(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization

International Bureau



(10) International Publication Number WO 2010/057968 A1

(43) International Publication Date 27 May 2010 (27.05.2010)

(51) International Patent Classification: A61K 8/04 (2006.01) A61K 8/92 (2006.01)

(21) International Application Number:

PCT/EP2009/065537

(22) International Filing Date:

A61K 8/26 (2006.01)

20 November 2009 (20.11.2009)

A61Q 15/00 (2006.01)

(25) Filing Language:

English

(26) Publication Language:

English

US

EP

(30) Priority Data:

24 November 2008 (24.11.2008) 61/117,288 09155203.4 16 March 2009 (16.03.2009)

- (71) Applicant (for AE, AG, AU, BB, BH, BW, BZ, CA, CY, EG, GB, GD, GH, GM, IE, IL, KE, KN, LC, LK, LS, MT, MW, MY, NA, NG, NZ, OM, PG, SC, SD, SG, SL, SZ, TT, TZ, UG, VC, ZA, ZM, ZW only): UNILEVER PLC [GB/ GB]; a company registered in England and Wales under company no. 41424, Unilever House, 100 Victoria Embankment, London Greater London EC4Y 0DY (GB).
- Applicant (for all designated States except AE, AG, AU, BB, BH, BW, BZ, CA, CY, EG, GB, GD, GH, GM, IE, IL, IN, KE, KN, LC, LK, LS, MT, MW, MY, NA, NG, NZ, OM, PG, SC, SD, SG, SL, SZ, TT, TZ, UG, US, VC, ZA, ZM, ZW): UNILEVER N.V. [NL/NL]; Weena 455, NL-3013 AL Rotterdam (NL).
- (71) Applicant (for IN only): HINDUSTAN UNILEVER LIMITED [IN/IN]; Hindustan Lever House, 165/166 Backbay Reclamation, Maharashtra, Mumbai 400 020 (IN).
- (72) Inventors; and
- (75) Inventors/Applicants (for US only): BUTTERWORTH, Andrew [GB/GB]; Unilever UK Limited, Coal Road, Seacroft, Leeds Yorkshire LS14 2AR (GB). FERRIER, Lindsay, Karen [GB/GB]; Unilever UK Limited, Coal Road, Seacroft, Leeds Yorkshire LS14 2AR (GB). FLETCHER, Neil, Robert [GB/GB]; Unilever R&D Port Sunlight, Quarry Road East, Bebington, Wirral

Mersevside CH63 3JW (GB). JONES, Shirley [GB/GB]; Unilever UK Limited, Coal Road, Seacroft, Leeds Yorkshire LS14 2AR (GB). MARRIOTT, Robert, Edward [GB/GB]; Unilever R&D Port Sunlight, Quarry Road East, Bebington, Wirral Merseyside CH63 3JW (GB). POLONKA, Jack [US/US]; Conopco Inc., d/b/a UNILEVER, 40 Merritt Boulevard, Trumbull, Connecticut 06611 (US). WILLIAMS, Jason, Richard [GB/GB]; Unilever R&D Port Sunlight, Quarry Road East, Bebington, Wirral Merseyside CH63 3JW (GB).

- (74) Agent: WHALEY, Christopher; Unilever PLC, Unilever Patent Group, Colworth House, Sharnbrook, Bedford Bedfordshire MK44 1LQ (GB).
- (81) Designated States (unless otherwise indicated, for every kind of national protection available): AE, AG, AL, AM, AO, AT, AU, AZ, BA, BB, BG, BH, BR, BW, BY, BZ, CA, CH, CL, CN, CO, CR, CU, CZ, DE, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, JP, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LY, MA, MD, ME, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PE, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW.
- (84) Designated States (unless otherwise indicated, for every kind of regional protection available): ARIPO (BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European (AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HR, HU, IE, IS, IT, LT, LU, LV, MC, MK, MT, NL, NO, PL, PT, RO, SE, SI, SK, SM, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG).

Published:

- with international search report (Art. 21(3))
- before the expiration of the time limit for amending the claims and to be republished in the event of receipt of amendments (Rule 48.2(h))



(57) Abstract: Anhydrous antiperspirant aerosol compositions to improve the soft focus of the film of antiperspirant deposited on skin, and particularly on underarm skin, comprise in addition to a particulate aluminium antiperspirant salt, an oil and a propellant, at least one smoothing aid selected from a low concentration of wax and mica and preferably both. Advantageously, the wax comprises stearyl alcohol and the mica is a mica interference pigment, preferably exhibiting blue highlights. In some preferred embodiments, the compositions include a moisturiser such as an occlusive agent and/or a triglyceride oil.





Antiperspirant Compositions

The present invention relates to antiperspirant compositions and particularly to aerosol antiperspirant compositions and to cosmetic methods of controlling perspiration from localised areas of the body, such as from the underarm.

Background and Prior Art

For many years, humans have employed cosmetic methods, sometimes alternatively referred to as non-therapeutic methods, to prevent or at least ameliorate bodily functions which local society at the time considers to be unsightly or otherwise undesirable. These methods have included controlling the appearance of sweat by topical application of an active ingredient which prevents egress of sweat from the eccrine glands. The active can be applied cosmetically and topically to the skin, broadly speaking, by one of two methods. Different consumers prefer one method or the other. In one method, sometimes called a contact method, a composition is wiped across the surface of the skin, depositing a fraction of the composition as it passes. In the second method, sometimes called the non-contact method, the composition is sprayed from a dispenser held proximate to the skin, often in the region of 10 to 20cms. The spray can be developed by mechanical means of generating pressure on the contents of the dispenser, such as a pump or a squeezable sidewall or by internally generated pressure arising from a fraction of a liquefied propellant volatilising, the dispenser commonly being called an aerosol.

25

20

5

10

15

In cosmetic aerosol compositions, the antiperspirant active is commonly an aluminium active. An aerosol dispenser has to withstand the internal pressure generated by the propellant so that it is usually made from metal. Aerosol compositions are commonly anhydrous, because the aluminium antiperspirant

- 2 -

active would dissolve in water-containing formulations generating an acidic solution that can cause or exacerbate corrosion of metal canisters.

5

10

15

20

25

30

Cosmetic products are employed by consumers in order to improve their appearance, and although for many years antiperspirant compositions in general and aluminium-containing antiperspirant aerosol compositions have been employed primarily to reduce visible wetness on various regions of the body, and especially in the underarm, some consumers are seeking even further benefits. A substantial number of female users of underarm antiperspirants shave their armpits, or pluck the hairs that grow there, an act which is considered by parts of society to improve their appearance. Shaving and plucking irritates human skin resulting often in localised reddening, which can manifest itself as blotchiness. The antiperspirant active itself can likewise irritate the skin by de-moisturisation in between the eccrine glands, and the two actions together combine to disfigure the armpit. Also, as has been long since recognised, an aluminium antiperspirant, and especially a particulate aluminium antiperspirant, tends to deposit a visible whiteness on the skin. Furthermore, as a person becomes older, skin progressively loses its inherent elasticity and becomes rougher and more wrinkled, even skin in the underarm that is typically often covered by clothing or least is usually out of the direct glare of harmful UV radiation. Moreover, the very materials that are necessary to inhibit localised perspiration themselves tend to de-moisturise the skin, reducing its elasticity and can even cause increased skin irritation, impairing the visual appearance of skin as well as its physical properties. Although a moisturising agent or agents has been incorporated in Dove antiperspirant compositions to counteract or ameliorate the impairment of skin elasticity, such as the incorporation of an occlusive agent, it has relatively little visual benefit.

To summarise, a need has been identified to counteract the impairment in visual appearance of underarm skin itself arising from performance of common acts to

- 3 -

enhance beauty, such as shaving together with applying an effective but nontherapeutic antiperspirant to counteract perspiration.

In the course of conducting investigations into how to improve the beauty of underarm skin whilst simultaneously applying a cosmetic antiperspirant composition, the team including the instant inventors recognised that the incorporation of a smoothness aid could assist, provided that it was chosen appropriately. A number of factors needed to be balanced or taken into account.

First, in order to remain useable as an aerosol, the viscosity of the composition had to be restrained to permit it to be sprayed through conventional dispensing apparatus. Secondly, the composition after its application and volatilisation of the propellant, should offer a pleasing appearance to the eye in such respects as tone, radiance and smoothness and not to exhibit excessive visible whiteness.

15

20

25

30

5

By way of reference, a commercial antiperspirant aerosol product (Dove) has been sold for several years which comprised a triglyceride emollient oil together with a moisturiser (an occlusive agent) in order to retain moisture and thereby enhance skin elasticity. Whilst this achieved its intended result, and helped to counteract shaving or plucking, it was comparatively ineffective at concealing skin imperfections, or improving the tone, radiance and smoothness of underarm skin. A number of materials were tried that had been suggested or incorporated in skincare products that were not anhydrous antiperspirant aerosols. Such materials included polymethylmethacrylate and silicone elastomers. Regrettably, they did not achieve the desired improvement in skin tone, radiance and/or smoothness when employed at a concentration that would have been desirable in aerosol compositions.

It is an object of the present invention to identify a smoothing aid for an aerosol antiperspirant composition that improves the beauty of underarm skin, or at last

- 4 -

ameliorates the effect of employing the antiperspirant itself or the effect of underarm challenges such as shaving or plucking of hair.

Brief summary of the present invention

5

10

15

20

25

According to one aspect of the present invention, there is provided an anhydrous antiperspirant composition comprising a base formulation and a propellant having a boiling point at standard pressure of below 10°C, which base formulation comprises a particulate astringent aluminium salt, an oil and optionally a particulate clay or silica suspending aid, and which further comprises a smoothing aid comprising a wax having a melting point of from 50°C to 90°C and optionally mica.

It has been found that by incorporating a wax in an aerosol composition, it is possible to improve the visible appearance of armpit skin, such as its tone, imperfections, smoothness and/or radiance. It is not necessary to employ a high concentration of the wax, and indeed it is possible to achieve a desired outcome at a concentration less than that at which the composition would be thickened to such an extent that could not be readily sprayed or would suffer a significant risk of blocking the spray outlet in use. Without being bound by any theory, hypothesis or conjecture, it is believed that when the composition containing the wax is sprayed onto human skin, the wax, or at least some of it, may form small globules that can act to refract or scatter light, thereby diffusing the appearance of the skin on which the wax has been deposited. By way of analogy, the diffusion can be compared with viewing an object through frosted glass – the object is visible but with a fuzzy outline.

In a second aspect of the invention, there is provided a means for enhancing the appearance of skin to which an astringent aluminium salt has been applied

WO 2010/057968

topically from an anhydrous aerosol composition as an antiperspirant which further comprises a selected smoothing aid.

In a third aspect of the present invention, there is provided a means for enhancing the appearance of skin to which an astringent aluminium salt has been applied topically from an anhydrous aerosol composition in which its base formulation contains an astringent aluminium salt that acts as an antiperspirant and a selected smoothing aid together with one or more moisturisers and/or a triglyceride oil.

In a further aspect of the present invention there is provided a method for mitigating the impaired visual appearance of skin, such as underarm skin, arising from topical application thereon of an aluminium-containing astringent antiperspirant salt by simultaneous spraying of a moisturiser and/or a triglyceride oil together with a wax and/or mica.

15

5

Other and further aspects of the present invention may be apparent from the subsequent text herein.

Detailed description of the present invention and preferred embodiments thereof.

20

The present invention in certain aspects contemplates incorporating a small concentration of wax in an anhydrous aerosol containing an astringent aluminium salt as an antiperspirant. The concentration of the wax is insufficient to solidify the base formulation.

25

The aerosol compositions herein comprise a base formulation which is mixed with a liquefiable propellant gas such that at the pressure in the dispenser, the propellant is a liquid or with a propellant that remains gaseous at elevated pressure.

- 6 -

A concentration of an ingredient given herein is based by weight on the base formulation, unless expressly specified otherwise.

The present invention seeks to ameliorate one or more visual impairments of underarm skin and/or improve positive attributes such as tone, smoothness and/or radiance, and thereby render the skin more attractive by creating a soft focus effect.

Herein, soft-focus effect of the film of antiperspirant composition on the skin applied by spraying is indicative of how strongly it scatters light. The soft-focus measurement is obtained by dividing the diffuse transmittance of the film by the total transmittance of a layer of that film.

Opacity is the degree to which the film of antiperspirant composition visibly covers
the skin upon which it has been applied. The opacity is determined by
measuring the diffuse reflectance of a film of similar thickness of the antiperspirant
composition on a black substrate and dividing it by the diffuse reflectance of the
film on a white substrate.

Gloss indicates shine in relation to the film. Herein, its gloss of the film is its reflectance measured at the specular angle of 60° from the normal to the film surface.

The instant invention in at least some preferred embodiments balances matt and gloss effects.

- 7 -

Base Formulation

5

10

15

20

25

Particulate Antiperspirant Active

Herein, the antiperspirant active is a particulate astringent aluminium salt and particularly a basic salt such as a basic aluminium salt. It is especially desirable to employ an aluminium chlorohydrate, by which is meant herein a material which satisfies the empirical formula $Al_2(OH)_xCl_y$ in which x + y = 6 and y is normally at least 0.5 and commonly not greater than 1.8, the material usually comprising bound water of hydration. The weight proportion of said water of hydration is conventionally not more than 12% and often lies in the range of from 3 to 10%. The term aluminium chlorohydrate herein encompasses materials with specified figures for x and y, such as aluminium sesquichlorohydrate and materials in which the chlorohydrate is present as a complex. It will be recognised that alternative names are sometimes used to indicate the presence of hydroxyl substitution, including aluminium hydroxychloride, aluminium oxychloride or basic aluminium chloride.

Aluminium chlorohydrate, as made, comprises a mixture of a number of different polymeric species in varying proportions, depending on the molar ratio of aluminium to chloride and the conditions employed during manufacture. All such mixtures are employable herein. It is especially desirable to employ what is commonly called activated aluminium chlorohydrate or enhanced activity aluminium chlorohydrate, sometimes abbreviated to AACH, in which the proportion of the more active species is higher by virtue of its method of manufacture. In one definition of activated, given in EP 6739, the material has greater than 20% Band III. Other methods of making AACH are given in EP 191628 and EP 451395. AACH is often made by recovery of an aluminium chlorohydrate from a dilute solution under strictly controlled

- 8 -

reaction/maturing/dewatering/drying conditions. AACH is commercially available by name from suppliers such as Reheis, and B K Giulini.

Aluminium chlorohydrate, whether it be activated or not activated, can be complexed, whereupon the CTFA name for the complex is concatenated to aluminium chlorhydrex, followed by the name of the molecule with which it is complexed. Commonly, such complexes include propylene glycol, that is representative of C_2 to C_6 glycols and glycine, that is representative of aminoacids. Any such complexed molecules remain complexed during formation of the invention base and aerosol compositions herein.

5

10

15

20

25

30

The particles of the astringent salt employed herein, such as an aluminium chlorohydrate, be it activated, complexed or otherwise, desirably have a diameter of below 125 microns, preferably \geq 99% by weight below 100 µm and especially \geq 95% by weight below 75 µm. Commonly, the astringent salt herein has a weight average mean particle size selected in the range of from 10 to 30 µm. In some embodiments, said mean is from 13 to 18 µm and in other embodiments, said mean is from 18 to 27µm. In a number of particularly preferred embodiments, the astringent salt is an activated aluminium chlorohydrate having said mean of from 14 to 26 µm.

Herein, unless otherwise stated, particle sizes (diameters) and distributions for the antiperspirant active are those that are obtainable by laser light scattering, for example obtained from the appropriate Mastersizer instrument for anhydrous suspensions, obtainable from Malvern Instruments set to produce a volume plot. The instrument is employed with a lens selected in accordance with the maker's instructions to accommodate the expected particle size distribution, (or various lenses can be tested until the best lens is identified) and is preferably operated employing cyclomethicone (DC245TM from Dow Corning) as the liquid dispersant for a sample of the base formulation to attain a particles concentration that

- 9 -

achieves obscuration, i.e. 10-30% light scattered. Using the Polydisperse analysis model and knowing the dispersant RI, the RI of the particulate material and imaginary RI factor of 0.1, the plot of the particles size (d) distribution and the average particle size D50 is obtained.

5

10

The weight proportion of antiperspirant salt, e.g. aluminium chlorohydrate of AACH in the base formulation is commonly selected in the range of from 5 to 60%, often not greater than 50% and in many suitable embodiments from 30 to 45% by weight, the weight including any water of hydration and complexed molecules. The proportion of such active in the base formulation is often selected in conjunction with the weight ratio of propellant to base formulation such that the weight proportion of aluminium salt in the full aerosol composition is from 1 to 15%, and particularly from 2 to 12 wt%.

15 **Oil**

The base formulation comprises an oil or preferably a mixture of oils in which particulate materials are suspended, so that the aerosol expels a pattern of liquid droplets when sprayed. The term oil herein indicates a hydrophobic material that is liquid at 20°C at 1 atmosphere pressure. It will be recognised that various of the oils contemplated herein can provide one or more functions in addition to acting as a carrier for particulate materials, for example some act as an emollient, or mask skin deposits to alter the appearance of the aerosol composition when applied topically, and/or some can mask odour of the composition itself or malodours generated on skin from secretions.

25

30

20

The proportion carrier fluids in the base formulation, including oils and optional or other functional ingredients which are liquid at 20°C and oil-miscible, is often at least 35%, and in many base formulations at least 45%, is commonly not higher than 85%, in many embodiments not higher than 75% and in several practical formulations not higher than 65%, %s being by weight of the base formulation. In

- 10 -

a number of highly desirable base formulations, the oil weight proportion is from 45 to 85%, particularly 45 to 70%, and in certain desirable base formulations is from 50 to 60%. The term carrier fluid is employed in regard to the base formulation and therefore excludes propellant which is subsequently mixed with the base formulation to create the aerosol composition.

The oils employed in the instant invention often desirably comprise one or more volatile silicone oils. By volatile herein is meant having a measurable vapour pressure at 20 or 25^oC of at least 1Pa. Typically the vapour pressure of a volatile silicone oil lies in a range from 1 or 10 Pa to 2 kPa at 25°C. Volatile silicone oils can be linear or cyclic siloxanes, usually containing from 3 to 9 silicon atoms, and commonly from 4 to 6 silicon atoms, the silicon atoms being substituted by methyl groups, so that their alternative names are methicones and cyclomethicones. It is especially desirable to employ volatile silicone oils in which at least 80% by weight and particularly at least 90% contain at least 5 silicon atoms, such as cyclopentadimethylsiloxane (D5), cyclohexadimethylsiloxane (D6), dodecamethylpentasiloxane and tetradecamethylhexasiloxane. The cylomethicone oils are especially preferred. Such oils are highly desirable for many consumers because they can evaporate without causing undue skin cooling. The volatile silicone oils often comprise at least 30% by weight of the carriers fluids and normally not higher than 95% thereof, and in a number of desirable formulations the weight proportion of the carrier fluids is at least 35%, and particularly at least 40% and in the same or other embodiments of the invention is up to 75%, especially up to 65% and particularly up to 55%.

25

30

5

10

15

20

The oils in the carrier fluid preferably comprise one or more non-volatile oils, which can be silicone oils and/or particularly non-silicone oils. Preferably, in some embodiments at least a fraction of the non-volatile oils has a refractive index of at least 1.45. Such oils in the base formulation can advantageously lessen the appearance of visible residues on skin, not only immediately on application but

- 11 -

also throughout the period typically from 6 to 24 hours before the antiperspirant formulation is washed off.

5

10

25

Non-volatile silicone oils employed herein preferably contain one or more unsaturated substituents such as phenyl or diphenylethyl in replacement of the corresponding number of methyl substituents in polycyclosiloxanes or more preferably in linear siloxanes, often having 2 or 3 silicon atoms. Such non-volatile oils have a higher refractive index than that of the volatile silicone oils and tend to mask the antiperspirant active when it is deposited on skin. The non-volatile oils can also comprise dimethiconols, which as the name suggests are hydroxylterminated. The proportion of non-volatile silicone oils in the carrier fluids is conveniently from 0 or 0.25 to 10% by weight and often from 0.5 to 5% by weight, such as in the range of from 1 to 3% by weight of the oils.

The oils in the carrier fluid can alternatively or additionally comprise one or more hydrocarbon fluids (oils), which can be either volatile or non-volatile. Suitable hydrocarbon fluids include liquid aliphatic hydrocarbons such as mineral oils or hydrogenated polyisobutene, desirably selected to exhibit a low viscosity. Further examples of liquid hydrocarbons are polydecene and paraffins and isoparaffins of at least 10 carbon atoms. Hydrocarbon fluids conveniently comprise from 0 to 25% by weight of the carrier.

In at least some advantageous embodiments, the carrier oils comprise liquid aliphatic or aromatic ester oils. Suitable aliphatic esters contain at least one long chain alkyl group, such as esters derived from C_1 to C_{20} alkanols esterified with a C_8 to C_{22} alkanoic acid or C_6 to C_{10} alkanedioic acid. The alkanol and acid moieties or mixtures thereof are preferably selected such that they each have a melting point of below $20^{\circ}C$. Aliphatic esters include isopropyl myristate, lauryl myristate, isopropyl palmitate, diisopropyl sebacate and diisopropyl adipate.

Further and very suitable ester oils include glyceride oils and in particular

- 12 -

triglyceride oils derived from glycerol and fatty acids containing at least 6 carbons, and especially natural oils. The weight proportion of aliphatic esters in the base formulation is often up to 10%, such as at least 1%.

It is particularly desirable to include a liquid aromatic ester include fatty alkyl benzoates. Examples of such esters include suitable C₈ to C₁₈ alkyl benzoates or mixtures thereof, including in particular C₁₂ to C₁₅ alkyl benzoates e.g. those available under the trademark Finsolv. An aryl benzoate, such as benzyl benzoate can also be used. Yet other suitable ester oils includes oils in which a short alkylene group of 1 to 3 carbons, optionally substituted by a methyl group, is interposed between benzene and benzoate residues. The weight proportion of such aromatic ester oils is often from 2.5 to 10% in the base formulation and particularly from 3.5 to 7%.

15 The total proportion of ester oils, including both aliphatic and aromatic ester oils (but excluding any fragrance oil) is often from 0 to 50% by weight of the oil mixture, is desirably at least 2% and especially is at least 5% by weight. Their total proportion is in many embodiments desirably up to 30% and particularly attractively up to 20% by weight of the oil mixture. In some especially desirable formulations, the ester oils constitute from 7.5 to 20% of the oils mixture. Expressed on the basis of the base formulation, the ester oils often constitute from 3.5 to 16% by weight and particularly from 4.5 to 13% by weight. The weight ratio of aromatic ester oil to aliphatic ester oil is often selected in the range of from 1:3 to 10:1 and especially from 1:2 to 2:1.

25

30

Natural oils which are most desirably employed herein comprise triglyceride oils such as in particular one or more unsaturated C₁₈ fatty acid glycerides. In many instances, the oils comprise one or more triglycerides. The fatty acid residues in the oils can comprise, commonly, from one to three olefinic unsaturated bonds and often one or two. Whilst in many instances the olefinic bonds adopt the trans

- 13 -

configuration, in a number of desirable products the bond or bonds adopt the cis configuration. If two or three olefinic unsaturated bonds are present, they can be conjugated. The fatty acid can also be substituted by an hydroxyl group. The natural oils employable herein desirably comprise one or more triglycerides of oleic acid, linoleic acid, conjugated linoleic acid, linolenic acid or ricinoleic acid. Various isomers of such acids often have common names, including linolenelaidic acid, trans 7-octadecenoic acid, parinaric acid, pinolenic acid punicic acid, petroselenic acid and stearidonic acid. It is especially desirable to employ glycerides derived from oleic acid, linoleic acid or petroselenic acid, or a mixture containing one or more of them.

5

10

15

20

Natural oils containing one or more of such triglycerides include coriander seed oil for derivatives of petroselinic acid, impatiens balsimina seed oil, parinarium laurinarium kernel fat or sabastiana brasilinensis seed oil for derivatives of cisparinaric acid, dehydrated castor seed oil, for derivatives of conjugated linoleic acids, borage seed oil and evening primrose oil for derivatives of linoleic and linolenic acids, aquilegia vulgaris oil for columbinic acid and sunflower oil, olive oil or safflower oil for derivatives of oleic acid, often together with linoleic acids. Other suitable oils are obtainable from hemp, which can be processed to derive stearadonic acid derivatives and maize corn oil. An especially convenient natural oil by virtue of its characteristics and availability comprises sunflower oil, ranging from those rich in oleic acid glycerides to those rich in linoleic acid glycerides, rich indicating that its content is higher than that of the other named acid.

The proportion of the natural oil, viz particularly the triglyceride oils of unsaturated fatty acids, in the formulation is often selected in the range of from 0.1 to 15% by weight of the oils mixture, desirably at least 0.25% by weight, especially in the range of from at least 0.5 of the oils mixture.

- 14 -

Expressed as a proportion of the base formulation, in some beneficial embodiments, the triglyceride oil represents from 1 to 8% by weight, particularly from 2 to 6% and advantageously from 3 to 5% by weight thereof.

Additionally or alternatively, the base formulation can comprise jojoba oil such as in a proportion of up to 5% w/w of the base formulation.

10

15

20

30

A further class of particularly suitable carrier oils comprise non-volatile liquid aliphatic ethers derived from at least one fatty alcohol that desirably contains at least 10 carbon atoms, such as myristyl ether derivatives e.g. PPG-3 myristyl ether or lower alkyl ($\leq C_6$) ethers of polygylcols (preferably polypropylene glycol and especially 10 to 20 units, such as an ether named as PPG-14 butyl ether in the CTFA. It is often convenient for the aliphatic ether to constitute at least 10%, and especially at least 15%, particularly up to 60% and very desirably up to 50% by weight of the oils mixture, including the range of from 30 to 45%. Expressed as a weight proportion of the base formulation, the ether oil constitute desirably from 15 to 30% and particularly from 20 to 25% by weight.

Such ethers, and especially those having a refractive index of above 1.46 can assist in masking the visibility of deposits on the skin, thereby complementing the positive skin conditioning properties of the overall formulation. It can be very desirable to select the ether in a weight ratio to the antiperspirant active of at least 0.3:1 such as up to 0.8:1, one practical range being from 0.5:1 to 0.7:1.

25 It is often convenient to select the aromatic ester oil plus non-volatile aliphatic ether to volatile silicone oil in a weight ratio of from 3:2 to 3:1, and especially from 1.75:1 to 2.5:1.

A further class of oils can be employed beneficially herein comprises waterimmiscible aliphatic alcohols, and particularly those having a boiling point of

higher than 100°C. These include branched chain alcohols of at least 10 carbon atoms and in many instances up to 30 carbon atoms, particularly 15 to 25, such as isostearyl alcohol, hexyl-decanol and octyl-dodecanol. It will be recognised that octyl dodecanol is an especially favoured water-immiscible aliphatic alcohol in the instant formulations, because it not only acts as an emollient oil but additionally moisturises skin by the mechanism of occlusion. Such alcohols can often constitute at least 0.1% and particularly at least 0.5% by weight of the oils mixture, and in many formulations are not more than 5% and particularly up to 3% by weight of the oils mixture. Expressed as a weight proportion of the base formulation, the branched alcohol desirable comprises from 0.25 to 2% by weight thereof, and particularly from 0.5 to 1.5% thereof. In some especially desirable embodiments, the branched fatty alcohol is present in a weight ratio to the smoothing aids of from 1:2 to 2:1.

In the invention base formulations, the total weight proportion of non-volatile oils is often falling in the range of from 30 to 50% and particularly from 35 to 45%. The total weight proportion of volatile oils in the invention base formulation is often chosen in the range of from 12.5 to 25%, and in many very suitable embodiments from 15 to 22.5%. The weight ratio of non-volatile to volatile oils is desirably chosen in the range of from 2:1 to 4:1, particularly from 3:2 to 5:2 and especially from 7:4 to 9:4.

Smoothing Aid

5

10

25

30

The base formulation of the aerosol compositions disclosed herein contains at least one smoothing aid, of which two classes are disclosed in more detail. One class comprises a wax and the other class comprises mica. Again without being bound by theory, mica is highly desirable because it can be considered to adopt the form of platelets. As such, it is a preferred representative of a broader class of minerals comprising alumino-silicate minerals that likewise adopt the form of platelets. It is particularly desirable to employ at least one member from both

- 16 -

classes, in that they complement each other. One tends to provide a matt finish, whereas the other increases reflectance, so that the two together enhance the soft-focus appearance of skin onto which has been sprayed the thin film of antiperspirant composition.

5

20

30

Without being bound by theory as to why the soft focus of the skin is improved, incorporation of a limited concentration of wax is observed to offer a matt finish to the antiperspirant film. The waxes that can be contemplated for employment herein normally have a melting point of up to 90°C and often up to 80°C.

Typically, desirable waxes have a melting point of at least 50°C, and especially at least 55°C. A number of highly desirable waxes melt in the range of up to about 70°C. The waxes are selected from:-

hydrocarbon waxes, such as mineral waxes or polyethylene, commonly having an average molecular weight of from 250 to 600;

aliphatic ester waxes, such as esters derived from either or both of a fatty acid or alcohol, the resultant ester often containing from 18 to 40 carbon atoms; natural waxes derived from plants or animals, such as beeswax, candelilla wax and caster wax

and, preferably, linear fatty alcohol waxes, commonly containing from 14 to 22 carbon atoms, such as cetyl alcohol, stearyl alcohol, or eicosyl alcohol.

The waxes can be natural or synthesised. Mixtures of two or more waxes can be employed, including mixtures from different subclasses.

It is particularly desirable to select stearyl alcohol or a mixture of waxes comprising stearyl alcohol, such as cetearyl alcohol.

When selecting the wax or waxes to employ, it is preferred to employ a wax that is amorphous rather than crystalline, and particularly effective results have been achieved from a wax that is not only amorphous, but also has a comparatively low melting point, such as in the range of from 55 to 65°C.

- 17 -

A low concentration of wax, and especially an amorphous wax, in the base formulation is deliberately chosen. It will be recognised that to at least some extent, during processing of the base formulation and the aerosol composition, intensive mixing operations can result in localised heat generation, leading to dissolution at least a fraction of the wax into the carrier oil (or blend of oil with the propellant, as the case may be). Localised melting of the wax can possibly also contemplate wax droplet formation.

5

20

25

As the concentration of wax increases, there is a tendency for the resultant aerosol composition to exhibit a higher viscosity, making it more difficult for the composition to be sprayed from the aerosol container and increasing the risk of blockage of the spray outlet (the spray nozzle) and/or the outlet control valve. Viscosity increase can also be compounded by the risk of particulates agglomeration as the proportion of wax increases in the base formulation.

Thus, in practice, there is a concentration window for the wax that takes into account both the desirability of providing a smoothing aid and the desirability of not increasing the viscosity of the composition too much. Desirably, the wax is present in the base formulation at a concentration of up to 2.5% by weight, often at up to 2% and in many preferred embodiments at up to 1.5% by weight. Advantageously, the wax is present at a concentration of at least 0.1% by weight of the base formulation, particularly at least 0.2% and especially at least 0.4% by weight. In the invention compositions contemplated herein, the weight ratio of total liquid to wax in the aerosol formulation, i.e. both liquids in the base formulation plus the propellant, is desirably greater than 140:1, and in many practical embodiments is greater than 200:1 such as in the range of from 250:1 up to 2000:1, and particularly from 500:1 to 1250:1.

- 18 -

The above limits for the concentration of the wax and its ratio to liquids in the base formulation are particularly suitable when the proportion of non-volatile oil in the base formulation comprises from 35 to 70% of the total of liquids in the base formulation.

5

10

15

The second class of smoothing aids contemplated herein comprises in particular mica, although may alternatively be considered to include other alumino-silicate minerals that are deemed to adopt the form of platelets. The mica desirably has a weight-averaged mean particle size of below 50µm and especially up to 30 µm. If desired more than one mica material can be employed. For practical reasons, the weight-averaged mean particle size of at least one mica material employed is preferably at least 7µm and in some advantageous embodiments is at least 15 µm. One convenient combination of mica materials comprises one material with -averaged mean particle size of 7 to 14 microns and a second material with weight-averaged mean particle size of 18 to 25 microns. In many desirable embodiments herein, the weight average mean particle size of the mica and the aluminium antiperspirant salt are both selected in the range of from 7 to 30 microns and are present in a ratio of from 2:1 to 1:2.

In some or other embodiments, herein, the mica can be employed without further treatment. In other and much favoured embodiments, the mica has been surface treated with titanium dioxide, and/or tin oxide and/or silicon dioxide, thereby forming a mica pigment. In particularly favoured embodiments of the present invention, the mica surface treatment further incorporates a transition metal oxide imparting a colour, so that the resultant material is an interference pigment. Suitable transition metal oxides include iron and chromium oxide. By the choice of the complementary oxide, the producer of the pigment is able to create pigments exhibiting the full visible spectrum of colour highlights, as well as silver. It is especially suitable to select mica interference pigments having blue, green or violet highlights, i.e. a wavelength of less than about 550nm, and particularly less

- 19 -

than 500nm. The interference pigment constitutes a fraction of the mica pigment, such as from 20 to 75% by weight.

In some embodiments of the present invention, the mica, including a mica pigment, is employed in conjunction with the above-mentioned wax. In other embodiments, which can provide at last some of the benefit arising from use of a combination of wax plus mica, the smoothing aid is a mica pigment exhibiting highlights of wavelength less than about 550 nm and particularly less than 500 nm.

10

5

The weight proportion of mica in the invention compositions herein is desirably at least 0.1% of the base formulation and preferably at least 0.25%. Its weight proportion is normally not greater than 5% and commonly not greater than 2%. When employed together with a wax, the weight ratio is often from 3:1 to 1:3, and in many practical embodiments from 2:1 to 1:2.

15

In a number of embodiments in accordance with the third aspect of the present invention, there is provided an anhydrous aerosol antiperspirant composition in which the base formulation comprises the smoothing aid comprising mica or/and wax in a concentration of at least 0.25%, together with a moisturiser at a concentration of at least 0.25% by weight and/or a triglyceride oil at a concentration of at least 1% by weight. Preferred smoothing aids, antiperspirant actives and triglyceride oils, and combinations thereof, as well as preferred amounts are described herein in greater detail.

25

30

20

The moisturiser for employment in conjunction with the third aspect can suitably be an occlusive agent, such as an occlusive oil, e.g. octyldodecanol. It will be recognised that as a general rule, most water-immiscible ester oils, such as isopropyl myristate or alkyl C_{12-15} benzoate act as an emollient oil, but do not act as an occlusive agent. The moisturising agent in the third aspect can alternatively

comprise polyethylene glycol (PEG) desirably having an average molecular weight of from 200 to 625 and particularly from 250 to 450, advantageously in a concentration of from 1 to 5% by weight of the base formulation.

5

10

15

20

In addition to the antiperspirant salt, the oil and the smoothing aid or aids, the base formulation preferably contains a particulate suspending aid. Suitable suspending aids include colloidal silicas, suitably pyrogenic silica, and clays such as montmorillonite clays such as bentonite and hectorite. Particularly desirably, the suspending aids have a hydrophobically treated surface. A particularly preferred bentonite is hydrophobic bentonite, e.g. aids which are commercially available under the trade mark Bentone, eg Bentone LT, Bentone 14, Bentone 27, Bentone 34, and Bentone 38 / 38V and is a bentonite treated with hydrophobic cationic materials. Other suitable clay suspending aids include colloidal magnesium aluminum silicates. Advantageously, the suspending aid is utilised at a level of from at least 1%, and preferably at least 2% by weight of the base formulation, often at up to 8% and in some desirable embodiments at up to 6%. It is often beneficial to calculate the proportion of suspending aid by reference to the antiperspirant active, and in particular in a weight ratio of antiperspirant active: suspending aid of up to 15:1. Said ratio is greater than 4:1 in many embodiments, and herein a particularly desirable ratio range is from 6:1 to 12:1.

Herein, it is advantageous to employ an activator for the clay suspending aid, for example to encourage variations in the manufacturing process from the base formulation. Clay activators include ethanol and especially propylene carbonate.

The amount of activator is preferably selected in the range of from 25 to 75 wt% of the weight of the suspending aid, is advantageously at least 35% and is often not more than 60%. Accordingly, herein, in some advantageous compositions, the base formulation comprises from 1 to 3% by weight propylene carbonate. It has been observed that the addition of the activator can further enhance the ability of the clay-suspending aid-containing composition after topical application to offer

- 21 -

observed smoothness and/or cover imperfections and/or improve the tone and/or the radiance of the skin.

Accordingly, in a further aspect of the present invention, the base formulation comprises in addition to the particulate aluminium antiperspirant salt, the oil and the clay suspending aid, (as described herein), a smoothing aid selected from the wax and/or the mica as described herein together with the propylene carbonate.

The base formulation can comprise one or more optional constituents which have hitherto been incorporated or proposed for incorporation in anhydrous aerosol antiperspirant compositions. Such optional constituents may be liquid (in which event they are desirably miscible with the oils therein) or solid, and normally comprise in total not more than 10% and often not more than 5% by weight of the base formulation. Such optional constituents can comprise non-antiperspirant deodorant actives, such as antimicrobial actives such as polyhexamethylene biguanides, e.g. those available under the trade name Cosmocil™ or chlorinated aromatics, e.g. triclosan available under the trade name Irgasan™, nonmicrobiocidal deodorant actives such as triethylcitrate, bactericides and bacteriostats. Yet other deodorant actives can include zinc salts such as zinc ricinoleate. The compositions can additionally or alternatively contain, as a bacteristat, an iron chelator such as pentenoic acid which hinders bacterial growth/reproduction. The proportion of the deodorant active in the base formulation is often selected in the range of from about 0.05 to 2% by weight of the base formulation and especially from 0.1 to 0.5%.

25

30

20

5

10

15

Other optional ingredients can include sensory modifiers, such as talc or finely divided high molecular weight insoluble polyethylene, such as in an amount of up to 3% by weight; skin cooling agents such as menthol often selected in an amount of up 0.5%, particularly up to 0.2% of the base formulation and wash-off agents such as non-ionic surfactants, and particularly polyethoxylated fatty alcohols or

- 22 -

acids, for example in an amount of up to about 3% of the base formulation. A further optional ingredient comprises a hair growth inhibitor, for example palmatine, such as at a concentration in the base formulation of up to 0.2% by weight, e.g. at least 0.01%.

5

10

20

25

The base formulation desirably contains a fragrance, typically selected for its aesthetic appeal, such as at a concentration in the base formulation of from 0.1 to 8% by weight, and often at least 1% such as from 2 to 6% by weight therein. A fraction of the fragrance, such as in the range of from 1 to 100%, particularly 25 to 75% can be encapsulated, for example in modified starch, or can be absorbed on and releasable from a cyclodextrin and/or can comprise a pro-fragrance.

Aerosol compositions

15 **Propellant**

The anhydrous aerosol composition herein comprises a propellant in addition to the base formulation described hereinabove. Commonly, the propellant is employed in a weight ratio to the base formulation of from 95:5 to 5:95, the weight proportion chosen depending in practice on the choice of propellants, the internal pressure which the manufacturer seeks to generate and the architecture of the aerosol canister. Depending on the propellant, in such aerosol compositions the ratio of propellant to base formulation is normally at least 20:80, generally at least 30:70, particularly at least 40:60, and in many formulations, the weight ratio is from 90:10 to 50:50. A ratio range of from 70:30 to 90:10 is sometimes preferred. Particularly suitable weight ratios of propellant:base formulation are in the regions of 3:1, 4:1 or 7:1, for example the propellant proportion of the total composition being within +/- 1% or 1.5% of 75%, 80% or 87% respectively.

- 23 -

Propellants herein generally accord with one of three classes; i) low boiling point gasses liquifided by compression, ii) volatile ethers and iii) compressed non-oxidising gases.

Class i) is conveniently a low boiling point material, typically boiling below -5°C. 5 and often below -15°C, and in particular, alkanes and/or halogenated hydrocarbons. This class of propellant is usually liquefied at the pressure in the aerosol canister and evaporates to generate the pressure to expel the composition out of the canister. Examples of suitable alkanes include particularly 10 propane, butane or isobutane, often in varying admixtures of the three components, and possibly containing a fraction of pentane or isopentane. Examples of halogenated hydrocarbons are fluorocarbons and chlorofluorocarbons such as, for example, 1,1-difluoroethane, 1-trifluoro-2fluoroethane, dichlorodifluoromethane, 1-chloro-1,1-difluoroethane, and 1,1-15 dichloro-1,1,2,2-tetrafluoroethane. It is particularly desirable to employ the first class of propellant. Such propellants are commonly employed at a weight ratio to the base formulation of from 1:2 and especially at least 1:1 up to 95:5.

The second class of propellant comprises a very volatile ether of which the most widely employed ether hitherto is dimethyl ether. This propellant can advantageously be employed at relatively low weight ratio of propellant to base formulation, for example to as low as 5:95. It can also be employed in admixture with, for example, compressible/liquefiable alkane gasses.

The third class of propellant comprises compressed non-oxidising gasses, and in particular carbon dioxide or nitrogen. Inert gases like neon are a theoretical alternative.

- 24 -

Aerosol Product Manufacture

An aerosol product according to the present invention can be made conveniently by

- first blending together the ingredients of the base formulation in a vessel, less any ingredient intended to be added later, agitating the mixture to suspend the particulate antiperspirant active,
 - charging an aerosol canister with the mixed base formulation and separately charging any other ingredient not blended previously into the base formulation,
- charging either before, after or simultaneously,
 - fitting and sealing the discharge line containing the valve onto the aerosol canister and
 - injecting propellant gas into the canister through the discharge line.
- By employing suitably intense mixing apparatus, such as by the use of a Sonolator or a Silverson mixer, at least some localised melting and/or dissolution of the wax ingredient can be effected, and particularly in respect of a wax having a comparatively low melting point such as up to 70°C.
- The selection of the aerosol canister is at the discretion of the manufacturer of the aerosol product. Conveniently, the canister itself can be made from tin plate or aluminium. The discharge line includes a valve biased to the closed position, and may be depressed or tilt valve. The valve can be opened by depression or lateral movement as is determined by the valve of an actuator.

25

30

The discharge line terminates in a spray outlet, typically comprising a nozzle which in conventional aerosol dispensers is moulded with the spray nozzle to form a depressor button or which in more recent developments such as in EP 1040055 or EP 1255682 comprises rotational or lateral relative motion of elements to prevent the valve being opened prematurely. The nozzle outlet has an internal

- 25 -

diameter that is usually selected within the range of from 300 to 800 microns, particularly not greater than 600 microns and in many embodiments from 350 to 450 microns. This is particularly beneficial in that it enables the composition to be sprayed without an undue risk of blockage through nozzles of similar internal diameter currently employable with corresponding otherwise similar compositions that lack the smoothing aid. This of course has the added advantage of allowing the canister to deliver a similar spray pattern, if desired, at a similar pressure to that of the smoothing aid-free composition.

According to a further aspect of the present invention there is provided an aerosol product comprising a base formulation substantially as described hereinbefore containing a smoothing aid selected from wax and/or mica, and a particulate antiperspirant active having >99% by weight of particles <75 microns and optionally containing a moisturiser and/or a triglyceride oil, disposed within an aerosol canister having a nozzle outlet of from 350 to 450 microns.

Topical application

5

20

25

30

The aerosol composition of the instant invention can be sprayed onto skin, and particularly into the underarm (axilla) in a conventional manner for spraying liquid compositions. Very conveniently, the base formulation can be mixed with a propellant prior to being sprayed from an aerosol canister. The canister is desirably held at a distance of between 12 and 18 cms from the armpit and the valve in the discharge line opened. The composition can be sprayed at the discretion of the user for a conventional period of time, such as from at least about 1 or 1.5 seconds and preferably from about 2 to 5 seconds per armpit. The spray deposits the droplets of mainly base formulation onto the skin surface, which tend to spread and coalesce, forming a film, from which the volatile oils evaporate. When present, globules, such as globules of the wax, and particulates, such as the mica, and especially the mica pigment, including the mica interference

- 26 -

pigment are exposed through the film and create the desired beneficial physical appearance.

Herein, except where the context demands otherwise, a numerical quantity, such as boundary or a ratio, can be qualified by the term "about".

Having described the invention, as well as preferred embodiments thereof, in general terms, particular embodiments thereof will described in more detail hereinafter by way of example only.

10

15

20

25

5

Comparison CA and Examples 1 to 4

In this Comparison and Examples, a base formulation as prepared having the composition summarised in Table 1 below by charging a vessel with the oils, and thereafter slowly introducing the particulate materials with energetic propeller mixing employing a Silverson mixer, including any smoothing aid (wax flakes and/or mica). The mixer blades tended to promote localised heating resulting in at least partial melting of the wax. When the formulation was visually homogenous, the fragrance oil was blended in and the entire composition was charged into metal 150g canisters. A discharge line was inserted through the mouth of the canister and a valve embedded in the line was mounted in a valve cup crimped over the open mouth of the canister. Thereafter, the canister was charged with the propellant through the discharge line, in a weight ratio of 13 parts base formulation to 87 parts propellant. The propellant comprised a mixture of propane, butane and isobutane (CAP 40 ex Calor). An actuator was fitted on to the canister having a spray outlet orifice (nozzle orifice) with an internal diameter of 400 microns.

- 27 -

Table 1

Comparison/Examples	CA	Ex 1	Ex 2	Ex 3	Ex 4
INCI name		%	by weigl	nt	•
Cyclomethicone ¹	18.52	15.96	15.24	15.24	15.36
Polydimethiconol in	1.54	1.54	1.54	1.54	1.54
cyclopentasiloxane ²					
PPG-14 Butyl Ether ³	22.24	22.24	22.24	22.24	22.24
Octyldodecanol ⁴	0.90	0.90	0.90	0.90	0.90
C12-15 Alkyl Benzoate ⁵	3.85	6.15	6.15	6.15	6.15
Helianthus Annuus Seed Oil 6	4.00	4.00	4.00	4.00	4.00
Disteardimonium Hectorite ⁷	4.23	3.85	3.85	3.85	3.85
ButylHydroxyToluene	0.77	0.77	0.77	0.77	0.77
Fragrance	5.38	5.38	5.38	5.38	5.38
Propylene Carbonate	0.12	0.00	0.12	0.12	0.00
Stearyl Alcohol	0.00	0.00	0.60	0.60	0.60
Titanium Dioxide & mica 8	0.00	0.37	0.37	0.37	0.37
Titanium Dioxide & mica 9	0.00	0.37	0.00	0.37	0.37
Titanium Dioxide & mica 10	0.00	0.00	0.37	0.00	0.00
Activated Aluminium Chlorohydrate Powder 11	38.46	38.46	38.46	38.46	38.46

The ingredients were as follows:-

	1	DC245	Dow Corning
5	2	DC1501	Dow Corning
	3	Fluid AP	Amerchol
	4	Eutanol G	Henkel
	5	Finsolv TN	Fintex
	6	AgriPure 80	Cargill PLC
10	7	Bentone 38V	Elementis

- 28 -

8	Timiron MP-115 Ultra-luster	Merck
9	Timiron Super Blue	Merck
10	Timiron Silk Blue	Merck
11	Summit 7172	Giulini

5

10

15

20

The various compositions were tested in the following manner and the results summarised in Table 2 below:-

- For each pair of compositions, 10 Female panellists aged between 18 and 66 and in good health who did not have a history of skin disease and were not undergoing treatment for a skin condition, shaved and washed their own axillae approx 12hrs before the study commenced and refrained from applying antiperspirant or other cosmetic composition to the axillae afterwards. Their underarms were assessed by an expert clinician in a well lit booth of a clinical test centre immediately before topical application of the test compositions, randomly one of the pair to the left and the other to the right axilla;
- Skin quality was scored on a 0 to 5 scale for tone, imperfections, radiance, visual smoothness and visual and the result recorded;
- Test product (about 250-300mg) was applied to each axillae by the assessor with a 2 second spray in a ventilated booth;
- The Expert clinician assessment reassessed the skin quality attributes directly after product application on the same 0 to 5 scales in the same booth and the results recorded;
- The difference between the score before and after application of the test product for each attribute was calculated and the average from the panellists is summarised in Table 2 below.

25

30

Increases in tone, radiance and smoothness are recorded as positive as are decreases in imperfections, if recorded. Accordingly, the data summarised in Table 2 indicates the effectiveness of the Example compositions at improving the

- 29 -

skin appearance and hence the perceived skin quality whilst applying a previously proven antiperspirant active.

The attributes were assessed by mapping onto the following scale (description of severity, coverage of axillae):-

0 none

5

- 1 light, up to 10%
- 2 mild, 11 to 25%
- 3 moderate, 26 to 50%
- 10 4 marked, 51 to 75%
 - 5 severe, 76 to 100%

The assessment of the attributes was developed in conjunction with the test subjects and was made by considering inter-alia the following skin features:-

Tone unevenness, freckles, Hyperpigmentation through to

evenness

Radiance dullness through healthy glow to brightness

Smoothness visible roughness, pits to velvety, peachy appearance.

- 30 -

Table 2

Comparison/Examples	CA	Ex 1	Ex 2	Ex 3	Ex 4
Tone	0.2	0.3	0.6	0.9	0.6
Radiance	0.35	0.6	0.6	0.8	0.9
Smoothness	0.3	0.4	0.6	0.9	1.0

From the data summarised in Table 2, it can be seen that the incorporation of at least one smoothing aid improved the perceived quality of the underarm skin 5 compared with a comparison aerosol product that itself had previously been demonstrated to exhibit improved moisture retention in the skin relative its predecessor, but without offering significant perceivable improvement in underarm skin appearance (Comparison A). In particular, Example 1 shows that incorporation of even a low concentration of two mica pigments, including in particular an interference pigment exhibiting blue highlights improved not only the 10 tone of the skin, but showed a greater improvement for both radiance and smoothness, indicating to the user a healthier and younger-looking skin. Examples 2, 3 and 4 show not only that incorporation of a wax improved the perception of the skin, but that the presence of both the wax and the mica pigment 15 offered a greater benefit than the mica pigment alone.

Claims:

- 1. An anhydrous antiperspirant composition comprising a base formulation and a propellant having a boiling point at standard pressure of below 10°C, which formulation comprises a particulate astringent aluminium salt, an oil and optionally a particulate clay or silica suspending aid, and which further comprises a smoothing aid comprising a wax having a melting point of from 50°C to 90°C and optionally mica.
- 2. A composition according to claim 1 in which the wax has a melting point in the range of from 55 to 90°C.
- 3. A composition according to claim 1 or claim 2 in which the wax has a melting point in the range of from 58 to 70°C.
- 4. A composition according to any one of the preceding claims in which the wax is amorphous.
- 5. A composition according to any of the preceding claims in which the wax comprises a linear fatty alcohol.
- 6. A composition according to claim 5 in which the wax comprises stearyl alcohol.
- 7. A composition according to any preceding claim in which the base formulation contains from 0.2 to 2.5% by weight of wax.
- 8. A composition according to claim 7 in which the base formulation contains from 0.4 to 1.5% by weight of the wax.

- 9. A composition according to any one of the preceding claims in which the base formulation contains at least 0.1% by weight of the mica.
- 10. A composition according to any preceding claim in which the base formulation contains up to 5% by weight of the mica.
- 11. A composition according to claim 10 comprising from 0.25% up to 2% by weight of the mica.
- 12. A composition according to any preceding claim in which the mica further comprises a coating with titanium dioxide tin oxide and/or silicon dioxide forming a mica pigment.
- 13. A composition according to claim 12 in which the coating further comprises a transition metal oxide or a bismuth salt.
- 14. A composition according to claim 13 in which the transition metal oxide is selected from one or more of iron and chromium
- 15. A composition according to any preceding claim in which the mica has a weight average mean particle size in the range of from 7 to 30 μm.
- 16. A composition according to claim 15 in which the mica has a weight average mean particle size in the range of from 18 to 25 μm.
- 17. A composition according to claim 15 or claim 16 in which the mica is a mixture of two materials, both materials having a weight average mean particle size in the range of from 18 to 25 μm, and at least one of which is a mica pigment.

- 18. A composition according to any one of claims 15 to 17 in which the mica is a mixture of two materials, one material having a weight average mean particle size in the range of from 18 to 25 μm and the other material having a weight average mean particle size in the range of from 7 to 14 μm, and at least one of which is a mica pigment.
- 19. A composition according to any preceding claim in which the mica comprises one or more interference pigments having silver, green, violet and/or blue highlights.
- 20. A composition according to claim 19 in which the highlights are blue.
- 21. A composition according to any preceding claim in which the base formulation contains from 5 to 60% by weight of the astringent antiperspirant.
- 22. A composition according to claim 21 in which the base formulation contains from 20 to 45% by weight of the astringent antiperspirant.
- 23. A composition according to any preceding claim in which the astringent antiperspirant is present in a weight ratio to the smoothing aid selected in the range of from 12.5:1 to 50:1.
- 24. A composition according to claim 23 in which the astringent antiperspirant is present in a weight ratio to the smoothing aid selected in the range of from 20:1 to 35:1.
- 25. A composition according to claim 23 or 24 in which the smoothing aid comprises from 30% to 55% by weight wax and from 70 to 45% by weight mica.

- 26. A composition according to any preceding claim in which the astringent antiperspirant salt comprises an aluminium chlorohydrate.
- 27. A composition according to claim 26 in which the aluminium chlorohydrate has a weight average mean particle size (D50) in the range of from 15 to 30μm and >99% by weight of its particles of below 100μm and preferably below 80μm diameter.
- 28. A composition according to any preceding claim in which the base formulation comprises a hectorite clay, optionally hydrophobically surface treated and further comprises propylene carbonate.
- 29. A composition according to claim 28 in which the base formulation comprises the hectorite clay and the propylene carbonate in a weight ratio of from 2:1 to 5:1.
- 30. A composition according to any preceding claim in which the oil in the base formulation comprises a volatile oil and a non-volatile and preferably from 25 to 50% by weight of the volatile oil and from 50 to 75% of the non-volatile oil.
- 31. A composition according to claim 30 in which the non-volatile oil comprises a triglyceride oil.
- 32. A composition according to claim 31 in which the base formulation contains from 1 to 8% by weight and preferably 2 to 6% by weight of the triglyceride oil.
- 33. A composition according to claim 30 or 31 in which the triglyceride oil and the smoothing aid are present in a weight ratio of from 1:1 to 4:1, and preferably 2:1 to 3:1.

- 34. A composition according to any of claims 30 to 33 in which the non-volatile oil comprises a branched fatty alcohol containing from 14 to 30 carbon atoms.
- 35. A composition according to claim 34 in which the base formulation contains from 0.5 to 1.5% by weight of the branched fatty alcohol.
- 36. A composition according to claim 35 in which the branched fatty alcohol and smoothing aid are present in a weight ratio of from 1:2 to 2:1.
- 37. A composition according to any of claims 30 to 36 in which the non-volatile oil comprises an aliphatic or aromatic mono or di ester oil.
- 38. A composition according to claim 37 in which the ester oil is an alkyl benzoate.
- 39. A composition according to claim 37 or 38 in which the ester oil comprises from 2.5% to 10% by weight of the base formulation and preferably from 3.5 to 7% by weight.
- 40. A composition according to any of claims 30 to 39 in which the non-volatile oil comprises a polypropylene glycol-alkyl ether.
- 41. A composition according to claim 40 in which the polypropylene glycol moiety contains 10 to 18 propylene glycol units and the alkyl moiety is C₃ to C₆.

- 42. A composition according to claim 40 or 41 in which the base formulation contains from 15 to 30% by weight of the polypropylene glycol alkyl ether and preferably from 20 to 25%.
- 43. A composition according to any of claim 30 to 42 in which the non-volatile oil comprises polydimethiconol
- 44. A composition according to any preceding claim in which the propellant and the base formulation are present in a weight ratio of from 23:2 to 1:1 and preferably from 9:1 to 7:3.
- 45. An aerosol product comprising an aerosol composition charged into a dispensing canister comprising a reservoir, a spray outlet, a passageway for fluid leading from the reservoir to the spray outlet including a valve and an actuator for opening and closing the valve.
- 46. A product according to claim 45 in which the spray outlet has an internal diameter of from 350 to 600 microns and preferably from 350 to 450 microns.
- 47. A method for reducing perspiration comprising topically applying to skin a composition according to any preceding claim and particularly in the underarm.

International application No PCT/EP2009/065537

A. CLASSIFICATION OF SUBJECT MATTER INV. A61K8/04 A61K8 A61K8/92 A61Q15/00 A61K8/26 ADD. 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) A61Q A61K Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched Electronic data base consulted during the international search (name of data base and, where practical, search terms used) EPO-Internal, WPI Data C. DOCUMENTS CONSIDERED TO BE RELEVANT Category* Citation of document, with indication, where appropriate, of the relevant passages Relevant to claim No. χ WO 2005/048966 A (PROCTER & GAMBLE [US]; 1-8,10,SWAILE DAVID FREDERICK [US]; ALTONEN GENE 21-24.MICHA) 2 June 2005 (2005-06-02) 26,44, 45,47 page 7, line 26 - page 10, line 14 1-47 Υ figure 1 examples A-E GB 2 299 024 A (UNILEVER PLC [GB]) X 1-3,7,21,22, 25 September 1996 (1996-09-25) 26,44,47 Υ page 1, line 3 - line 7 1 - 47page 1, line 29 - line 35 page 3, line 28 - page 4, line 24 page 5, line 16 - line 34 page 9, line 1 - page 10, line 14 χ Further documents are listed in the continuation of Box C. X See patent family annex. Special categories of cited documents: "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 "A" document defining the general state of the art which is not considered to be of particular relevance invention "E" earlier document but published on or after the international "X" document of particular relevance: the claimed, invention cannot be considered novel or cannot be considered to document which may throw doubts on priority claim(s) or involve an inventive step when the document is taken alone which is cited to establish the publication date of another "Y" document of particular relevance; the claimed invention citation or other special reason (as specified) cannot be considered to involve an inventive step when the document is combined with one or more other such docu-"O" document referring to an oral disclosure, use, exhibition or ments, such combination being obvious to a person skilled other means "P" document published prior to the international filing date but later than 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 20 April 2010 04/05/2010 Name and mailing address of the ISA/ Authorized officer European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016

Lenzen, Achim

International application No PCT/EP2009/065537

C(Continua	tion). DOCUMENTS CONSIDERED TO BE RELEVANT	
Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Α	ANONYMOUS: "MIRASIL WAX-B" INTERNET ARTICLE, [Online] XP002537447 Retrieved from the Internet: URL:http://www.delfino.com/rhodiapc/text_a nd_graphics/pdf/Mirasil_WAX_B.pdf> [retrieved on 2009-07-15]	1-47
X	WO 2005/074877 A (PROCTER & GAMBLE [US]; SMITH SCOTT EDWARD [US]) 18 August 2005 (2005-08-18)	1,2,7, 21-24, 26,44,47
Υ	page 1, paragraphs 1,3 page 4, paragraph 8 page 5, line 12 - line 18 page 6, line 4 - line 8 page 9, paragraph 4 page 12, paragraph 3 examples 1-26	1-47
A	ANONYMOUS: "Sonneborn - White Oils, Petrolatums, Petroleum Distillates, Microcrystalline Waxes" INTERNET ARTICLE, [Online] XP002537448 Retrieved from the Internet: URL:http://www.sonneborn.com/products/pdf/sonneborn_brochure.pdf> [retrieved on 2009-07-16]	1-47
X	US 6 375 938 B1 (CLOTHIER JR JAMES G [US] ET AL) 23 April 2002 (2002-04-23)	1,7, 21-23, 26,44,47
Y	column 1, line 7 - line 14 column 1, line 60 - column 2, line 27 column 4, line 59 - column 5, line 60 column 6, line 10 - line 45 column 6, line 53 - line 60 column 7, line 15 - line 29 column 8, line 26 - line 27 examples 5,6	1-47
Υ	US 3 923 971 A (SEILINGER ALEXANDRE) 2 December 1975 (1975-12-02) column 1, line 7 - line 10 column 1, line 25 - line 32 column 1, line 45 - line 52 column 1, line 62 - column 2, line 30 column 2, line 49 - line 51 examples 6,7	9,11-20, 25
Y .	US 4 889 711 A (KAI MASANOBU [JP] ET AL) 26 December 1989 (1989-12-26) column 1, line 9 - line 12 column 1, line 60 - line 66 column 2, line 16 - line 24 column 3, line 11 - line 14 example 4	9,11-20, 25
	-/	

International application No PCT/EP2009/065537

	1 0 1/ 11 2003	,, 003331
tion). DOCUMENTS CONSIDERED TO BE RELEVANT		
Citation of document, with indication, where appropriate, of the relevant passages		Relevant to claim No.
EP 0 334 203 A (PROCTER & GAMBLE [US]) 27 September 1989 (1989-09-27) page 2, line 4 - line 5 page 2, line 31 - line 34 page 2, line 41 - line 51 page 3, line 27 - line 29 page 5, line 26 - page 6, line 9 examples I,II		28
WO 2007/031137 A1 (UNILEVER PLC [GB]; UNILEVER NV [NL]; LEVER HINDUSTAN LTD [IN]; FLETCHE) 22 March 2007 (2007-03-22) page 1, line 15 - line 21 page 30, line 22 - page 31, line 13		45,46
EP 1 428 521 A (UNILEVER PLC [GB]; UNILEVER NV [NL]) 16 June 2004 (2004-06-16) 		1-47
		•
		·
	EP 0 334 203 A (PROCTER & GAMBLE [US]) 27 September 1989 (1989-09-27) page 2, line 4 - line 5 page 2, line 31 - line 34 page 2, line 41 - line 51 page 3, line 27 - line 29 page 5, line 26 - page 6, line 9 examples I, II W0 2007/031137 A1 (UNILEVER PLC [GB]; UNILEVER NV [NL]; LEVER HINDUSTAN LTD [IN]; FLETCHE) 22 March 2007 (2007-03-22) page 1, line 15 - line 21 page 30, line 22 - page 31, line 13 EP 1 428 521 A (UNILEVER PLC [GB]; UNILEVER NV [NL])	EP 0 334 203 A (PROCTER & GAMBLE [US]) 27 September 1989 (1989–09–27) page 2, line 4 - line 5 page 2, line 31 - line 34 page 2, line 41 - line 51 page 3, line 27 - line 29 page 5, line 26 - page 6, line 9 examples I,II W0 2007/031137 A1 (UNILEVER PLC [GB]; UNILEVER NV [NL]; LEVER HINDUSTAN LTD [IN]; FLETCHE) 22 March 2007 (2007–03–22) page 1, line 15 - line 21 page 30, line 22 - page 31, line 13 EP 1 428 521 A (UNILEVER PLC [GB]; UNILEVER NV [NL])

Information on patent family members

International application No
PCT/EP2009/065537

					CI/EI	2009/009557
Patent document cited in search rep		Publication date		Patent family member(s)		Publication date
WO 20050489	066 A	02-06-2005	CA	2544215	A1	02-06-2005
			EP	1694283		30-08-2006
			MX	PA06005512		17-08-2006
 GB 2299024	A	 25-09-1996	 AU	695096		 06-08-1998
ub 2299024	А	25-09-1990	AU	4802496		03-10-1996
			ZA	9601867		08-09-1997
WO 20050748	377 A	18-08-2005	CA	2553638		18-08-2005
			EP	1722862		22-11-2006
			ΜX	PA06008589		28-08-2006
			US	2005169851	A1	04-08-2005
US 6375938	B1	23-04-2002	AT	409454	T	 15-10-2008
			ΑÜ	2002250089		11-11-2002
			CA	2442958		07-11-2002
			EP	1383477		28-01-2004
			ES	2314046		16-03-2009
			MX	PA03009717		29-01-2004
			WO	02087521	A2	07-11-2002
US 3923971	Α	02-12-1975	NONE			
 US 4889711	Α	26-12-1989	 EP	0229640	A2	22-07-1987
		20 12 1000	JΡ	1456779		09-09-1988
		•	ĴΡ	62164615		21-07-1987
			JP	63005376		03-02-1988
			PH	22779	Α	12-12-1988
EP 0334203	Α	27-09-1989	CA	1328817	С	26-04-1994
WO 20070311	.37 A1	22-03-2007	AR	057697	A1	 12-12-2007
			AT	432112		15-06-2009
			AU	2006291807	A 1	22-03-2007
			CA	2615546		22-03-2007
			CN	101272825		24-09-2008
			EP	1909916		16-04-2008
			ES	2325872		22-09-2009
			JP	2009502824		29-01-2009
			ZA 	200800902	A 	26-08-2009
EP 1428521	Α	16-06-2004	AU	2003268846	A1	01-07-2004
			BR	0305539		31-08-2004
			CA	2451296		13-06-2004
			CN	1506039		23-06-2004
			MX	PA03011578	Α	26-08-2005
EP 1428521	Α		PL	364007	A1	14-06-2004
		•	RŪ	2333741		20-09-2008
			US	2004120909	A1	24-06-2004
			ZA	200309676		13-06-2005