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(54) Title: LIQUID DETERGENT COMPOSITION (57) Abstract Liquid detergent compositions for softening and washing fabrics comprising an aqueous base, a detergent active material such as a mixture of anionic and nonionic materials, a detergency builder, such as sodium tripolyphosphate, and a special fabric softening clay which swells in water but not in sodium tripolyphosphate solution. The use of such clays enable the product to be prepared without an undesirably high viscosity.		

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LIQUID DETERGENT COMPOSITION

5 This invention relates to a liquid detergent composition, in particular to a liquid detergent composition for washing fabrics and imparting a softness thereto.

10 British Patent Specification No GB 2 132 629-A describes a fabric softening heavy duty liquid detergent which contains finely divided swelling bentonite as a fabric softening material. A number of suitable bentonite materials is suggested for use, including Wyoming bentonite.

15 We have found that these recommended bentonites when incorporated in liquid detergent compositions, especially those such compositions which exist as structured liquids, significantly increase the viscosity of the product.

20 Viscosity is an important property. Too low a viscosity can result in long term product instability when the product contains undissolved material in suspension, whereas too high a viscosity makes product processing and use by the consumer difficult.

25

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We have surprisingly found a class of fabric softening clay materials which do not significantly increase product viscosity in such products but yet still provide a softness benefit on fabrics treated with the product.

Thus, according to the present invention, there is provided a liquid detergent composition for washing fabrics and imparting softness thereto, the composition comprising

- (i) an aqueous base;
- (ii) a detergent active material or a mixture thereof;
- (iii) at least 5% by weight of a detergent builder; and
- (iv) a fabric softening clay material having a swellability in water (determined as herein described) of more than 36% and a swellability in an 8% sodium tripolyphosphate solution of less than 25%.

The fabric softening clay materials which are useful in the compositions are characterised by their swelling behaviour, which is quantified by the following test.

Two dispersions are prepared at room temperature containing respectively:

- A - 475g water and 25g of clay material;
- B - 435g of water, 40g sodium tripolyphosphate and 25g of clay material (the sodium tripolyphosphate is

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completely dissolved in the water before the addition of the clay).

The dispersions are stirred for 5 minutes with a magnetic stirrer and then placed in a 1000 ml measuring cylinder. The dispersions are then left to stand, undisturbed for two weeks. After this time the dispersions are examined. Generally some separation will have occurred. A lower layer of dispersion or gel containing the clay will be visibly distinguishable from a relatively clear upper layer. The height of the lower layer (h) and the overall height of the total liquid (H) are determined and percentage swellability (S) is calculated using the expression

15

$$S = \frac{h}{H} \times 100$$

We have found that clay materials having a swellability (S) of more than 36%, preferably more than about 75% in dispersion A, and less than 25%, preferably less than about 20% in dispersion B are useful in the present composition.

25

The following Table identifies a number of useful clay materials in this respect.

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TABLE

		<u>S (g)</u>		
	<u>TRADE NAME</u>	<u>CLAY TYPE</u>	<u>DISPERSION A</u>	<u>DISPERSION B</u>
5	CLARSOL STF	) Kaolinite	9	14
	MKIC	)	9	15
	CLARSOL ATC	Attapulgate	95	63
10	BENTONE EW	Hectorite	100	65
	CLARSOL W100	Na Bentonite	100	98
	CLARSOL KC1	)	36	86
15	MDO 77/84	) Ca Bentonite	9	73
	LAUNDROSIL DG	)	1	75
	CP 103	)	10	95
	MARMORA	)	15	10
20	CLARSOL KC1	) Ca Bentonite	100	98
	MDO 77/84	) accurately ¹	98	48
	LAUNDROSIL DG AC	) activated with	100	51
	CP103	) sodium carbon-	100	100
	MARMORA	) ate	100	15
25	CLARSOL KC2	) Ca Bentonite	100	68
	MDO 81/84	) commercially ¹	100	36
	LAUNDROSIL DG AC	) activated with	100	25
	DOKUM KARAKAYA	) sodium carbonate	95	16
30	BENTONITE DC	) acid activated	10	10
		) calcium bentonite		
	STEETLEY NO 1	) white	68	14
35	STEETLEY NO 2	) bentonite	75	20

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1 - commercial activation with sodium carbonate usually results in the presence of excess sodium carbonate in the treated material. Accurate activation is carried out with the objective of leaving no excess sodium carbonate.

5

Of the clay materials listed in this Table, only DOKUM KARAKAYA, accurately activated MARMORA and STEETLEY NOS 1 and 2 meet the requirements of the present invention.

10

We have found it indeed surprising that some clay materials which have a low swellability in sodium tripolyphosphate solutions will swell considerably in water and will, when incorporated in products as described herein, provide fabrics washed therewith with a softening benefit.

It would appear that the clay materials which are useful in the present invention are some lamella smectite clays containing exchangeable sodium and calcium cations. Clay materials which are free of these ions, such as acid activated clays, do not swell sufficiently in water and do not provide a fabric softening benefit. It will be appreciated that in a practical liquid detergent product, the exchangeable cationics of the clay will exchange with those of the electrolyte system of the product prior to its actual use in the wash process. The softening benefit will therefore relate to this exchanged form.

30 The level of fabric softening clay material in the product is preferably at least 1% by weight, but not more than 10% by weight. A most preferred level is from 3% to 7% by weight.

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The detergent compositions of the present invention necessarily contain one or more detergent active materials.

5 The detergent compounds may be selected from anionic, nonionic, zwitterionic and amphoteric synthetic detergent active materials. Many suitable detergent compounds are commercially available and are fully described in the literature, for example in "Surface Active Agents and
10 Detergents", Volumes I and II, by Schwartz, Perry and Berch.

The preferred detergent compounds which can be used are synthetic anionic and nonionic compounds. The former
15 are usually water-soluble alkali metal salts of organic sulphates and sulphonates having alkyl radicals containing from about 8 to about 22 carbon atoms, the term alkyl being used to include the alkyl portion of higher acyl radicals. Examples of suitable synthetic anionic
20 detergent compounds are sodium and potassium alkyl sulphates, especially those obtained by sulphating higher (C_8-C_{18}) alcohols produced for example from tallow or coconut oil, sodium and potassium alkyl (C_9-C_{20}) benzene sulphonates, particularly sodium linear secondary alkyl
25 ($C_{10}-C_{15}$) benzene sulphonates; sodium alkyl glyceryl ether sulphates, especially those ethers of the higher alcohols derived from tallow or coconut oil and synthetic alcohols derived from petroleum; sodium coconut oil fatty monoglyceride sulphates and sulphonates; sodium and
30 potassium salts of sulphuric acid esters of higher (C_8-C_{18}) fatty alcohol-alkylene oxide, particularly ethylene oxide, reaction products; the reaction products of fatty acids such as coconut fatty acids esterified with isethionic acid and neutralised with sodium hydroxide;
35 sodium and potassium salts of fatty acid amides of methyl taurine; alkane monosulphonates such as those derived by

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reacting alpha-olefins (C_8-C_{20}) with sodium bisulphite and those derived from reacting paraffins with SO_2 and Cl_2 and then hydrolysing with a base to produce a random sulphonate; and olefin sulphonates, which term is used to describe the material made by reacting olefins, particularly $C_{10}-C_{20}$ alpha-olefins, with SO_3 and then neutralising and hydrolysing the reaction product. The preferred anionic detergent compounds are sodium ($C_{11}-C_{15}$) alkyl benzene sulphonates and sodium ($C_{16}-C_{18}$) alkyl sulphates.

Suitable nonionic detergent compounds which may be used include in particular the reaction products of compounds having a hydrophobic group and a reactive hydrogen atom, for example aliphatic alcohols, acids, amides or alkyl phenols with alkylene oxides, especially ethylene oxide either alone or with propylene oxide. Specific nonionic detergent compounds are alkyl (C_6-C_{22}) phenols-ethylene oxide condensates, generally 5 to 25 EO, ie 5 to 25 units of ethylene oxide per molecule, the condensation products of aliphatic (C_8-C_{18}) primary or secondary linear or branched alcohols with ethylene oxide, generally 5 to 40 EO, and products made by condensation of ethylene oxide with the reaction products of propylene oxide and ethylenediamine. Other so-called nonionic detergent compounds include long chain tertiary amine oxides, long chain tertiary phosphine oxides and dialkyl sulfoxides.

Amounts of amphoteric or zwitterionic detergent compounds can also be used in the compositions of the invention but this is not normally desired due to their relatively high cost. If any amphoteric or zwitterionic detergent compounds are used it is generally in small amounts in compositions based on the much more commonly

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used synthetic anionic and/or nonionic detergent compounds.

Mixtures of detergent active materials may be used.

- 5 In particular, we prefer a mixture of an anionic detergent active, a nonionic detergent active and soap, particularly when the product is in the form of a structured liquid.

- 10 Where the detergent active material is soap, this is preferably selected from alkali metal salts of fatty acids having 12 to 18 carbon atoms. Typical such fatty acids are oleic acid, ricinoleic acid, and fatty acids derived from castor oil, rapeseed oil, groundnut oil, coconut oil, palmkernel oil or mixtures thereof. The sodium or
15 potassium salts of these acids can be used.

- The level of detergent active material in the product is preferably at least 2% by weight, but not more than 45% by weight, most preferably from 6% to 15% by weight.
20

- The products according to the invention necessarily contain a detergency builder material to reduce the level of free calcium ions in the wash liquor and thereby improve detergency. This material may be selected from
25 precipitating detergency builder materials such as alkali metal carbonates and ortho-phosphates, ion-exchange builder materials such as alkali metal aluminosilicates and sequestering builder materials such as alkali metal tripolyphosphates, citrates and nitrilotriacetates.
30 Particularly preferred is sodium tripolyphosphate for reasons of product structure and building efficiency. At least 5% by weight of the detergency builder material is required to provide a noticeable effect upon detergency.

- 35 It is particularly preferred that the product be in the form of a structured liquid, that is a liquid which

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contains a detergent in the lamella phase, which provides the product with rheological properties such that any undissolved material is held in stable homogeneous suspension. This enables the product to contain

5 relatively high levels of detergency builder. The lamella phase is obtained by a critical choice of detergent active materials. European patent specification No EP-A-38101 (UNILEVER) describes such a product which contains sodium tripolyphosphate and a

10 detergent active mixture containing an anionic detergent active material, a nonionic detergent active material and a soap, the level of the sodium tripolyphosphate being more than would be soluble in the product, with the result that some of the tripolyphosphate remains undissolved but

15 stably suspended in the product.

It is a preferred feature of the present invention that the level of detergency builder material in the product is more than would dissolve at 20°C. In the case

20 of sodium tripolyphosphate, a preferred level is from 22 to 35% by weight, based on the weight of the product.

The liquid detergent composition of the invention may further contain any of the adjuncts normally used in

25 fabric washing detergent compositions, eg sequestering agents such as ethylenediamine tetraacetate; buffering agents such as alkali silicates; soil suspending and anti-redepositon agents such as sodium carboxymethyl cellulose and polyvinylpyrrolidone; fluorescent agents;

30 perfumes; germicides; and colourants.

Further, the addition of lather depressors such as silicones, and enzymes, particularly proteolytic and amylolytic enzymes; and peroxygen bleaches, such as sodium

35 perborate and potassium dichlorocyanurate, including bleach activators, such as N,N,N',N',- tetraacetyl

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ethylene diamine, may be useful to formulate a complete heavy duty detergent composition suitable for use in washing machines.

5 Also particularly beneficial are agents for improving the thermal stability of the product, such as sodium toluene sulphonate, xylene sulphonate or cumene sulphonate, at levels of up to 1% by weight, such as from 0.4% to 0.5%.

10

 The products of the present invention may be prepared by a variety of methods. However, we have found that benefits arise from mixing ingredients in a particular order. Thus, it is preferable to add a portion of the
15 detergency builder to water, before adding the clay and the detergent active material. In this way products having uniform rheological properties from batch to batch can be obtained. In particular, a preferred method is to add the necessary quantity of water at an elevated
20 temperature of say 40°C-80°C to a mixing vessel provided with a stirrer. An amount of between one part in twenty and one part in four of the detergency builder is then added, with stirring. Where the detergency builder is water-soluble, this amount will dissolve in the water and
25 prevent the clay material from swelling but will not be sufficient to impair the stability of the surfactant. The clay material is then added and dispersed with stirring. Anionic and nonionic detergents, including soap where this is present, are then added. The remaining part of the
30 detergency builder is then added while maintaining this elevated temperature with stirring until a homogeneous mass is obtained.

 Finally, the mixture is cooled under constant
35 agitation and water is added, if necessary, to compensate

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evaporation loss. Thereafter perfume may be added when the product is at substantially ambient temperature.

When, alternatively, the clay is added to the water before any detergency builder, the clay swells producing a composition which may have a viscosity which is higher than desired. If all the detergency builder is added before the clay a product may result which separates on standing. If both the detergent active material and the detergency builder are added before the clay, the product may already have a high viscosity so that the powdered clay cannot easily be added without at the same time introducing air into the product resulting in a product having a density lower than may be desired.

The compositions of the invention should have a viscosity of less than 3000, preferably less than 1500 cPs measured at 20°C and at a shear rate of 21 sec⁻¹. Most preferably the viscosity is between 650 and 850 cPs. Viscosities below 650 cPs can result in a loss of product stability.

The invention will now be illustrated by the following examples.

EXAMPLE 1

A liquid detergent composition was prepared according to the following formulation:

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	<u>Ingredient</u>	<u>% (by weight)</u>
	Sodium C ₁₂ -alkyl benzene sulphonate	6.5
5	Soap	1.0
	Alcohol ethoxylate 7EO	2.5
	Clay	5.0
	Sodium carboxymethyl cellulose (SCMC)	0.1
	Sodium tripolyphosphate (STP)	22.8
10	Sodium silicate	1.0
	Fluorescent agent	0.1
	Glycerol	4.85
	Borax	3.1
	Silicone	0.16
15	Perfume	0.29
	Proteolytic enzyme	0.80
	Water	balance

20

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This composition was made by the following method:

The water is heated to 60°C and maintained at that temperature. 2% STP is added, followed by the clay, SCMC, fluorescer, sodium hydroxide, silicate, glycerol, borax, fatty acid and sulphonic acid (which with the sodium hydroxide generate the soap and the anionic detergent active respectively) and nonionic active while stirring is continued. After 10 minutes agitation the remaining 20.8% STP was added and the mixture was then cooled with further stirring. When cool, the silicone, perfume and enzymes were added.

A number of such compositions were prepared containing different clay materials. In each case the product viscosity was measured at 20°C and 21 sec⁻¹. Each composition was used to wash cotton test cloths using the following wash method:

Cotton terry towelling test cloths which have been preharshened by 10 washes in a commercially available fabric washing powder product SKIP (ex Lever, France), are washed in the test product for 20 minutes at 40°C using tap water with a hardness of 48°FH. A laboratory scale apparatus having a capacity of 1 litre is used, and three test cloths of size 15cm x 15cm are washed together. After washing, the cloths are rinsed twice in tap water, wrung out and line dried for 24 hours.

After drying, the cotton test cloths were assessed for softness by a panel of 12 experts, each clay being compared against White bentonite, ex Steetley. The results were as follows:

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	<u>Clay material</u> (Before incorporation)	<u>supplier</u>	<u>softening</u>	<u>viscosity</u> (cPs)
5	BENTONITE DC	SUD-CHEMIE	)	870
	CLARSOL ATC	CECA	)	1700
	MKIC	KAOLINS DU	) significantly	850
		MORBIHAN	) poor	
	LAUNDROSIL DG AC	SUD-CHEMIE	)	1250
10	CLARSOL STF	CECA	)	750
	BENTONE EW	NATIONAL LEAD)		1800
	MDO 81/84	ECC	)	1300
	LAUNDROSIL DG	SUD-CHEMIE	)	1600
15	CLARSOL KC 1	CECA	)	1500
	CLARSOL KC 2	CECA	) no	1200
	CLARSOL W 100	CECA	) significant	1500
	CP 103	LAPORTE	) difference	1800
	MDO 77/84	ECC	)	1350
20	WHITE BENTONITE	STEETLEY	)	850
		(contol)	)	
	CALCIUM	MARMORA	)	750
	BENTONITE			

25

It will be seen from these results that a number of clay materials provide a softening benefit which is not significantly different to WHITE BENTONITE. However, in most cases the viscosity of the products is higher than desirable. Two clay materials, MKIC and CLARSOL STF provide product viscosities equal or less than WHITE BENTONITE. However, in both cases the softening benefit is less preferred.

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The only clay material which is comparable to WHITE BENTONITE both in terms of softening and viscosity is the bentonite from MARMORA.

5

EXAMPLE 2

A mixture of fabric test cloths were washed with detergent compositions as set out below at 40°C in water having a hardness of 30°FH (3×10^{-3} molar free calcium ions). Some test cloths consisted of new terry towelling, some consisted of new acrylic fibres and a third group consisted of terry towelling pieces which had been pre-harshened by washing 30 times in a commercially available softener-free powder product SKIP (ex Lever, France) in 45°FH water. After the mixed test cloths were washed 10 times, rinsed and dried in a conventional manner (without the use of a post-wash fabric softening agent) they were divided into fabric types and assessed for softness by a panel of expert assessors (lower softness scores indicate better softness).

The formulations tested and the results obtained were as follows:

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EXAMPLE NO:		2WB	2W	2P	2L
Ingredient (%)					
	Anionic detergent active	6.5	6.5	5.5	14.1
5	Nonionic detergent active	2.5	2.5	2.0	12.8
	Soap	1.0	1.0	-	15.8
	Dialkylmethethylamine	-	-	3.8	-
	Coconut trimethylammonium chloride	-	-	1.2	-
10	Sodium tripolyphosphate	22.8	22.8		-
	Sodium perborate	-	-	11.7	-
	Sodium silicate	1.0	1.0	5.7	-
	Sodium sulphate	-	-	22.6	-
	Sodium carbonate	-	-	1.4	-
15	Sodium carboxymethyl cellulose	0.1	0.1	0.8	-
	Clay	5.0	-	3.4	-
	Water and miscellaneous	-----balance-----			
20	Product form	Liquid	Liquid	Powder	Liquid
	Product dosage (per 20 l)g	215	215	234	155
25	<u>SOFTENING SCORES</u>				
	- New terry towelling	6	19	18	14
	- Preharshened terry towelling	5	19	14	13
30	- New acrylic	7	22	9	14

Formulation 2WB is substantially identical to Example 1, utilises STEETLEY NO 1 White Bentonite as the clay component. Formulation 2W is identical except that the clay component has been omitted. Formulations 2P and 2L are intended to represent commercially available fabric washing products used at recommended dosages for those products.

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As will be seen from the softening results,
formulation 2WB is preferred to all other formulations
tested.

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CLAIMS

1. A liquid detergent composition for washing fabrics
and imparting softness thereto, the composition
5 comprising:
- (i) an aqueous base;
 - (ii) a detergent active material or a mixture
10 thereof;
 - (iii) at least 5% by weight of a detergency builder;
and
 - 15 (iv) a fabric softening clay material having a
swellability in water (determined as herein
described) of more than 36% and a swellability
in an 8% sodium tripolyphosphate solution of
less than 25%..
20
2. A liquid detergent according to claim 1, comprising
from 6% to 15% by weight detergent active material.
3. A liquid detergent according to claim 1, wherein the
25 detergent active material is selected from anionic,
nonionic, zwitterionic and amphoteric detergent active
materials and mixtures thereof.
4. A liquid detergent according to claim 1, comprising
30 from 22 to 35% by weight detergency builder.
5. A liquid detergent according to claim 1, wherein the
detergency builder is selected from precipitating
detergency builder materials, ion-exchange builder
35 materials and sequestering builder materials.

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6. A liquid detergent according to claim 5, wherein the detergency builder comprises sodium tripolyphosphate.

7. A liquid detergent according to claim 1, comprising 1%
5 to 10% by weight of the fabric softening clay material.

8. A liquid detergent according to claim 1, wherein the fabric softening clay material is a lamella smectite clay.

10 9. A process for preparing a liquid detergent composition comprising an aqueous base, one or more detergent active materials, at least 5% by weight of a detergency builder and a fabric softening clay material, said process comprising the steps of:

15

(i) adding at least a portion of the detergency builder to water;

20

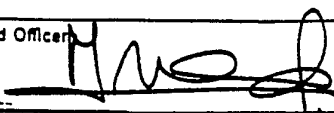
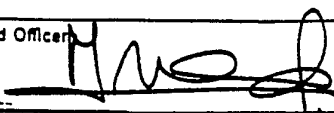
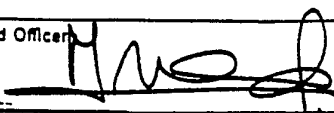
(ii) thereafter adding the fabric softening clay material and the one or more detergency active materials;

25

(iii) thereafter adding the remaining portion (if any) of the detergency builder.

INTERNATIONAL SEARCH REPORT

International Application No PCT/GB 86/00712

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 IPC ⁴ : C 11 D 3/12; C 11 D 17/00											
II. FIELDS SEARCHED Minimum Documentation Searched ⁷ <table border="1"> <thead> <tr> <th>Classification System</th> <th>Classification Symbols</th> </tr> </thead> <tbody> <tr> <td>IPC⁴</td> <td>C 11 D</td> </tr> </tbody> </table> Documentation Searched other than Minimum Documentation to the Extent that such Documents are Included in the Fields Searched ⁸			Classification System	Classification Symbols	IPC ⁴	C 11 D					
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III. DOCUMENTS CONSIDERED TO BE RELEVANT ⁹ <table border="1"> <thead> <tr> <th>Category ¹⁰</th> <th>Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²</th> <th>Relevant to Claim No. ¹³</th> </tr> </thead> <tbody> <tr> <td>A</td> <td>FR, A, 2537595 (COLGATE-PALMOLIVE) 15 June 1984 see the claims & GB, A, 2132629 (cited in the application) --</td> <td>1</td> </tr> <tr> <td>P,A</td> <td>GB, A, 2168717 (COLGATE-PALMOLIVE) 25 June 1986 see the claims -----</td> <td>1</td> </tr> </tbody> </table>			Category ¹⁰	Citation of Document, ¹¹ with indication, where appropriate, of the relevant passages ¹²	Relevant to Claim No. ¹³	A	FR, A, 2537595 (COLGATE-PALMOLIVE) 15 June 1984 see the claims & GB, A, 2132629 (cited in the application) --	1	P,A	GB, A, 2168717 (COLGATE-PALMOLIVE) 25 June 1986 see the claims -----	1
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IV. CERTIFICATION <table border="1"> <tr> <td> Date of the Actual Completion of the International Search 12th February 1987 </td> <td> Date of Mailing of this International Search Report 25 FEB. 1987 </td> </tr> <tr> <td> International Searching Authority EUROPEAN PATENT OFFICE </td> <td> Signature of Authorized Officer M. VAN MOL  </td> </tr> </table>			Date of the Actual Completion of the International Search 12th February 1987	Date of Mailing of this International Search Report 25 FEB. 1987	International Searching Authority EUROPEAN PATENT OFFICE	Signature of Authorized Officer M. VAN MOL 					
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ANNEX TO THE INTERNATIONAL SEARCH REPORT ON

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