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(21) International Application Number: PCT/NL91/00240 (22) International Filing Date: 28 November 1991 (28.11.91) (30) Priority data: 9002606 29 November 1990 (29.11.90) NL (71) Applicant (for all designated States except US): DSM N.V. [NL/NL]; Het Overloon 1, NL-6411 TE Heerlen (NL). (72) Inventors; and (75) Inventors/Applicants (for US only) : VAN DE MOESDIJK, Cornelis, Gerardus, Maria [NL/NL]; Op 't Veldje 33, NL-6176 BL Beek (NL). JANSSEN, Hendrik, Jan [NL/ NL]; Salviusstraat 7, NL-6141 LL Sittard (NL). SCHROIJEN, Josephus, Cornelus [NL/NL]; Achter de Heggen 6, NL-6373 XC Landgraaf (NL).		(74) Agent: ALFENAAR, Marinus; Octrooibureau DSM, P.O. Box 9, NL-6160 MA Geleen (NL). (81) Designated States: AT (European patent), AU, BB, BE (European patent), BF (OAPI patent), BG, BJ (OAPI patent), BR, CA, CF (OAPI patent), CG (OAPI patent), CH (European patent), CI (OAPI patent), CM (OAPI patent), CS, DE (European patent), DK (European pa- tent), ES (European patent), FI, FR (European patent), GA (OAPI patent), GB (European patent), GN (OAPI patent), GR (European patent), HU, IT (European pa- tent), JP, KP, KR, LK, LU (European patent), MC, MG, ML (OAPI patent), MR (OAPI patent), MW, NL (Euro- pean patent), NO, PL, RO, SD, SE (European patent), SN (OAPI patent), SU ⁺ , TD (OAPI patent), TG (OAPI patent), US. Published <i>With international search report.</i>
(54) Title: PROCESS FOR THE PREPARATION OF MELAMINE FROM UREA (57) Abstract The invention relates to a selective removal of impurities from the gas stream leaving a melamine synthesis reactor in the preparation of melamine from urea by means of a treatment with steam at elevated temperature in the presence of a catalyst which is shape-selective. For instance a molecular sieve is chosen.		

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PROCESS FOR THE PREPARATION OF MELAMINE FROM UREA

The invention relates to the steam treatment of the gas stream leaving the synthesis reaction in the preparation of melamine from urea or the thermal decomposition products thereof.

Such steam treatment is disclosed in US-A-3,386,999 and has the object of producing, after quenching of the treated gas stream, highly pure melamine which is free of cyanic acid and other melamine precursors.

In the single example of this patent publication the gas stream leaving the synthesis reactor is treated with steam for an unspecified period of time, the melamine obtained after a quenching and crystallisation is characterised by "dry and pure". Only 253.0 kg/hours melamine are obtained from 790 kg urea-feed per hour. Further NL-A-8105027, example 3 discloses that under the practical conditions of conventional melamine synthesis processes, temperature 360-400°C, and contact time 0.1-2.0 seconds, the conversion of cyanic acid is small if no catalyst is present and the uncatalysed process of US-A-3,386,999 is of no practical use. Therefore NL-A-8105027 proposes the use of a catalyst in the steam treatment to decompose gaseous urea, thermal decomposition products thereof and melamine containing gas streams into carbondioxide and ammonia. Preferred catalysts are chosen from the group of active carbon, oxides and phosphates of boron and aluminum. EP-A-51156 claims the use of iron and copper oxide to catalyze the steam treatment of gas stream containing melamine, cyanic acid etc. At temperatures in the range 100-150°C a virtually complete conversion of HCN, melamine and isocyanic acid into ammonia and carbon dioxide is realized.

These known catalytic treatments make only sense when all or virtually all of the above-mentioned compounds are to be removed from off-gas streams. In practice, however, there appears to be a demand for selective removal of HCN, isocyanic acid, cyanamide and similar compounds from process gas streams still containing melamine in higher concentrations without significant amounts of melamine being converted. This applies, for instance, to the product stream leaving the synthesis reactor in the melamine process. This stream contains in the fluidizing gas, ammonia, the carbon dioxide and ammonia released in the reaction, besides higher concentrations of melamine product and, to a much lesser degree, isocyanic acid, cyanamide and any hydrogen cyanide formed. When melamine is recovered from the process stream by cooling with water, an aqueous medium or a cold gas stream, these compounds give rise to the formation or precipitation of undesirable by-products such as ureido melamine, melam, melem, cyanuric acid, melamine cyanurate, ammelide, guanidine and ammeline. These contaminate either the sublimated melamine or the melamine-containing suspension, and as a consequence either melamine of too low purity is obtained, or a further purification step, for instance by recrystallization, as disclosed in US-A-3496177, becomes necessary. It may also prove necessary to prevent the build-up of impurities in the aqueous process stream obtained after separation, for instance by filtration, of the melamine, as this restricts recycling of this aqueous process stream in the process (see for instance US-A-44084046). For the several variants of the known melamine synthesis processes and the gas and liquid streams occurring in these, the general process literature is referred to as well as the many patent applications relating to this literature, for instance US-A-3321603, 3300493, 3513167, 3700672, 4348520 and GB-A-1309275. The above-mentioned restrictions raise the costs or necessitate the discharge of streams with a relatively high environment-burdening effect.

It is therefore the object of this invention to obtain a process for the treatment of gaseous product streams containing isocyanic acid, cyanamide and similar compounds in the preparation of melamine in which these compounds under the practical conditions of the process are converted into non-interfering by-products without melamine being decomposed to any significant degree.

10 The non-catalytic hydrolysis treatment being too slow, and the known catalyzed processes being non-selective, the inventors nevertheless have succeeded in developing a catalytic hydrolysis process that selectivity converses cyanic acid and other precursors of melamine into carbon
15 dioxide and ammonia without appreciably converting melamine, even though the latter may be present in a more than 1000-fold excess.

 The steam treatment of the gas stream in the presence of a catalyst according to the present invention is
20 characterized in that the catalyst is shape-selective.

 Shape-selective catalyst means in this respect, that the shape of the catalyst is such that its active sites are better accessible to molecules to be hydrolyzed than those that are not.

25 Shape selectivity of a catalyst is mainly determined by two factors, i.e. (1) the total number and location of catalytically active sites, and (2) the get-at-ability of the different reactants to these locations.

30 Complete specificity of the hydrolysis can in principle be realized if the catalytically active sites are located in such a way that they are only accessible for the by-products to be hydrolysed. This would be the case if the catalytic sites are located at the inner surface of the catalyst only,
35 and the melamine molecules are prevented of entering the inner surface of the catalyst.

 Generally the "inner" surface of the catalyst is defined as the active surface located in the micropores of the catalyst and the outer surface as the active surface

located in the mesopores of the catalyst.

The diameter of the micropores are smaller than 2.0 nm, preferably smaller than 1.2 nm and even more preferably smaller than 1.0 nm. Highest selectivity can be attained if the micropores are smaller than 0.8 nm. However pores with a diameter less than about 0.4 nm do not participate in the hydrolysis process.

10 The catalytic active surface should be located mainly in the micropores. Preferably less than 1/6 of the active sites is located in the mesopores, more preferably this fraction is smaller than 1/8, best results are obtained if this fraction is smaller than 1/10.

15 The above requirements with respect to pore diameter and active site location are indicative for catalysts that in principle are usefull as shape selective catalyst, however in the practice of industrial processes transport phenomena can play an important role, and selectivity may be affected by catalyst particle size, and the methods (for instance sintering or pressure) and materials (binders) used in shaping the catalyst into granulate, beads, spheres etc. for practical use in large scale processes. (Micro)pores with inert surfaces may be introduced and other pores (partly) obstructed, thereby influencing the get-at-ability of the active sites. Therefore catalysts having similar micropore diameter and distribution of active sites over inner and outer surface may show a different selectivity.

30 It is preferred that the process of the present invention is performed under such conditions that at least 60%, more preferably at least 80% and most preferably at least 90% of the by-product, but that at most 10%, more preferably at most 5% and most preferably at most 3% of the melamine in the gas stream is converted into carbondioxide and ammonia.

35 The catalyst for the hydrolysis reaction of melamine and by-products should contain acid cites. Oxidic materials are very usefull as hydrolysis catalyst.

Zeolites and molecular sieves are especially usefull as
shape selective catalyst material, as these show micropores
5 with well defined diameters.

The invention will now further be elucidated by the
example of zeolites, however the invention is not limited
thereto, and every material possessing acidic active groups
that is or can be brought into a shape selective form forms
10 part of the invention.

Zeolites are naturally occuring or synthetically
produced crystalline alkali metal alumino-silicates with a
three-dimensional interconnecting network of structure of
silica and alumina tetrahedra and form part of the state of
15 the art of catalysis. Examples are mordenite and erionite
and the A-type zeolite, which basically is in the sodium
form. Zeolites are commercially available and can be
obtained in many forms.

Zeolites are normally characterized by their nominal pore
20 diameter. This is the pore opening that can be calculated on
the basis of the crystallography of the zeolite. However
there are generally inhomogenities and other factors
disturbing the regular crystallographic structure. Therefor
the effective pore diameter should be preferred to
25 characterize the zeolite for practical reasons.

The effective pore diameter is the diameter of the largest
inert molecules that still enter the cavities in the zeolite
structure. This parameter is normally determined by
measuring the absorption of small model molecules as funtion
30 of their molecular diameter.

However this measurement being indicative for the
determination of the actual micropore diameter distribution
in the catalytic material, does not give the information
about the number of the active sites in the micropores vs.
35 those outside thereof.

The inventors have found that the use of
temperature programmed desorption (TPD) of trialkylamine
molecules is a representative method for determining the
available number of active sites and the distribution of the

active sites over inner and outer surface of inorganic catalyst. In TPD a catalyst sample is at room temperatures
5 equilibrated with the vapour of the trialkylamine. Consequently the temperature is increased gradually to about 500°C, and the weight loss as a function of the temperature is determined. The weight of adsorbed trialkylamine at 100°
10 is taken as indication of the number of acid sites that are accessible to the amine. If the TPD measurements are performed with ammonia (radius 0.3-0.4 nm) and triethylamine (radius about 0.8 nm), the number of acid sites accessible to the small by-product molecules and those accessible to melamine respectively are determined.

15 The catalyst's selectivity is further improved when the active groups on the outer surface of the catalyst are deactivated. This can be achieved by various known processes. Examples are inertization by ion exchange, loading with inert metals, for instance gold, or with inert
20 groups, for instance silicon compounds such as SiCl_4 , tetraalkyl orthosilicate, etc. The use of tetraethyl orthosilicate is preferred.

Depending on the technological conditions, such as reactor type, temperature and gas load, the catalyst can be
25 applied in the form of powder, tablets, rings, spheres, rods, etc. of varying diameters,

The temperature and pressure at which the conversion according to the invention is applied may vary within very wide limits, for instance 100-500°C and 0.1-100
30 bar, and are determined mainly by the temperature and pressure of the gas stream supplied, so that in most cases they will depend on the melamine synthesis process and the place in the process where the conversion is initiated. The lower limit of the temperature range is formed for practical
35 reasons by the desublimation point of the melamine in the gas stream. Very high temperatures, for instance more than 425°C, should generally be avoided to prevent depolymerization of melamine in cyanamide.

To achieve a high selectivity of the conversion the

amount of water is chosen so that the ratio between water and compounds to be converted is approximately
5 stoichiometric. Generally, a steam excess promotes complete removal of the substances to be converted, but its disadvantage is that melamine may also be hydrolysed. Preferably, therefore, the steam excess is limited to approximately a three-fold excess. When there is a steam
10 deficiency, of course not all impurities are converted. However, depending on the concentration of the compounds to be converted relative to the melamine concentration, if desired a higher or lower excess of water may be present in the gas stream or may be dosed. If only relatively very low
15 concentrations of substances to be converted are present, the excess may be relatively high because in that case in an absolute sense only little melamine can be converted with the residual steam. It should be noted that in most cases the stoichiometric amount of steam cannot exactly be
20 calculated because it is not possible to establish unequivocally the identity and concentration of all substances in the melamine process streams. By systematic research the average person skilled in the art will be able to determine the optimum amount of steam for a given
25 situation.

The catalyst can be applied in fixed bed or fluidized bed, preferably a fixed bed reactor is applied to prevent attrition.

The contact time of the gas stream with the
30 catalyst bed may vary within wide limits and is in part determined by the temperature of the treatment, the activity and amount of the catalyst. The average person skilled in the art, however, can simply establish this parameter for each individual case by systematic research. Generally, this
35 contact time will be between 0.25 and 1.5 sec.

The invention will now be elucidated with reference to the following examples, without being limited thereto.

Examples

- In a laboratory set-up the melamine synthesis was carried out at atmospheric pressure, subsequently the gas stream leaving the melamine synthesis reactor was subjected to the hydrolysis according to the invention and the melamine was recovered from the gas stream in a quench section as substantially depicted in Figure 1.
- 5 The synthesis section comprises two series-arranged fluidized bed reactors (1) and (2) of approximately 300 and 700 cm³, with a catalyst bed cross section of 7 and 12.5 cm², respectively. Both reactors contained 80 g of a customary SiO₂-Al₂O₃ catalyst.
- 10 As fluidization medium ammonia (3) was fed through the bottom plate of the first reactor. Urea (4) was dosed in the form of prills to the bed of the first reactor (dosing of a urea melt is possible in principle). The gas stream (5) leaving the second reactor is of a composition that is representative of commercial processes with a so-called wet melamine catch. The degree of conversion from urea to melamine is of the order of 90 to 95%.

Subsequently water (6) was dosed to the melamine-rich gas stream (5) leaving the second reactor and the steam-enriched gas stream (7) was passed over a fixed catalyst bed (8), with a height of 5-15 cm, depending on the type of catalyst, and a cross section of 10 cm², into the hydrolysis reactor (9).

The gas stream (10) leaving the hydrolysis reactor was subsequently cooled very rapidly in a quench system (11) using an amount of water that allowed all melamine present to dissolve. For proper contact the liquid and the gas were passed co-currently over a packed scrubber (13) (wet catch). The solution obtained (14) was recycled (17), via a buffer vessel with a continuous drain, to this quench system. The continuous drain (16) was set so that the input and output of the entire system per unit of time were in equilibrium.

Experiments were performed using different hydrolysis catalysts at different levels of the steam supply

to and the temperature of the hydrolysis reactor.

As soon as the system had stabilized, samples were
5 taken from the buffer vessel, which were analyzed for the
presence of melamine and by-products.

Of the catalysts the active surface area and its
distribution over inner and outer surface were determined by
means of temperature programmed desorption. The results are
10 presented in Table I.

The catalysts were applied as an extrudate, bead or
in pellet form.

The Zeolith 4A powder was mixed with 20% silica as inert
binder and then tabletted.

15

Example I

In Example I the experiments were carried out
according to the above process.

The samples were analyzed by liquid chromatography.
20 Depending on the component to be analyzed, detection took
place by the UV, fluorescence, conductivity or polarographic
method.

In the experiments 1-8 the ammonia flow rate was 72
25 Nl/hr, and the urea flow rate 35.5 g/hr. Water flow rate and
temperature were varied as indicated in the Table II. In
experiment 1-5 the UCAR 60343 catalyst was used. Zeolon 400
and Grace 522 were used in 6 and 7-8 respectively.

In experiment 9-10 the urea flow rate was increased
30 to 47.6 g/hr. As catalyst Grace 544 was used (table III).
Clearly no selectivity at all can be obtained with this
catalyst of which the active sites on the inner surface are
less than 50% of the total available number of active sites.

Table I

	Catalyst Manufacturer	Type	Form	TPD mol/g	TEA mol/g	Micropore size nm
5						
10	PG Corp. (Norton)	Zeolon 400	extrudate	2000	160	Mordinite nom 0.4-0.8 effective 0.35
15	Union Carbide	UCAR 60343	extrudate	2400	160	nom 0.36-0.52
20	Grace	522	bead	3100	290	Ca-alumino- silicate
25	Grace	544	bead	2600	1380	Nominal pore size 0.5 mm
30	Bayer	Zeolith 4A + 20% SiO ₂ *) tablet				nom 0.4
35	Norit	Active Carbon Supra A-8859	extrudate	250	3700	na **)

30

*) Zeolith 4A Supra A-8859 particles as obtained from the manufacturer were mixed with 20% (wt) highly pure and in active SiO₂ and compressed to tablets. It should be remarked that it is not exactly known which binder material are used by the other manufacturers. Generally these materials are alumina and or clay, of which the activity for the hydrolysis process is not to predict so that they may cause at this stage some deviations from the predicted behavior for the present shape selective catalyst.

**) It is apparent that TPD of ammonia and TEA (triethyl-amine) cannot be used to characterize active carbon. Further despite the high surface of active carbon the number of active sites as determined by NH₃-TPD is small.

45

Table II

5	Experiment	1	2	3	4	5
	water flow rate [g/hr]	1.14	2.28	0.57	1.14	2.28
	temperature [°C]	375	375	360	360	360
10	% melamine conversion	2	6	1	2	4
	reduction of impurity [%]	84	91	82	84	91
15	Experiment	6	7	8		
	water flow rate [g/hr]	1.15	1.14	1.70		
	temperature [°C]	390	370	370		
	% melamine conversion	3.6	5	8		
20	reduction of impurity [%]	67	90	94		

Table III

25	Experiment	9	10
	water flow rate [g/hr]	1.15	2.30
30	temperature [°C]	390	390
	% melamine conversion	9.3	13.9
	reduction of impurity [%]	17	15.7

35 Experiments 3, 4 and 5 clearly illustrate the effect of the excess of injected steam on the conversion. 0.57 g/hr steam approximately corresponds to the amount of water stoichiometrically needed at 95% urea conversion into melamine.

Example II

In processes involving cooling of the gas stream leaving the reactor with a cold ammonia/carbon dioxide mixture in the quench section and melamine recovery by sublimation, it is possible to operate the hydrolysis reactor under even more favourable conditions by recycling a portion of the NH_3/CO_2 gas mixture leaving the quench section to the hydrolysis reactor and diluting the gas stream before hydrolysis, so that a lower melamine partial pressure is obtained and, without additional means, the temperature of the hydrolysis reactor can be raised to the desired level. In Example II this situation is simulated by dosing mixed gas, $\text{NH}_3 : \text{CO}_2$ ratio = 2 : 1, upstream of the hydrolysis reactor in the feed circuit (250°C) upstream of the hydrolysis steam supply. Recycle gas of 250°C was added to the residue gas of 390°C from the synthesis reactor in such an amount that the temperature of the hydrolysis reactor was 355°C . (Ratio of reaction gas : recycle gas = approximately 3 : 1).

After stationary conditions had been reached, samples from the buffer vessel were analyzed. The results are given in Table IV.

Table IV

Experiment	12	13
NH_3 flow rate [Nl/hr]	72	72
urea flow rate [g/hr]	35.5	35.5
water flow rate [g/hr]	1.14	2.28
temperature [$^\circ\text{C}$]	355	355
% melamine conversion	1	3
reduction of impurity [%]	81	90
catalyst	UCAR 60343	

Example III

Using the conditions of experiment 11, with Zeolith
 5 4A as catalyst, in this example melamine separation from the
 process stream by desublimation was simulated, to which end
 a cooled rod (18) was introduced into the quench section
 (exp. 13).

The same experiment was conducted while the hydrolysis
 10 reactor was bypassed (blank experiment).

The solid samples obtained were analyzed for the most
 abundant impurities.

The results are presented in Table V.

15

Table V

Impurity	concentration, wt. %	
	blank	exp. 13
20 ammeline	0.33	< 0.02
ammelide	0.49	0.017
cyanuric acid	0.21	0.011
cyanourea	0.60	0.02
25 cyanomelamine [ppm]	160	25
melem [ppm]	460	10

Example IV

30 The Zeolon 400 catalyst was treated with
 tetraethylorthosilicate, through adsorption at between 300
 and 350°C of the tetraethylorthosilicate from a nitrogen
 stream, which is saturated at 85°C with the silicate, during
 a period of one hour.

35 Consequently the treated catalyst is calcined at 500°C in
 air during 2 hours.

Of the treated calcined catalyst the TPD of ammonia and
 triethylamine is determined. The result is given in Table
 VI.

TABLE VI

5	NH ₃	TEA
	10 ⁻⁶ mol/g	10 ⁻⁶ mol/g
	Zeolon 400	2000
10	(untreated)	160
	Zeolon 400	2120
	(treated)	89

15

The number of active sites accessible for TEA has decreased by nearly factor two, whereas the total number of active sites has remained nearly constant.

Experiment 6 was repeated with the treated Zeolon 400.

20 The melamine conversion was about 2.5% and 80% of the impurities was converted, showing the positive effect of deactivation of the outer surface on the selectivity of the catalyst.

C L A I M S

- 5
1. Process for the treatment with steam in the presence of a catalyst of a gas stream in the preparation of melamine from urea or from thermal decomposition products thereof, characterized in that the catalyst is
10 shape selective.
 2. Process according to claim 1, characterized in that the catalyst is chosen from the group of natural and synthetic zeolites and molecular sieves.
 3. Process according to claim 1, characterized in that the
15 effective diameter of the micropores in the shape-selective catalyst is between 0.3 and 2.0 nm.
 4. Process according to claim 1, characterized in that the active sites of the catalyst that are located within the micropores is at least 50% of total active sites.
 - 20 5. Process according to claim 4, characterized in that at least 80% of the active sites is located within the micropores.
 6. Process according to claim 5, characterized in that at least 90% of the active sites is located within the
25 micropores.
 7. Process according to claims 3 and 4.
 8. Process according to any one of claims 1-7, characterized in that the active sites on the outer surface of the catalyst are deactivated.
 - 30 9. Process according to claim 8, characterized in that the catalyst has been treated with triethyl orthosilicate.
 10. Process according to any one of claims 1-9, characterized in that the temperature at which the steam treatment takes place is between 100 and 500°C.
 - 35 11. Process according to claim 10, characterized in that the temperature is above approximately 300°C.

12. Process according to any one of claims 1-9,
5 characterized in that the amount of steam used for the treatment is 1 to 3 times the amount of steam needed for complete conversion of the by-products.
13. Process as substantially described in the specification,
the examples and the figure.

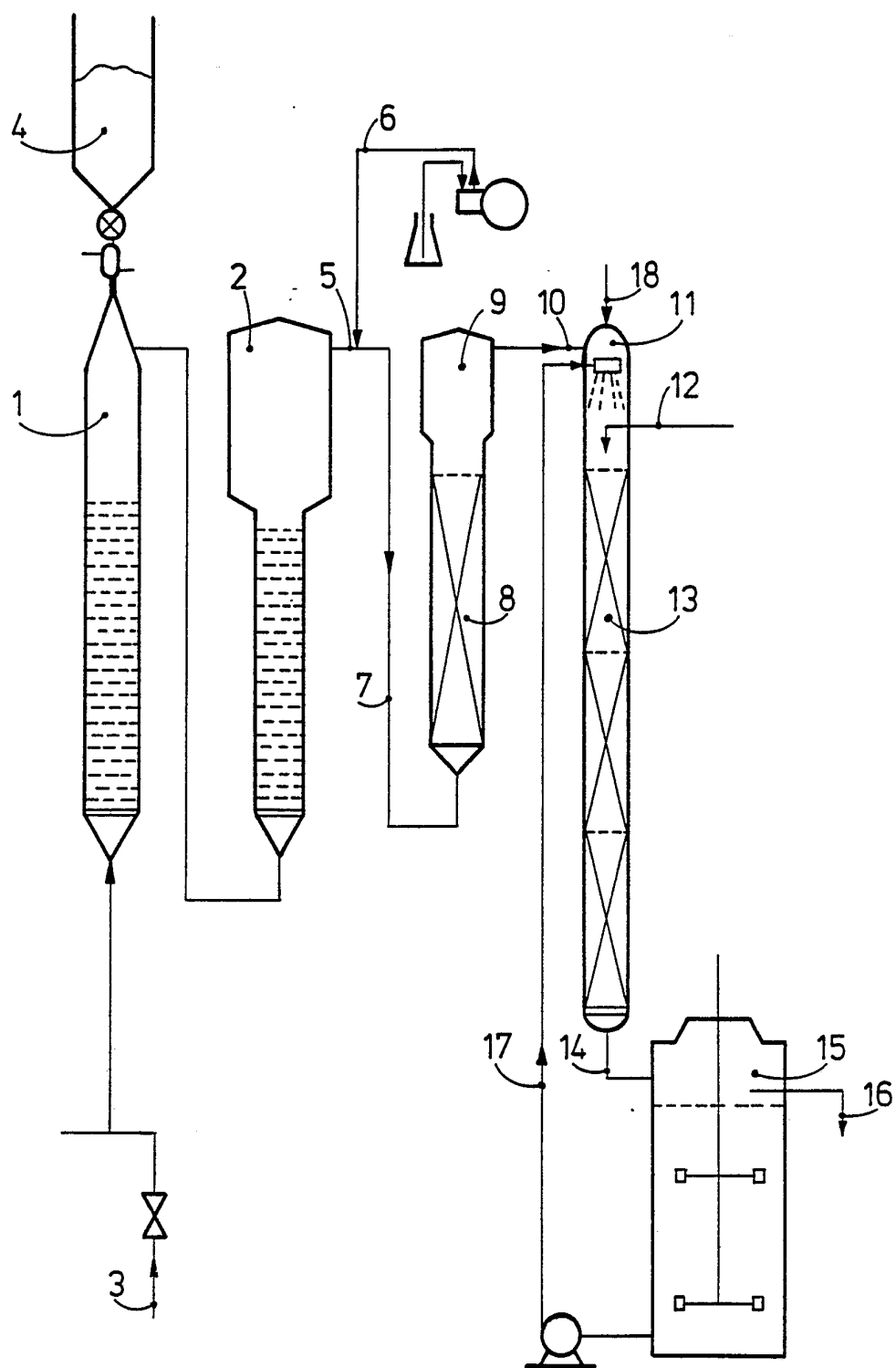


Fig. 1

INTERNATIONAL SEARCH REPORT

PCT/NL 91/00240

International Application No

I. CLASSIFICATION OF SUBJECT MATTER (if several classification symbols apply, indicate all) ⁶		
According to International Patent Classification (IPC) or to both National Classification and IPC Int.Cl. 5 C07D251/60; C07D251/62		
II. FIELDS SEARCHED		
Minimum Documentation Searched ⁷		
Classification System	Classification Symbols	
Int.Cl. 5	C07D ; C07C	
Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸		
III. DOCUMENTS CONSIDERED TO BE RELEVANT⁹		
Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³
A	NL,A,8 105 027 (STAMICARBON B.V.) 1 June 1983 cited in the application see the whole document; especially page 8-9, example 6, table 5, line 6 - line 8 ---	1-3, 10-12
A	US,A,3 386 999 (M. MANES) 4 June 1968 cited in the application see column 5 - column 6; claims see column 4, line 71 - column 5, line 60; example ---	1,10-12
A	EP,A,0 051 156 (BASF AKTIENGESELLSCHAFT) 12 May 1982 cited in the application see the whole document ---	1,10
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¹⁰ 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 "L" 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 "T" 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 "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		
IV. CERTIFICATION		
Date of the Actual Completion of the International Search	Date of Mailing of this International Search Report	
21 FEBRUARY 1992	11.8.92	
International Searching Authority	Signature of Authorized Officer	
EUROPEAN PATENT OFFICE	FINK D.G. <i>[Signature]</i>	

III. DOCUMENTS CONSIDERED TO BE RELEVANT (CONTINUED FROM THE SECOND SHEET)		
Category °	Citation of Document, with indication, where appropriate, of the relevant passages	Relevant to Claim No.
A	US,A,3 414 571 (H.W. HAINES) 3 December 1968 see the whole document; especially column 2, line 52 - line 61 ---	1,10

**ANNEX TO THE INTERNATIONAL SEARCH REPORT
ON INTERNATIONAL PATENT APPLICATION NO.**

NL 9100240
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