

FORM 1

613792

COMMONWEALTH OF AUSTRALIA

PATENTS ACT 1952

APPLICATION FOR A STANDARD PATENT

I\We,

UNILEVER PLC

of

UNILEVER HOUSE, BLACKFRIARS
LONDON EC4
ENGLAND

hereby apply for the grant of a standard patent for an
invention entitled:

FABRIC TREATMENT COMPOSITION.

which is described in the accompanying complete specification

Details of basic application(s):

Number of basic application	Name of Convention country in which basic application was filed	Date of basic application
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8822206.2

GB

21 SEP 88

My/our address for service is care of GRIFFITH HACK & CO.,
Patent Attorneys, 601 St. Kilda Road, Melbourne 3004,
Victoria, Australia.

DATED this 20th day of September 1989

UNILEVER PLC

GRIFFITH HACK & CO.

TO: The Commissioner of Patents.

012.00 200989

AUSTRALIA

B

PATENTS ACT 1952

APPLICATION
BY ASSIGNEE
OF INVENTOR

DECLARATION IN SUPPORT OF AN APPLICATION
FOR A PATENT

In support of an application made by: UNILEVER PLC
for a patent for an invention entitled:

FABRIC TREATMENT COMPOSITION

I, Peter Geoffrey Mole
of Unilever House, Blackfriars, London EC4, England,

do solemnly and sincerely declare as follows:

1. I am authorised by the above mentioned applicant for the patent to make this declaration on its behalf.
2. The name and address of each actual inventor of the invention is as follows:

Christine MORRIS and Graham Andrew TURNER, both British Subjects of respectively 31 St James Avenue, Upton by Chester, Chester, Cheshire CH2 1NB Great Britain and 2 Redbrook Close, Bromborough, Wirral, Merseyside, L62 6EA Great Britain.

3. The facts upon which the applicant is entitled to make this application are as follows:

The applicants are the assignee of the employer of the actual inventors, namely UNILEVER UK CENTRAL RESOURCES LIMITED. UNILEVER UK CENTRAL RESOURCES LIMITED was entitled to have assigned to it a patent granted to any of the actual inventors in respect of the said invention.

4. The basic application as defined by Section 141 of the Act was made as follows:

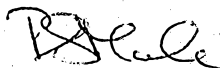
In Great Britain on 21 September 1988
by UNILEVER PLC

5. The basic application referred to in the preceding paragraph was the first application made in a Convention country in respect of the invention the subject of this application.

Declared at London, England

this 2nd. day of May 1991

Signed



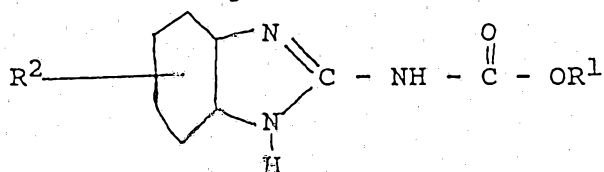
Position:

Authorised Signatory

(12) PATENT ABRIDGMENT (11) Document No. AU-B-41536/89
(19) AUSTRALIAN PATENT OFFICE (10) Acceptance No. 613792

- (54) Title
BENZIMIDAZOLE CARBAMATE ANTI-MICROBIAL
- International Patent Classification(s)
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- (21) Application No. : 41536/89 (22) Application Date : 20.09.89
- (30) Priority Data
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8822206 21.09.88 GB UNITED KINGDOM
- (43) Publication Date : 29.03.90
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- (71) Applicant(s)
UNILEVER PLC
- (72) Inventor(s)
CHRISTINE MORRIS; GRAHAM ANDREW TURNER
- (74) Attorney or Agent
GRIFFITH HACK & CO , GPO Box 1285K, MELBOURNE VIC 3001
- (57) Claim

1. Fabric treatment composition comprising a fabric-softening material and a benzimidazolyl-alkyl-carbamate.
2. Fabric treatment composition according to Claim 1, characterised in that the benzimidazolyl-alkyl-carbamate is of the following formula:



where R¹ is a C₁₋₄ alkyl, alkoxy or phenoxy alkyl; and
R² is a C₁₋₄ alkyl, hydrogen, nitrogen or halogen.

AUSTRALIA

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PATENTS ACT 1952

Form 10

COMPLETE SPECIFICATION

(ORIGINAL)

FOR OFFICE USE

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Related Art:

TO BE COMPLETED BY APPLICANT

Name of Applicant:

UNILEVER PLC

Address of Applicant: UNILEVER HOUSE
BLACKFRIARS
LONDON EC4
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Actual Inventor:

Address for Service: GRIFFITH HACK & CO.,
601 St. Kilda Road,
Melbourne, Victoria 3004,
Australia.

Complete Specification for the invention entitled:
FABRIC TREATMENT COMPOSITION.

The following statement is a full description of this invention
including the best method of performing it known to me:-

FABRIC TREATMENT COMPOSITION

This invention relates to a fabric treatment composition for treating fabrics during the rinse step of a fabric laundry process, and to a method for using said composition. In particular, the invention relates to a fabric treatment composition comprising a fabric-softening material, which is preferably a cationic fabric softening material and an antimicrobial agent.

It is known that fungal growth can develop on fabrics stored under certain physiological conditions, such as in a humid atmosphere or where there is poor air circulation. These growths can damage the fabrics by causing them to discolour. The fabrics can also develop an unpleasant smell. It has been found that the discolouration or staining of the fabrics is difficult to remove either by normal laundering or after treatment with a strong bleaching agent.

It has previously been proposed (EP 86 423, Henkel KGaA) to incorporate specific antimicrobial substances in fabric treatment compositions.

We have now surprisingly discovered that benzimidazolyl-alkyl-carbamates, when combined with fabric-softening materials into a fabric treatment composition, are particularly effective in inhibiting fungal growth on fabrics which have been rinsed with said fabric treatment composition.

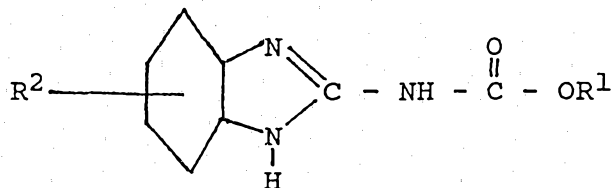
The incorporation of benzimidazolyl-alkyl-carbamates in textile treatment compositions according to the invention has also been found advantageous in that benzimidazolyl-alkyl-carbamates are:

1. effective against a wide range of micro-organisms, particularly fungi;
2. active during the life of the product;
3. of low mammalian toxicity and non-toxic to humans
- 5 at the concentrations used;
4. colourless and odourless;
5. effective at low concentrations;
6. inexpensive and easy to apply;
7. resistant to sunlight;
- 10 8. not affecting fabric handle or strength;
9. compatible with water-repellent and flame-retardant agents, dyes and other textile auxiliaries; and
10. do not sensitize the fabric to damage by light or other influences.

15 Thus, according to the invention there is provided a fabric treatment composition comprising a softening material and a benzimidazolyl-alkyl-carbamate. Preferably the softening material comprises a cationic
20 fabric softener.

Benzimidazolyl-alkyl-carbamates are well-known fungicides for use in agriculture. It has also been suggested in Japanese patent application 49 093 537
25 (Yoshitomi) to treat textiles with a composition containing Me-2-benzimidazole-carbamate in a solvent. This treatment, however, requires a separate processing step for applying the benzimidazolyl-alkyl-carbamate and a further separate processing step for removing the
30 undesirable solvent compound.

The benzimidazolyl-alkyl-carbamate as used in a textile treatment composition of the invention is preferably of the following formula:



wherein:

- R^1 is a C_{1-4} alkyl, alkoxy or phenoxy alkyl; and
 R^2 is a C_{1-4} alkyl, hydrogen, nitrogen or halogen.

Particular examples of suitable benzimidazolyl-alkyl-carbamate substances are: benzimidazolyl-methyl-carbamate, benzimidazolyl-ethyl-carbamate, 5-ethyl-benzimidazolyl-ethyl-carbamate, benzimidazolyl-isopropyl-carbamate, 4-ethyl-benzimidazolyl-isopropyl-carbamate, 4-propyl-benzimidazolyl-isopropyl-carbamate, 4-butyl-benzimidazolyl-isopropyl-carbamate, 4-isobutyl-benzimidazolyl-isopropyl-carbamate, benzimidazolyl-ethylmethoxy-carbamate, 4-ethyl-benzimidazolyl-ethylmethoxy-carbamate, 5-ethyl-benzimidazolyl-ethylmethoxy-carbamate, benzimidazolyl-ethylethoxy-carbamate, benzimidazolyl-ethyl-propoxy-carbamate, benzimidazolyl-ethylphenoxy-carbamate, 4-ethyl-benzimidazolyl-ethylphenoxy-carbamate and 5-ethyl-benzimidazolyl-ethylphenoxy-carbamate.

The preferred benzimidazolyl-alkyl-carbamate is benzimidazolyl-methyl-carbamate.

It is also possible to employ mixtures of different benzimidazolyl-alkyl-carbamates.

Particularly preferred is the use of Intace B 350 (75% carbendazim ex Intace), and Intace B 300 (100% carbendazim ex Intace).

Other antimicrobial substances may be incorporated in a fabric treatment composition according to the invention. By antimicrobial substances are meant both antibacterial and antifungal products. Examples of such material are

formaldehyde, halogenated phenols, nitrated mono- or polyalcohols and antimicrobial surface-active tensides.

Particularly suitable for use in conjunction with
5 benzimidazolyl-alkyl-carbamates are tetra-methylthiuram-disulphid, 1-methylol-2-thiono-2-dihydrobenzthiazol, N-dimethyl-N'-phenyl-(N'-fluordichloromethyl-thio) sulfphamid and 3-isothiazolone compounds such as for instance described in US 4,424,134. Generally, the
10 amount of additional antimicrobial substances in the textile treatment composition is less than two times the amount of benzimidazolyl-alkyl-carbamate.

The fabric softening materials may be selected from
15 cationic, nonionic, amphoteric or anionic fabric softening materials.

Suitable amphoteric fabric conditioning material for use in a composition according to the invention are fabric
20 substantive amphoteric materials forming a particulate dispersion at a concentration of less than 1 g/l at at least one temperature between 0 and 100°C. Preferably at at least one temperature between 10 and 90 °C, more preferred between 20 and 80 °C. For the purpose of this
25 invention a fabric substantive amphoteric material is preferably an amphoteric or zwitterionic tertiary or quaternary ammonium compound having either one single long hydrocarbyl side chain or two long hydrocarbyl chains. From these compounds the use of amphoteric or
30 zwitterionic ammonium compounds having two long hydrocarbyl chains is particularly preferred for many reasons including costs, ease of processing and better stability and performance. Suitable amphoteric materials are for example disclosed in EP 89200545.5.

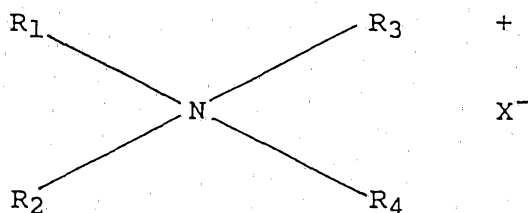
35 In this specification the expression hydrocarbyl chain refers to linear or branched alkyl or alkenyl chains optionally substituted or interrupted by functional

groups such as -OH, -O-, -CONH-, -COO-, etc.

Preferably the amphoteric fabric substantive materials are water insoluble and have a solubility in water at pH 2.5 at 20°C of less than 10 g/l. The HLB of the amphoteric fabric substantive material is preferably less than 10.0.

Suitable cationic fabric softener materials for use in a composition according to the present invention are cationic materials which are water-insoluble in that the material has a solubility in water at pH 2.5 and 20°C of less than 10 g/l. Highly preferred materials are cationic quaternary ammonium salts having two C12-24 hydrocarbyl chains.

Well-known species of substantially water-insoluble quaternary ammonium compounds have the formula



wherein R_1 and R_2 represent hydrocarbyl groups from about 12 to about 24 carbon atoms; R_3 and R_4 represent hydrocarbyl groups containing from 1 to about 4 carbon atoms; and X is an anion, preferably selected from halide, methyl sulfate and ethyl sulfate radicals.

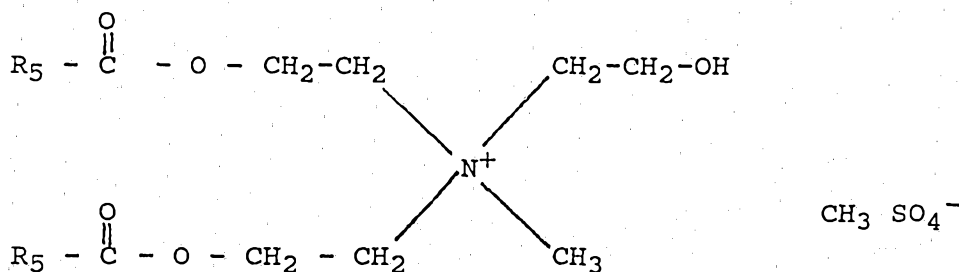
Representative examples of these quaternary softeners include ditallow dimethyl ammonium chloride; ditallow dimethyl ammonium methyl sulfate; dihexadecyl dimethyl ammonium chloride; di(hydrogenated tallow) dimethyl ammonium methyl sulfate; dihexadecyl diethyl ammonium chloride; di(coconut) dimethyl ammonium chloride.

Ditallow dimethyl ammonium chloride, di(hydrogenated tallow) dimethyl ammonium chloride, di(coconut) dimethyl ammonium chloride and di(coconut) dimethyl ammonium methosulfate are preferred.

5 Suitable materials also include dialkyl ethoxyl methyl ammonium methosulphate based on soft fatty acid, dialkyl ethoxyl methyl ammonium methosulphate based on hard fatty acid, and a material in which R_3 and R_4 represent methyl, R_1 is C_{13-15} , R_2 is CH_2CH_2OCOR , where R is stearyl, and X is methosulphate. Ditallow dimethyl ammonium chloride, di(hydrogenated tallow alkyl) dimethyl ammonium chloride, di(coconut alkyl) dimethyl ammonium chloride and di(coconut alkyl) dimethyl ammonium methosulfate are preferred.

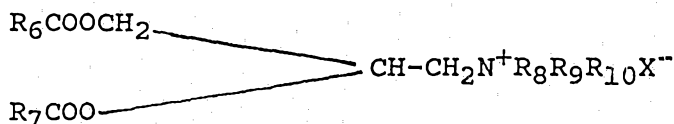
Other preferred cationic compounds include those materials as disclosed in EP 239,910 (P&G), which is included herein by reference.

Other preferred materials are the materials of formula



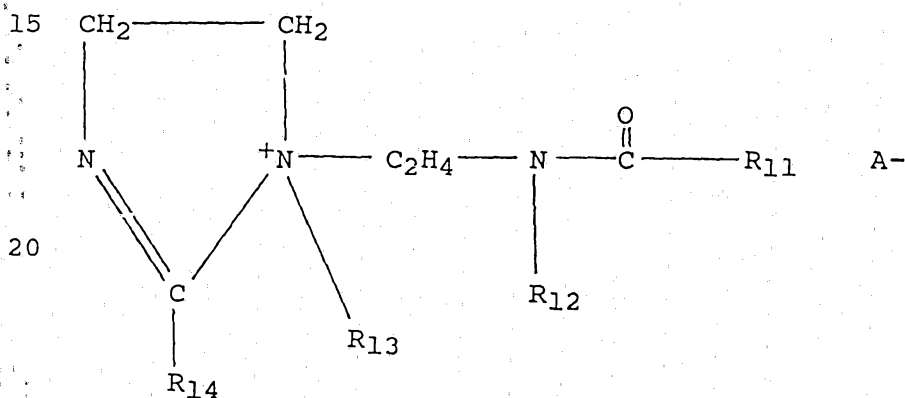
R_5 being tallow, which is available from Stepan under the tradename Stepantex VRH 90

and



where R_8 , R_9 and R_{10} are each alkyl or hydroxyalkyl groups containing from 1 to 4 carbon atoms, or a benzyl group. R_6 and R_7 are each an alkyl or alkenyl chain containing from 11 to 23 carbon atoms, and X^- is a water soluble anion. These materials and their method of preparation are described in US 4,137,180 (LEVER BROTHERS).

- 10 Another class of preferred water-insoluble cationic materials are the hydrocarbylimidazolinium salts believed to have the formula:



wherein R_{13} is a hydrocarbyl group containing from 1 to 4, preferably 1 or 2 carbon atoms, R_{11} is a hydrocarbyl group containing from 8 to 25 carbon atoms, R_{14} is an hydrocarbyl group containing from 8 to 25 carbon atoms and R_{12} is hydrogen or an hydrocarbyl containing from 1 to 4 carbon atoms and A^- is an anion, preferably a halide, methosulfate or ethosulfate.

Preferred imidazolinium salts include 1-methyl-1-(tallowylamido-) ethyl -2-tallowyl- 4,5-dihydro imidazolinium methosulfate and 1-methyl-1-(palmitoylamido)ethyl -2-octadecyl-4,5- dihydro- imidazolinium chloride. Other useful imidazolinium

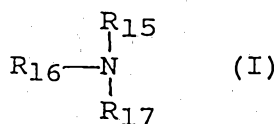
materials are 2-heptadecyl-1-methyl-1- (2-stearyl-amido)-
ethyl-imidazolinium chloride and 2-lauryl-1-
hydroxyethyl-1-oleyl-imidazolinium chloride. Also
suitable herein are the imidazolinium fabric softening
5 components of US patent No 4 127 489, incorporated by
reference.

Representative commercially available materials of the
above classes are the quaternary ammonium compounds
10 Arquad 2HT (ex AKZO); Noranium M2SH (ex CEKA); Aliquat-
2HT (Trade Mark of General Mills Inc), Stepantex Q185
(ex Stepan); Stepantex VP85 (ex Stepan); Stepantex VRH90
(ex Stepan); Synprolam FS (ex ICI) and the imidazolinium
compounds Varisoft 475 (Trade Mark of Sherex Company,
15 Columbus Ohio) and Rewoquat W7500 (Trade Mark of REWO).

The compositions according to the invention may also
possibly in addition to the above mentioned softening
agents, one or more amine softening materials.

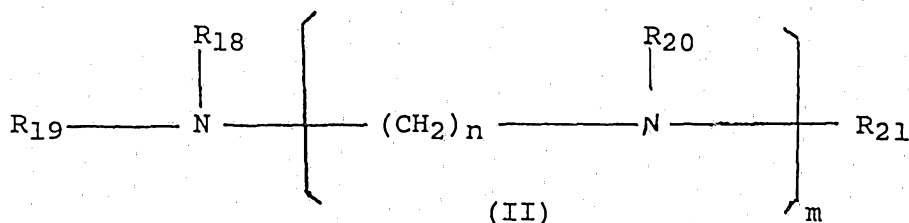
20 The term "amine" as used herein can refer to

(i) amines of formula



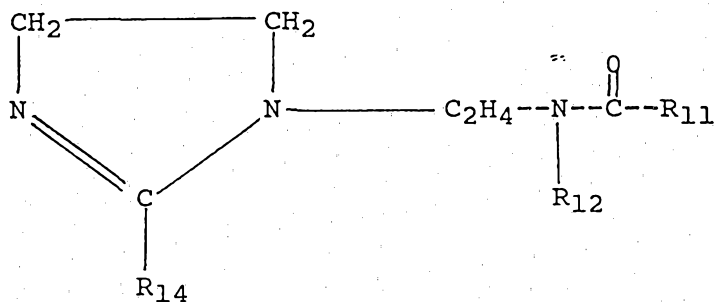
wherein R₁₅, R₁₆ and R₁₇ are defined as below;

30 (ii) amines of formula



wherein R_{18} , R_{19} , R_{20} and R_{21} , m and n are defined as below.

5 (iii) imidazolines of formula



15 wherein R_{11} , R_{12} and R_{14} are defined as above.

(iv) condensation products formed from the reaction of fatty acids with a polyamine selected from the group consisting of hydroxy alkylalkylenediamines and dialkylenetriamines and mixtures thereof. Suitable materials are disclosed in European Patent Application 199 382 (Procter and Gamble), incorporated herein by reference.

25 When the amine is of the formula I above, R_{15} is a C_6 to C_{24} , hydrocarbyl group, R_{16} is a C_1 to C_{24} hydrocarbyl group and R_{17} is a C_1 to C_{10} hydrocarbyl group.

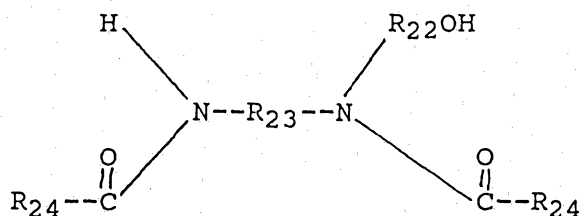
Suitable amines include those materials from which the quaternary ammonium compounds disclosed above are derived, in which R_{15} is R_1 , R_{16} is R_2 and R_{17} is R_3 . Preferably, the amine is such that both R_{15} and R_{16} are C_6 - C_{20} alkyl with C_{16} - C_{18} being most preferred and with R_{17} as C_1 - C_3 alkyl, or R_{15} is an alkyl or alkenyl group with at least 22 carbon atoms and R_{16} and R_{12} are C_1 - C_3 alkyl. Preferably these amines are protonated with hydrochloric acid, orthophosphoric acid (OPA), C_1 - C_5 carboxylic acids or any other similar acids, for use in the fabric conditioning compositions of the invention.

When the amine is of formula II above, R_{18} is a C_6 to C_{24} hydrocarbonyl group, R_{19} is an alkoxyated group of formula $-(CH_2CH_2O)_yH$, where y is within the range from 0 to 6, R_{20} is an alkoxyated group of formula $-(CH_2CH_2O)_zH$ where z is within the range from 0 to 6 and m is an integer within the range from 0 to 6, and is preferably 3. When m is 0, it is preferred that R_{18} is a C_{16} to C_{22} alkyl and that the sum total of z and y is within the range from 1 to 6, more preferably 1 to 3. When m is 1, it is preferred that R_{18} is a C_{16} to C_{22} alkyl and that the sum total of x and y and z is within the range from 3 to 10.

Representative commercially available materials of this class include Ethomeen (ex Armour) and Ethoduomeen (ex Armour).

Preferably the amines of type (ii) or (iii) are also protonated for use in the fabric conditioning compositions of the invention.

When the amine is of type (iv) given above, a particularly preferred material is



where R_{22} and R_{23} are divalent alkenyl chains having from 1 to 3 carbon atoms, and R_{24} is an acyclic aliphatic hydrocarbon chain having from 15 to 21 carbon atoms. A commercially available material of this class is Ceranine HC39 (ex Sandoz).

The compositions according to the invention may also

contain nonionic fabric-softening agents. Suitable nonionic fabric-softening agents include glycerol esters, such as glycerol monostearate, fatty alcohols, such as stearyl alcohol, alkoxylated fatty alcohols, 5 fatty acids, lanolin and derivatives thereof. Suitable materials are disclosed in European Patent Applications 88 520 (Unilever), 122 141 (Unilever) and 79 746 (Procter & Gamble), the disclosures of which are incorporated herein by reference. Prefererably, such 10 materials are included at a level within the range of from 0.5% to 10% by weight of the composition.

The compositions can also contain one or more optional ingredients selected from non-aqueous solvents such as 15 C₁-C₄ alkanols and polyhydric alcohols, pH-buffering agents such as weak acids, e.g. phosphoric, benzoic or citric acids (the compositions preferably have a pH of less than 6.0 more preferred between 2.0 and 5.0), re-wetting agents, viscosity modifiers such as 20 electrolytes, included at levels from about 20 to 6000 ppm, aluminium chlorohydrate, antigelling agents, perfumes especially body odour reducing perfumes, perfume carriers, hydrocarbons, fluorescers, colourants, hydrotropes, antifoaming agents, 25 antiredeposition agents, enzymes, optical brightening agents, opacifiers, stabilisers such as guar gum and polyethylene glycol, anti-shrinking agents, anti-wrinkle agents, soil-release agents, antioxidants, anti-corrosion agents, preservatives such as Bronopol (Trade 30 .k), a commercially available form of 2-bromo-2-nitropropane-1,3-diol, to preserve the fabric treatment composition, dyes, bleaches and bleach precursors, drape-imparting agents, antistatic agents and ironing aids.

35 Silicones can be included in the compositions as the ironing aid, re-wetting agent or the antifoaming agent. Suitable silicones for use in the compositions according

to the invention include predominantly linear polydialkyl or alkylaryl siloxanes in which the alkyl groups contain one to five carbon atoms. The siloxanes can be amido- or amino-substituted. When the siloxane is amine-substituted the amine group may be quaternised.

These optional ingredients, if added, are each present at levels up to 5% by weight of the composition.

10 The compositions according to the invention may take a variety of forms. They may for example be liquids containing an aqueous base.

Alternatively, the compositions according to the invention may be in the form of a powder, creams, pastes, a block or a tablet. Suitable types of blocks and tablets are disclosed in European Patent Specification EP-A-255 779 (Unilever).

20 When the compositions are in the form of liquid products, the total level of the fabric softening material and the benzimidazolyl-alkyl-carbamate is preferably from 1 to 30% by weight. Based on the total weight of the composition, the weight of the fabric-softening agent is preferably from 1 to 20% and the weight of the benzimidazolyl-alkyl-carbamate is preferably from 0.05 to 10%, more preferably from 0.1 to 5%, most preferably from 0.5 to 1.5%. The weight ratio of softening agent to benzimidazolyl-alkyl-carbamate is preferably between 10:1 and 100:1, more preferably around 20:1. Preferably more than 50 wt % of the softening materials are cationic softening materials.

35 When the compositions are in the form of powders, blocks or tablets, they may contain means for restraining the release of the cationic fabric-softening agent and benzimidazolyl-alkyl-carbamate in water, such as

paraffin and tallow alcohol. Other suitable materials are disclosed in the European Patent Specification EP-A-255 779 (Unilever), the disclosure of which is incorporated herein by reference. The compositions may also contain an electrolyte to aid dispersion of the cationic fabric-softening agent and the benzimidazolyl-alkyl-carbamate, after the release inhibitor has ceased to have effect, and an antifoam material. A suitable antifoam granule may be of the type described in our European Patent Specification EP-A-94 250.

In use, the compositions are preferably added to a large volume of water to form a liquor with which the fabrics to be treated are contacted. The liquor can be formed by adding the fabric treatment composition to water during the rinse stage of a hand-washing process.

Alternatively, the composition can be added by hand or by way of an automatic dispensing device to rinse water in the tub of a washing machine of the type which is common in Europe, that is a machine in which the fabrics are rinsed in a tub which is filled with rinse water, agitated to achieve efficient rinsing and then drained. Generally, the total concentration of the fabric-softening agent and the benzimidazolyl-alkyl-carbamate in the rinse liquor will be between 30 and 500 ppm.

Alternatively, the compositions, especially when being in the form of a tablet or block, can be used in a washing machine which is designed in such a manner that the fabrics are rinsed in running water. With a machine of this type, the fabrics will normally have been previously washed in the same tub of the machine, or they may be placed in the tub specifically for rinsing. When previously washed, they will have at least some of the wash liquor containing a detergent active still adsorbed on or otherwise associated with them, the running water then serving to flush this wash liquor out

of the fabrics. The rinse water running into the tub will be substantially free of any detergent actives. The composition can be added to the fabrics in the tub before the rinse water flows into the tub,
5 simultaneously with the rinse water or part-way through the rinse cycle.

The invention will now be illustrated by the following non-limiting examples. In these examples, all parts and percentages are by weight. Where commercially available materials are referred to, the percentages quoted are those of the active detergent therein.

EXAMPLES 1-6

Aqueous compositions were prepared according to the formulations in the following tables. The compositions were prepared by co-melting the cationic fabric-softening agent and benzimidazolyl-alkyl-carbamate at 80°C. The co-melt thus formed was added to water at 80°C to form a dispersion. In examples containing 2.5 and 5.0% carbendazim respectively, the benzimidazolyl-alkyl-carbamate was post-dosed to a dispersion containing the cationic fabric-softening agent.

The compositions were tested by the following method.

Two pieces of a 50/50 resinated polyester cotton fabric and two pieces of a 100% cotton poplin fabric (each measuring 20 cm x 20 cm) were washed at 50°C with 5 grams of a commercially available washing powder, Persil Automatic, in a litre of water in a tergotometer. After the wash liquor was separated, the fabrics were rinsed twice with a litre of demineralised water. 2 ml of an aqueous test composition was added to the second rinse. After line dry, the treated fabrics were assessed, as follows, to evaluate the microbial activity present.

Two pieces, each measuring 5 cm x 5 cm, were cut from each piece of treated fabric. Both pieces were placed on Petri dishes containing nutrient agar. 0.1 ml of an inoculum of either Aspergillus niger or Penicillium spores was spotted onto the centre of each piece. Care was taken not to allow the inoculum to spread onto the surrounding agar. The inoculum concentration was 10^8 spores ml^{-1} . The plates were incubated at 28°C and evaluation of fungal growth assessed after 3 days.

The scale used to assess fungal growth was:

	0	=	No growth on fabric.
	10	=	Very slight growth.
5	20	=	Visible, light, patchy growth.
	30	=	Moderate growth on whole sample.
	40	=	Heavy growth on most of sample.
	50	=	Completely overgrown.

- 10 Owing to discolouration of the fabric, caused by spore inoculum, a fungal growth score of 0 is never attained. A score of less than 10 is indicative of no growth.

EXAMPLE		50/50 POLYESTER COTTON		COTTON POPLIN	
N°		<u>A. niger</u> <u>Penicillium</u>		<u>A. niger</u> <u>Penicillium</u>	
Reference					
	Water	41	39	50	49
	1. 5%				
20	Arquad 2HT	40	37	44	49
	2. 5% 2HT +				
	0.1% C*	35	28	40	15
	3. 5% 2HT +				
25	0.25% C	30	30	35	< 10
	4. 5% 2HT +				
	0.50% C	25	< 10	30	< 10
30	5. 5% 2HT +				
	2.5% C	18	< 10	< 10	< 10
	6. 5% 2HT +				
35	5.0% C	< 10	< 10	< 10	< 10

* C = Carbendazim (Intace B 350)

The results show that even if carbendazim is incorporated in a fabric treatment composition containing Arquad 2HT, at low levels, such as 0.1%, there is a significant reduction in fungal growth.

5

EXAMPLES 7-11

Compositions according to the following tables were prepared as described in Example 1. Fabrics were evaluated as described for Example I. The following results were obtained.

EXAMPLE N°	<u>50/50 POLYESTER COTTON</u>		<u>COTTON POPLIN</u>	
	<u>A. niger</u>	<u>Penicillium</u>	<u>A. niger</u>	<u>Penicillium</u>
15	Water	50	47.5	50
20	7. 5%			
	Arquad 2HT	37.5	35	50
	8. 5%			
	Arquad 2HT			
	+ 0.5% C*	25	< 10	30
25	9. 5%			
	Arquad 2HT			
	+ 0.5% K**	32.5	42.5	42.5
30	10. 5%			
	Arquad 2HT			
	+ 10% A***	25	25	47.5
35	11. 5%			
	Arquad 2HT			
	+ 0.5% P****	35	50	50

* C = Carbendazim (Intace B 350)

** K = Kathon (2-n-octyl-4-isothiazolin-3-one)

ex Rohm & Haas)

*** A = Actifresh L 8110, which comprises benzyl-alkyl-bis(hydroxy alkyl)ammonium, ex British Sanitized

5 **** P = Preventol R 89 (alkyl dimethyl benzyl ammonium chloride, ex Bayer)

These examples clearly show the surprisingly better performance of textile treatment compositions comprising
10 benzimidazolyl-alkyl-carbamate antimicrobial substances.

EXAMPLES 12-16

15 Tablets comprising 8 g sodium bicarbonate, 2 g Adogen and 0.4 g carbendazim (Intace B 300) were prepared and compared with tablets comprising 8 g sodium bicarbonate and 2 g Adogen.

20 Pieces of 50/50 resinated polyester cotton fabrics and pieces of a 100% cotton poplin fabric were washed in a Japanese washing machine at room temperature with 31 g of a commercially available wash powder. After a 1 minute spin to separate the wash liquor, the fabrics were rinsed for ten minutes in running water. One tablet
25 was added and the rinse continued for a further 5 minutes. After line drying, the treated fabrics were assessed for microbial activity as in Example 1.

The results were the following:

Fabric	Treatment	<u>Fungal Growth</u>	
		Aspergillus	Penicillium
50/50 Polycotton	Control*	40-45	40
100% Cotton poplin	Control	50	50
50/50 PC	Softener + 0.4 g Carb.	< 10	< 10
10 100% CP	Softener + 0.4 g Carb.	< 10	< 10

* Control = Tablet containing Adogen and NaHCO_3 .

Examples 17-30

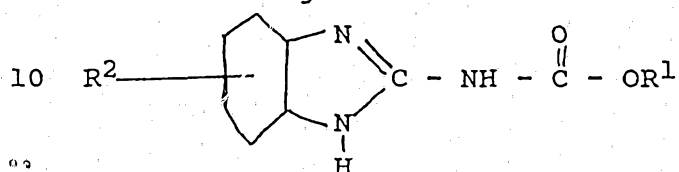
The following aqueous compositions were prepared as in examples 1-6 and tested on cotton as in these examples. The carbendazim used was Intace B300. These examples again show a reduction in fungal growth when carbendazim is added to fabric softening compositions of the invention.

		<u>Fungal Growth</u>	
<u>Product</u>		<u>A Niger</u>	<u>Penicillium</u>
	17. Water	50	50
5	18. 7.56% Stepantex VRH 90	50	50
	19. 7.56% Stepantex VRH 90 + 1.0% Carbendazim	0-10	5
	20. 5% DTTMAPC (1,2 ditallow oxytrimethylammonium propane chloride)	50	50
10	21. 5% DTTMAPC + 1% Carbendazim	0-10	0
	22. 5% Imidazolinium (Rewoquat W7500H)	50	50
	23. 5% Imidazolinium + 1.0% Carbendazim	0-5	0
15	24. 10.8% Dihardened tallow dimethyl ammonium chloride (DHTMAC) 2.7% Hardened tallow fatty acid	40	50
	25. 10.8% DHTMAC 2.7% Hardened tallow fatty acid 1.0% Carbendazim	20	0-10
20	26. 6% DHTDMAC	50	50
	27. 6% DHTDMAC + 1.0% Aluminium Chlorohydrate (ACH)	50	50
25	28. 6% DHTDMAC + 1.0% ACH + 1.0% Carbendazim	10	0
	29. 3.8% DHTDMAC 1.9% Ceranine HC39 (ex Sandoz) 1.9% DHTM amine oxide (Lilaminox M44, ex Berol Nobel) 0.38% Orthophosphoric Acid	50	50
30	30. 3.8% DHTDMAC 1.9% Ceranine HC39 1.9% DHTM Amine Oxide 0.38% Orthophosphoric acid + 1.0% Carbendazim	30	10-15
35			

THE CLAIMS DEFINING THE INVENTION ARE AS FOLLOWS:

1. Fabric treatment composition comprising a fabric-softening material and a benzimidazolyl-alkyl-carbamate.

2. Fabric treatment composition according to Claim 1, characterised in that the benzimidazolyl-alkyl-carbamate is of the following formula:



where R^1 is a C_{1-4} alkyl, alkoxy or phenoxy alkyl; and
 R^2 is a C_{1-4} alkyl, hydrogen, nitrogen or halogen.

3. Fabric treatment composition according to Claim 1 or 2, characterised in that it comprises:
 - 1 - 30% by weight of the fabric-softening material, and
 - 0.1 - 5% by weight of the benzimidazolyl-alkyl-carbamate.

4. Fabric treatment composition according to claim 1, 2 or 3 comprising a cationic fabric softening material.

5. Fabric treatment composition according to ^{any one of} Claims 1 to 4 ~~1-4~~, characterised in that it further comprises at least one other antimicrobial ^{agent} ~~substance~~.

6. Method for treating fabrics by rinsing the fabrics in an aqueous bath comprising a textile treatment composition comprising a fabric-softening material and a benzimidazolyl-alkyl-carbamate, wherein the concentration of the fabric softening material plus benzimidazolyl-alkyl-carbamate composition in the bath is from 30 to 500 ppm.

DATED THIS 20TH DAY OF SEPTEMBER 1989

UNILEVER PLC

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