near atmospheric pressure or below. Oxygen depleted air leaves the unit 16 as stream 2. Oxygen stream 7 is then cooled to below approx. 40°C by first producing steam in heat recovery unit 17 and further by means of cooling water in heat exchanger 18. The oxygen stream 8 is then compressed to required process pressure in compressor 19 comprising one or more stages with intermediate cooling resulting in stream 9 which is fed to unit 20 together with a gas mixture 10 containing one or more reagents. Possibly also a reagent stream 11 is fed to unit 20. The resulting product from unit 20 is stream 12, which can be split such that part thereof is drawn off as streams 13 and 14. Gas mixture 10 could be made in unit 21 by mixing reactant 3, like e.g. natural gas, CO and H₂ containing synthesis gas,

ammonia, methanol etc. together with e.g. steam (stream 4) or inert gases (stream 5) and possibly also an oxygen containing gas stream 6 or a recycled process stream 15 which could be part of product stream 12. Unit 21 could be a catalytic or non-catalytic reactor or a static gas mixer.

The process as described in Figure 1 will however require undesired energy consumption in connection with the oxygen supply.

Figure 2 shows a chemical process, which can be a catalytic or non-catalytic process, according to the invention and where the oxygen reactant is delivered from a mixed conducting membrane 14 and where a sweep gas is fed to the permeate side of the membrane.

Compressed air stream 1 is fed to a mixed conducting membrane unit 14. To the permeate side of the membrane unit 14 there is supplied a sweep gas stream 7 which picks up oxygen from unit 14 and forms an oxygen containing gas stream 8 which is fed to a chemical reactor unit 15 to which is supplied a reactant stream 9, and giving the resulting product stream 10 which can be split such that part thereof is drawn off as streams 11 and 12. To unit 16 which is a catalytic or non-catalytic reactor a reactant stream 3 is supplied, like e.g. natural gas, a CO and H₂ containing gas, ammonia, methanol etc. together with e.g. steam (stream 4) or inert gases (stream 5) and possibly also an oxygen containing gas stream 6 or a recycled process stream 13 which could be part of product stream 10. The product formed in unit 16 is then used as sweep gas in the mixed conducting membrane unit 14. By applying the above described system for supplying oxygen, units 17, 18 and 19 of Figure 1 will not be necessary resulting in lower investment costs and reduced energy consumption.

Example 1

This example shows a chemical process performed in a system of units as described in Figure 2. Hot compressed air stream 1 at 1-30 bara and 500-1200°C is fed to the membrane unit to which a sweep gas stream 7 at 1-30 bara and 400-1300°C, produced in the reactor 16, is supplied to the permeate side of the membrane and recovering oxygen therefrom. The reactor 16 can be a catalytic or non-catalytic reactor. The sweep gas, which now contains the oxygen (stream 8), can then be fed to a catalytic or non-catalytic reactor 15 for reaction with reactant 9 producing the desired product of the total process. However, part of the said product can be recycled to the sweep gas producing reactor 16. Then this recycle gas must be cooled and recompressed and its oxygen content being controlled according to that required in the reactor 16.

An alternative configuration of the chemical process according to Figure 2 comprises that the desired product is produced in unit 16 and that the said product stream 7 (the sweep gas) is supplied to the permeate side of the membrane. The oxygen containing product stream 8 is cooled and separated in one product stream 11 and in one stream 12 containing oxygen. Thus unit 15 is not required. Stream 12 is then recycled by means of a compressesor or fan and the resulting oxygen containing stream 13 is fed to reactor 16 as an oxidant.

Example 2

This example shows application of the present invention on supply of oxygen to a syngas process where some nitrogen is acceptable in the syngas. The syngas can be used in ammonia production or in a once through or partly once through methanol or Fisher Tropsch process.

Steam is preheated by mixing steam with hot combusted fuel. The mixture consisting of mainly H_2O , N_2 and CO_2 is used as sweep gas in the membrane unit. Alternatively, air, fuel and steam are mixed and passed over a catalyst to burn

the fuel. The composition of the produced sweep gas could be as shown in the following table. The composition of the gas leaving the membrane unit (exit membrane) is also shown. The compositions are stated as volume%.

Component	Sweep gas %	Exit Membrane %
CO_2	2.0	1.85
N_2	16.2	15.00
O_2	0.50	7.87
Argon	0.20	0.19
H ₂ O	81.20	75.09
Temperature, °C	800	800
Pressure, bara	30	29.5

This example shows that the gas leaving the membrane unit has picked up enough oxygen (7.87 vol.%) for the production of syngas by burning this gas with a hydrocarbon feed. In e.g. production of ammonia the ratio between N_2 and O_2 in the sweep gas is regulated to give the desired ratio of H_2 and N_2 in the final syngas product.

Example 3

This example shows application of the present invention on supply of oxygen to a closed loop nitric acid process. The supply of oxygen in the present invention comprises application of the product gas leaving the ammonia burner as a sweep gas to pick up oxygen from the permeate side of a mixed conducting membrane. As in the above referred US Patent No. 3,927,182, the main part of the tail gas is recycled, but according to the present invention recycled tailgas contains at least 15% oxygen and is applied as an oxidant in the ammonia burner. Compared to conventional methods for supplying oxygen to a closed loop nitric acid process the present invention will reduce the energy requirement because recompression of the oxygen stream is no longer required.

10

By the present invention the inventors have arrived at a flexible process which compared to conventional processes promises lower investment and better utilisation of energy. This is obtained by application of a sweep gas for the permeate side of a mixed conductive membrane and where the sweep gas is made up of available hot process streams. Thus the application of a sweep gas in said processes does not require supply of additional gases or any additional supply of fuel or energy. The described application of mixed conductive membranes can in principle be used in the same way for several types of processes. Since the oxygen enriched sweep gas is leaving the mixed conducting membrane unit 16 of Figure 1 at elevated pressure and temperature it can be transferred directly to a reactor for the final production of the desired products without intermediate cooling, recompression and reheating, thus saving large compressors like unit 16 and heat exchangers like units 19 and 15 of Figure 1.

Claims

- 1. A method for performing a catalytic or non-catalytic process wherein oxygen is one of the reactants and being supplied from an oxygen containing gas mixture, and where oxygen is picked up from the permeate side of a mixed oxygen ion and electron conducting membrane by means of a sweep gas at elevated temperature and where the thus formed oxygen containing sweep gas is a reactant for a catalytic or non-catalytic oxidation process.
- 2. Method according to claim 1
 characterized in that
 the sweep gas is preheated in a catalytic or non-catalytic reactor
 defined as a sweep gas preheater by burning of fuel.
- 3. Method according to claim 1,
 c h a r a c t e r i z e d i n t h a t
 the process gas used as sweep gas is the gas phase product or part of
 the gas phase product from the reactor downstream the membrane unit.
- 4. Method according to claim 1,
 c h a r a c t e ri z e d i n t h a t
 the gas phase product or part of the gas phase product from the reactor
 downstream the membrane unit contains oxygen not utilised in the main
 reactor and that the said process gas is applied as oxidant in the sweep
 gas preheater.

5. Method according to claim 1,

characterized in that

the process gas used as sweep gas is a steam containing gas and where steam is heated by mixing with about stoichiometric amounts of fuel and air and feed the thus formed mixture to the sweep gas preheater to increase the temperature of the steam containing sweep gas to between 400 and 1300°C.

6. Method according to claim 1,

characterized in that

the composition and oxygen content of the sweep gas leaving the membrane unit is regulated to give the desired temperature rise and gas composition of the product stream from the catalytic or non-catalytic reactor by regulation of the feed streams to the sweep gas preheater and the amount of oxygen transferred to the permeate side of said membrane unit.

7. Method according to claim 1,

characterized in that

the temperature rise in the sweep gas preheater is controlled by addition of a coolant such as steam, water, inert gas or gas mixtures not taking part in the reaction in the catalytic reactor or non-catalytic reactor.

8. Method according to claim 1,

characterized in that

the sweep gas containing oxygen picked up from the membrane unit is used in production of syngas for ammonia formation by being reacted with a hydrocarbon fuel in a catalytic reactor.

- 9. Method according to claim 1, c h a r a c t e r i z e d i n t h at the process gas used as sweep gas is the NO-containing gas leaving the ammonia burner in a nitric acid process and that the then formed oxygen containing gas is used in the production of nitric acid.
- 10. Method according to claim 1, characterized in that the main product is formed in the sweep gas preheater and that the gas leaving the sweep gas preheater is applied as sweep gas in the membrane unit, and that the sweep gas containing oxygen is cooled and separated in one product stream and one stream containing oxygen and that the said stream containing oxygen is recycled to the sweep gas preheater as an oxidant.

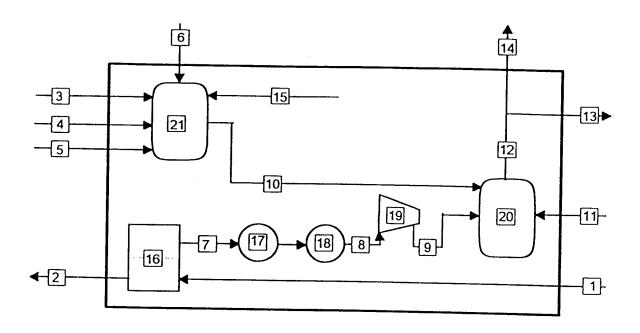


Fig. 1

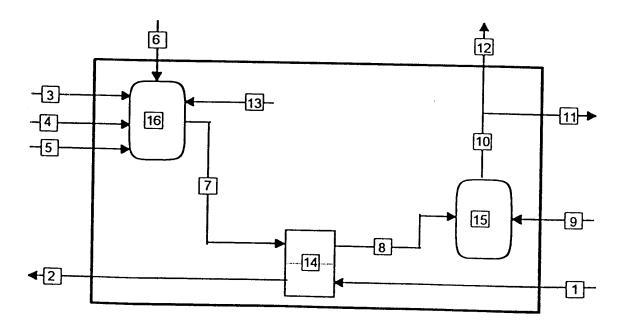


Fig. 2

INTERNATIONAL SEARCH REPORT

International application No.

1 - 10

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

document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive

PCT/NO 97/00170

A. CLASSIFICATION OF SUBJECT MATTER IPC6: C01B 13/02, B01D 53/22, B01D 53/32 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) IPC6: C01B, B01D 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) EPODOC, WPI C. DOCUMENTS CONSIDERED TO BE RELEVANT Citation of document, with indication, where appropriate, of the relevant passages Category* Relevant to claim No. Х EP 0743088 A2 (PRAXAIR TECHNOLOGY, INC.), 20 November 1996 (20.11.96) X US 5160713 A (TERRY J. MAZANEC ET AL), 1 3 November 1992 (03.11.92)

E,A	WO 9741060 A1 (ELTRON RESEARCH, INC.), 6 November 1997 (06.11.97), page 7, line 23 - line 30; page 11, line 6 - line 24; page 23, line 19 - line 29, figure 8	1-10
X Furth	er documents are listed in the continuation of Box C. X See patent family annex	

US 5245110 A (CHRISTIAN P VAN DIJK ET AL),

14 Sept 1993 (14.09.93)

document defining the general state of the art which is not considered

erlier document but published on or after the international filing date

document which may throw doubts on priority claim(s) or which is

document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other	step when the document is taken alone
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	"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
the priority date claimed	"&" document member of the same patent family
Date of the actual completion of the international search	Date of mailing of the international search report
	0 6 -04- 1998
1 April 1998	
Name and mailing address of the ISA	Authorized officer
Swedish Patent Office	
Box 5055, S-102 42 STOCKHOLM	Marianne Bratsberg
Facsimile No. +46 8 666 02 86	Telephone No. + 46 8 782 25 00

Special categories of cited documents:

to be of particular relevance

A

"E"

INTERNATIONAL SEARCH REPORT

International application No.
PCT/NO 97/00170

ategory*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No
A	EP 0658367 A2 (AIR PRODUCTS AND CHEMICALS, INC.), 21 June 1995 (21.06.95)	1

INTERNATIONAL SEARCH REPORT

Information on patent family members

02/03/98

International application No. PCT/NO 97/00170

Patent document cited in search report		Publication Patent family date member(s)		Publication date			
EP	0743088	A2	20/11/96	CA CN JP	2172301 1136467 8309142	A	19/11/96 27/11/96 26/11/96
US	5160713	A	03/11/92	CA JP WO	2093332 5508381 9205862	T	10/04/92 25/11/93 16/04/92
US	5245110	A	14/09/93	AU AU CA EP NO WO	662928 2660492 2119354 0604554 940996 9306041	A A A	21/09/95 27/04/93 01/04/93 06/07/94 18/05/94 01/04/93
0	9741060	A1	06/11/97	AU	6979196	A	19/11/97
 EP	0658367	A2	21/06/95	US US	5516359 5565017		14/05/96 15/10/96