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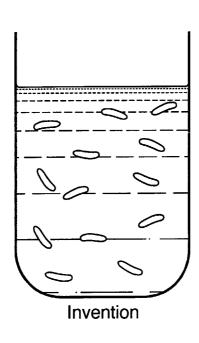
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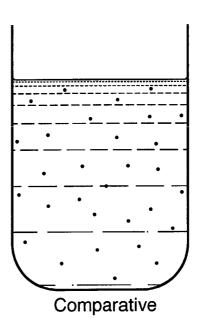
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[Continued on next page]

(54) Title: PERSONAL WASH COMPOSITIONS CONTAINING PARTICLE-IN-OIL DISPERSION





'O 02/067888 A1

(57) Abstract: The invention provides personal wash liquids comprising stable particle-in-oil dispersion designed to better deliver functional ingredients from such liquids. Specifically, the invention defines compositions for delivering solid particles and/or capsules.

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# PERSONAL WASH COMPOSITIONS CONTAINING PARTICLE-IN-OIL DISPERSION

5 The present invention relates to liquid personal wash compositions to effectively deposit solid particles comprising (a) surfactant (synthetic surfactant with or without soap); (b) large particle-in-oil dispersions comprising (i) a solid particle (0.1 to 150 microns, preferably 0.2 to 100 microns) or (ii) a capsule in either 10 petrolatum or in a hydrocarbon oil which is thickened or structured with an oil-mixable polymer; and (c) water soluble thickener mixable with the surfactant (e.g., concentrated surfactant) which can prevent physical separation of the large particle-in-oil dispersions 15 (dispersions having size greater than 50 microns) and are capable of preventing separation of large particle-in-oil dispersions for over 3 months.

Deposition of emollient oils from liquid cleansers can be achieved using small droplet deposition assisted with cationic polymer or by using large drops. However, deposition of solid particles (e.g., hydrophilic particle) from a cleansing product (e.g., wash-off cleanser) remains technically challenging.

U.S. Patent No. 5,661,189 to Grieveson discloses an aqueous liquid and a moisturizing composition containing emollients thickened with a hydrophobic thickener which emollients have a particle in the range of 50-500: These are thickened particles, not particle in oil dispersions. Moreover, the reference does not teach or suggest

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preparation of particle-in-oil disposition and subsequently mixing with surfactant solution. Rather, all ingredients are apparently mixed in a static mixer. If such process were used in the subject invention, particles would be mixed throughout the solution and deposit poorly in contrast to high particle/capsule deposition of the invention.

U.S. Patent No. 5,804,540 to Tsaur discloses personal wash liquids comprising low viscosity oils pre-thickened by non-antifoaming hydrophobic polymers.

Again, the reference fails to teach or suggest solid cosmetic particles or capsules as particle-in-oil dispersions. Further, there is no disclosure of a particle-in-oil dispersion in a surfactant system comprising water soluble thickener mixable with surfactant.

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U.S. patent No. 5,869,070 to Dixon et al. discloses stable liquid cleansing compositions with skin moisturizing agent and gel forming polymer. Skin moisturizing agents are petrolatum, polybutene and mixtures of petrolatum/polybutene with ratio of 3:1 to 5:1.

The reference fails to disclose solid, particle (e.g., cosmetic particle) in oil or capsules in oil. Further, although petrolatum/polybutylene is disclosed as moisturizer, there is no teaching or suggestion of a separate oil thickened by petrolatum/polybutylene. Finally, there is no teaching or suggestion of particle-in-oil found in a thickened/concentrated surfactant system.

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U.S. Patent No. 5,912,002 to Grieveson et al. discloses cleansing composition comprising surfactant and internal emulsion comprising cosmetic agent, emulsifier and carrier.

Grieveson differs from the compositions of the invention in that compositions of the subject invention require that the oil (of "particle in oil") is capable of separating and entrapping solid particles in presence of concentrated surfactant solution and without need of emulsifier. By contrast, Grieveson is an oil-in-oil emulsion where stability is achieved only by using an emulsion. Further, the oil (carrier oil) of Grieveson can be any oil in the range of 200 to 500,000 centistokes at 25°C while, in the subject invention, oil viscosity must be higher than 10,000 centistokes at 0.1 S<sup>-1</sup>.

U.S. Patent No. 6,066,608 to Glenn Jr. discloses compositions comprising surfactant, petrolatum, fumed silica and xanthan gum. Again, prior preparation of

20 particle/capsule-in-oil dispersion is not taught or suggested. The moisturizer rather is prepared by encapsulating moisturizing agents within a complex coacervate to protect drop integrity. There is no disclosure of first forming particle-in-oil dispersion; (b)

25 adding to surfactant base (e.g., by injecting); and (c) ensuring there is a large particle-in-oil dispersion which can suspend particle/capsule for enhanced delivery (e.g., by gently mixing surfactant and dispersion and then passing composition mixture through a screen).

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Unexpectedly, applicants have now found a superior way to deliver (a) solid particles; and/or (b) capsule (e.g., perfume-containing capsules) from a wash-off liquid by forming a particle-in-oil dispersion; wherein particle refers to any of (a) or (b) above; wherein the oil is petrolatum; or emollient oils which contain an oil mixable polymer; and wherein the surfactant system in which the particle-in-oil dispersion is found is thickened/concentrated (e.g., with xanthan gum or other thickener). The particle-in-oil dispersion remains stable 10 at room temperature for greater than 3 months with no visible physical separation and, in use, deposits far greater amounts of solid particle/capsule from a wash-off liquid composition than would be possible if the particles 15 were added alone rather than as particle-in-oil dispersion.

More specifically, the invention comprises:

A personal liquid composition comprising:

(a) a surfactant system comprising:

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- (i) 5% to 30%, preferably 8% to 25% by wt. of a surfactant selected from the group consisting of anionic surfactants, amphoteric surfactants, nonionic surfactants, cationic surfactants and mixtures thereof;
- (ii) 0.1% to 5% by wt. of a water soluble thickener mixable with said surfactant(s) and which prevents physical separation of a particle-in-oil dispersion as noted in (b) below from the liquid cleanser for three

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months or more as measured at room temperature; and

- (b) 2% to 20% by wt. of said particle-in-oil dispersion comprising:
  - (i) 60% to 99% by wt. of said dispersion of petrolatum or thickened emollient oil which contains oil mixable polymers, wherein said petrolatum or thickened oil can suspend solid particles having density greater than 1.05 (without need of an emulsifier, although such emulsifier may be optionally added if desired); and
  - (ii) 1% to 40% by wt. of a "particle" selected from the group consisting of (a) solid particles with particle size in range 0.1 to 250, preferably 0.2 to 200 microns, more preferably 0.5 to 150 microns; and (b) capsules with size in the range of 1-200 microns,

wherein the particle-in-oil dispersion is itself stable at room temperature for over 3 months; wherein the particle-in oil dispersion has size of about 50 to 5000 microns, preferably 100-1000 microns; and

wherein particle-in-oil dispersion is stable in the composition for three months or more at R.T.

Figure 1 is a schematic representation of Example 1A of the invention wherein the liquid is prepared in the order of the invention, i.e., by first mixing particle of (b)(ii) with petrolatum or thickened oil of (b)(i) to form particle-

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- 6 -

in-oil dispersion (b) and subsequently adding said dispersion into surfactant system (a) to form particle-in-oil in surfactant solution dispersion. As noted, sparkles are clumped or aggregated together in a much more visually noticeable manner.

By contrast, the comparative of Example 1, which is also schematically shown, has small filter particles rather than such large agglomerated lumps.

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Because of these clumps, applicants have demonstrated that many more glitters are deposited on substrate (e.g., skin or hair).

# DETAILED DESCRIPTION OF THE INVENTION

The present invention relates to compositions for delivering solid particles (preferably hydrophilic solid particles) and/or capsules (e.g., perfume-containing capsules) from wash off liquid compositions. These "particles" (referring generally to the solid particles, and capsules) are delivered from relatively large size (50 to 5000 microns), stable (for at least three months at room temperature), particle-in-oil dispersions found in thickened/concentrated surfactant compositions wherein the entire composition (containing particle-in-oil dispersion) is itself stable for three months at room temperature.

The compositions, as described below, will be defined by (1) surfactant system; (2) thickening system for said surfactant system; and (3) particle-in-oil composition

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wherein the particle-in-oil composition is itself defined by (a) particles (e.g., solid particle or capsules) and (b) oil in which particles are found. The oil is petrolatum; or thickened emollient oil which contains oil-mixable polymers to help further suspend/stabilize particles.

The invention is thus defined in more detail below.

### 10 SURFACTANT SYSTEM

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The composition of the invention comprises 5 to 30% by wt. surfactant. The surfactants in the composition may be anionic, nonionic, amphoteric/zwitterionic, cationic or mixtures thereof.

The anionic surfactant may be, for example, an aliphatic sulfonate, such as a primary alkane (e.g.,  $C_8-C_{22}$ ) sulfonate, primary alkane (e.g.,  $C_8-C_{22}$ ) disulfonate,  $C_8-C_{22}$  alkane sulfonate,  $C_8-C_{22}$  hydroxyalkane sulfonate or alkyl glyceryl ether sulfonate (AGES); or an aromatic sulfonate such as alkyl benzene sulfonate.

The anionic may also be an alkyl sulfate (e.g.,  $C_{12}$ - $C_{18}$  alkyl sulfate) or alkyl ether sulfate (including alkyl glyceryl ether sulfates). Among the alkyl ether sulfates are those having the formula:

RO(CH2CH2O)nSO3M

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wherein R is an alkyl or alkenyl having 8 to 18 carbons, preferably 12 to 18 carbons, n has an average value of greater than 1.0, preferably between 2 and 3; and M is a solubilizing cation such as sodium, potassium, ammonium or substituted ammonium. Ammonium and sodium laurel ether sulfates are preferred.

These differ from ether sulfates of the invention in that they are not branched.

The anionic may also be alkyl sulfosuccinates (including mono- and dialkyl, e.g., C6-C22 sulfosuccinates); alkyl and acyl taurates, alkyl and acyl sarcosinates, sulfoacetates, C8-C22 alkyl phosphates and phosphates, alkyl phosphate esters and alkoxyl alkyl phosphate esters, acyl lactates, C8-C22 monoalkyl succinates and maleates, sulphoacetates, and acyl isethionates.

20 Sulfosuccinates may be monoalkyl sulfosuccinates having the formula:

R<sup>4</sup>O<sub>2</sub>CCH<sub>2</sub>CH (SO<sub>3</sub>M) CO<sub>2</sub>M;

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25 amido-MEA sulfosuccinates of the formula

R<sup>4</sup>CONHCH<sub>2</sub>CH<sub>2</sub>O<sub>2</sub>CCH<sub>2</sub>CH (SO<sub>3</sub>M) CO<sub>2</sub>M

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wherein  $\mbox{R}^4$  ranges from  $\mbox{C}_8-\mbox{C}_{22}$  alkyl and M is a solubilizing cation;

amido-MIPA sulfosuccinates of formula

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RCONH (CH<sub>2</sub>) CH (CH<sub>3</sub>) (SO<sub>3</sub>M) CO<sub>2</sub>M

where M is as defined above.

Also included are the alkoxylated citrate sulfosuccinates; and alkoxylated sulfosuccinates such as the following:

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|| R-O-(CH<sub>2</sub>CH<sub>2</sub>O)<sub>n</sub>CCH<sub>2</sub>CH(SO<sub>3</sub>M)CO<sub>2</sub>M

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wherein n = 1 to 20; and M is as defined above.

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Sarcosinates are generally indicated by the formula  $\hbox{RCON\,(CH_3)\,CH_2CO_2M, wherein R ranges from $C_8$ to $C_{20}$ alkyl and $M$ is a solubilizing cation. }$ 

25 Taurates are generally identified by formula

R<sup>2</sup>CONR<sup>3</sup>CH<sub>2</sub>CH<sub>2</sub>SO<sub>3</sub>M

wherein  $\mbox{R}^2$  ranges from  $\mbox{C}_8-\mbox{C}_{20}$  alkyl,  $\mbox{R}^3$  ranges from  $\mbox{C}_1-\mbox{C}_4$  alkyl and M is a solubilizing cation.

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Another class of anionics are carboxylates such as follows:

 $R-(CH_2CH_2O)_nCO_2M$ 

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wherein R is  $C_8$  to  $C_{20}$  alkyl; n is 0 to 20; and M is as defined above.

Another carboxylate which can be used is amido alkyl polypeptide carboxylates such as, for example, Monteine LCQ<sup>(R)</sup> by Seppic.

Another surfactant which may be used are the  $C_8$ - $C_{18}$  acylisethionates. These esters are preferably prepared by reaction between alkali metal isethionate with mixed aliphatic fatty acids having from 6 to 18 carbon atoms and an iodine value of less than 20. At least 75% of the mixed fatty acids have from 12 to 18 carbon atoms and up to 25% have from 6 to 10 carbon atoms.

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Acyl isethionates, when present, will generally range from about 0.5-15% by weight of the total composition. Preferably, this component is present from about 1 to about 10%.

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The acyl isethionate may be an alkoxylated isethionate such as is described in Ilardi et al., U.S. Patent No. 5,393,466, hereby incorporated by reference into the subject application. This compound has the general formula:

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Wherein R is an alkyl group having 8 to 18 carbon atoms, m is an integer from 2 to 4, R' and R" are the same or different and are hydrogen or an alkyl group having from 1 to 4 carbon atoms and  $M^{+}$  is a monovalent cation such as, for example, sodium, potassium or ammonium.

In general the anionic component will comprise from

10 about 1 to 20% by weight of the composition, preferably 2 to
15%, most preferably 5 to 12% by weight of the composition.

It may also include a C8 to C14 unbranched fatty acid
hydrocarbon (e.g., lauric acid, palmitic acid, capric,
etc.).

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## ZWITTERIONIC AND AMPHOTERIC SURFACTANTS

Zwitterionic surfactants are exemplified by those which can be broadly described as derivatives of aliphatic quaternary ammonium, phosphonium, and sulfonium compounds, in which the aliphatic radicals can be straight or branched chain, and wherein one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate. A general formula for these compounds is:

$$(R^3)_x$$

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$$R^2 - Y^{(+)} - CH_2 - R^4 Z^{(-)}$$

wherein R<sup>2</sup> contains an alkyl, alkenyl, or hydroxy alkyl radical of from about 8 to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties and from 0 to about 1 glyceryl moiety; Y is selected from the group consisting of for example nitrogen, atoms; R<sup>3</sup> is an alkyl or monohydroxyalkyl group containing about 1 to about 3 carbon atoms; X is 1 when Y is a sulfur atom, and 2 when Y is a nitrogen or phosphorus atom; R<sup>4</sup> is an alkylene or hydroxyalkylene of from about 1 to about 4 carbon atoms and Z is a radical selected from the group consisting of carboxylate, sulfonate, sulfate, phosphonate, and phosphate groups.

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Amphoteric detergents which may be used in this invention include at least one acid group. This may be a carboxylic or a sulphonic acid group. They include quaternary nitrogen and therefore are quaternary amido acids. They should generally include an alkyl or alkenyl group of 7 to 18 carbon atoms. They will usually comply with an overall structural formula:

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30 where R<sup>1</sup> is alkyl or alkenyl of 7 to 18 carbon atoms;

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 $\mbox{R}^2$  and  $\mbox{R}^3$  are each independently alkyl, hydroxyalkyl or carboxyalkyl of 1 to 3 carbon atoms;

n is 2 to 4;

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m is 0 to 1;

X is alkylene of 1 to 3 carbon atoms optionally substituted with hydroxyl, and

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Y is 
$$-CO_2$$
- or  $-SO_3$ -

Suitable amphoteric detergents within the above general formula include simple betaines of formula:

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and amido betaines of formula:

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$$R^{2}$$
 $R^{1}$  - CONH  $(CH_{2})_{m}-N^{+}-CH_{2}CO_{2}^{-}$ 
 $R^{3}$ 

where m is 2 or 3.

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In both formulae  $R^1$ ,  $R^2$  and  $R^3$  are as defined previously.  $R^1$  may in particular be a mixture of  $C_{12}$  and  $C_{14}$  alkyl groups derived from coconut so that at least half,

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preferably at least three quarters of the groups  $R^1$  have 10 to 14 carbon atoms.  $R^2$  and  $R^3$  are preferably methyl.

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A further possibility is that the amphoteric detergent is a sulphobetaine of formula

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$$R^{1} - CONH (CH_{2})_{m} - N^{+} - (CH_{2})_{3}SO_{3}^{-}$$

where m is 2 or 3, or variants of these in which -  $(CH_2)_3SO_3$  is replaced by

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In these formulae  $R^1$ ,  $R^2$  and  $R^3$  are as discussed previously.

Amphoacetates and diamphoacetates are also intended to be covered in possible zwitterionic and/or amphoteric compounds which may be used.

The surfactant system may also optionally comprise a nonionic surfactant.

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

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The nonionic may also be a sugar amide, such as a polysaccharide amide. Specifically, the surfactant may be one of the lactobionamides described in U.S. Patent No. 5,389,279 to Au et al. which is hereby incorporated by reference or it may be one of the sugar amides described in Patent No. 5,009,814 to Kelkenberg, hereby incorporated into the subject application by reference.

Other surfactants which may be used are described in U.S. Patent No. 3,723,325 to Parran Jr. and alkyl polysaccharide nonionic surfactants as disclosed in U.S. Patent No. 4,565,647 to Llenado, both of which are also incorporated into the subject application by reference.

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Preferred alkyl polysaccharides are alkylpolyglycosides of the formula

 $R^2O(C_nH_{2n}O)_t(glycosyl)_x$ 

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wherein R<sup>2</sup> is selected from the group consisting of alkyl, alkylphenyl, hydroxyalkyl, hydroxyalkylphenyl, and mixtures thereof in which alkyl groups contain from about 10 to about 18, preferably from about 12 to about 14, carbon atoms; n is 0 to 3, preferably 2; t is from 0 to about 10, preferably 0; and x is from 1.3 to about 10, preferably from 1.3 to about 2.7. The glycosyl is preferably derived from glucose. To prepare these compounds, the alcohol or alkylpolyethoxy alcohol is formed first and then reacted with glucose, or a source of glucose, to form the glucoside (attachment at the 1-position). The additional glycosyl units can then be attached between their 1-position and the preceding glycosyl units 2-, 3-, 4- and/or 6-position, preferably predominantly the 2-position.

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Nonionic generally comprises 0 to 10% by wt. of the composition.

Cationic synthetic surfactant should not serve as the

sole surfactant in this product, but can be used as a cosurfactant at a lower level of from about 0.5% to about 6%
by weight. The more preferred types of cationic surfactant
are selected from the group consisting of: alkyl trimonium
chloride and methosulfate, and dialkyldimonium chloride and
methyl sulphate, and alkyl alkonium chloride and methyl

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sulphate and mixtures thereof. These surfactants contain  $C_{12}$  to  $C_{24}$  carbon atoms per alkyl chain. The most preferred cationic is selected from the group consisting of stearalkonium chloride, stearyltrimonium chloride. Distearyl-dimonium chloride, and mixtures thereof.

#### THICKENING SYSTEM

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Another required component of the invention is organic,
inorganic or polymeric stabilizer. Specifically, the
compositions comprise 0.1 to 10% by wt. of an organic,
inorganic or polymeric stabilizer which provides physical
stability of the large oil droplets, (droplets of
polymer/oil blend) in the surfactant composition at 40°C for
over four weeks.

Generally the organic polymeric stabilizer of the invention include, but are not limited to any of several long chain acyl derivatives or mixtures thereof. Included are the glycol mono- di- and triesters having about 14 to about 22 carbon atoms. Preferred glycol esters include the ethylene glycol mono- and distearates, glyceryl stearates, palm oil glyceride, tripalmitin, tristearin and mixtures thereof. Another example of suspending agent useful in the present invention include the alkanolamides having from about 14 to about 22 carton atoms. Preferred alkanolamides are stearic monoethanolamide, stearic diethanolamide stearic monoisopropanolamide, stearic monoethanolamide stearate and mixtures thereof. Still another example of a suspending agent useful in the present invention include the long chain fatty acid esters such as stearyl stearate, stearyl

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palmitate, palmityl palmitate, trihydroxystearylglycerol and tristearylglycerol. Still another example of a suitable suspending agent useful in the present invention include the long chain amine oxides having from about 14 to about 22 carbon atoms. Preferred amine oxides are hexadecyldimethylamine oxide and octadecyldimethylamide oxide.

Example of a suitable polymeric suspending agent (or thickening agent) useful in the present invention include 10 the carbohydrate gums such as cellulose gum, microcrystalline cellulose, cellulose gel, hydroxyethyl cellulose, hydroxypropyl cellulose, sodium carboxymethylcellulose, hydroxymethyl carboxymethyl cellulose, hydroxymethyl carboxypropyl cellulose, methyl 15 cellulose, ethyl cellulose, guar gum, gum karaya, gum tragacanth, gum arabic, gum acacia, gum agar, xanthan gum and mixtures thereof. Preferred carbohydrate gums are the cellulose gums and xanthan gum. Of all the above described types of suspending agents, preferred compounds include the 20 long chain glycol ester and the carbohydrate gums. Other stabilizers which may be used are set forth in U.S. Patent No. 5,854,293 to Glenn, Jr. at column 4, line 36 to column 6, line 65. This reference is hereby incorporated by reference into the subject application. 25

The thickener may also be a cationic polymer.

Suitable cationic polymers include Guar

hydroxypropyltrimonium chloride, Quaternium-19, -23, -40, 
57, poly(dimethyldiallylammonium chloride), poly (dimethyl
butenyl ammonium chloride)-, w- bis (triethanolammonium

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chloride), poly (dipropyldiallylammonium chloride), poly (methyl-beta propaniodiallylammonium chloride), poly (diallylpiperidinium chloride), poly (vinyl pyridinium chloride), quaternised poly (vinyl alcohol), quaternized poly (dimethylaminoethylmethacrylate) and mixtures thereof.

The suspending agent or mixtures of agent may be present from about 0.1 to 10% of the composition.

While not wishing to be bound by theory, thickener is delivered to aid in suspension of the particle-in-oil dispersion.

Specifically, the thickened surfactant system prevents

15 physical separation of particle-in-oil dispersion for over
three months.

# PARTICLE-IN-OIL DISPERSION

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20 The particle-in-oil dispersion comprises solid particles and well-structured oil that can stably suspend and entrap the solid particles in a concentrated surfactant solution. Stability of the particle-in-oil dispersion is achieved by the viscosity and the suspension power of the oil.

The suspended solid particles can be either inorganic or organic particles. Examples of inorganic include but are not limited to silica, talc, mica; and examples organic particles are silicone powders or capsules such as perfume capsules or vitamin E capsules. Capsules suitable for this

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invention are free flow powders prepared by various encapsulation processes such as interfacial polymerization encapsulation, coacervation encapsulation, spray drying/cooling encapsulation or spray coating encapsulation. These solid particles should be cosmetic grade approved for personal cleansing application. Typically the solid particles will have particle size of 0.1 to 150 microns, preferably 0.2 to 200 microns, more preferably 0.5 to 150 microns.

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The well-structured oil is defined as emollient oil that has a viscosity higher than 10,000 cps at 0.1 S<sup>-1</sup> (measured at room temperature (25°C)) and can stably suspend the solid particles at room temperature for over 3 months without visible physical separation. Preferably wellstructured oil is petrolatum or emollient oil structured/thickened with oil mixable polymers. Examples of emollient oils are mineral oils, triglycerides oils such as sunflower seed oil, castor oil or soybean oil, alkyl esters such as isopropylpalmitate or isopropylmyristate and silicone oils. These emollient oils alone are not useful for the invention due to its poor suspension/entrapping To suspend and entrap dense solid particles, properties. oil mixable polymer is required to add into these emollient oils. The oil mixable polymer works as a thickener and also works as structuring agent to suspend the solid particles. Examples of possible oil mixable polymers are hydrogenated or non-hydrogenated polymer of alkylene or isoalkylene such as polybutene, poly-alphaolefin, or polyester, polyacrylate and it's copolymer, and rubber thermoplastic block copolymers such as butadiene/styrene or styrene/butylene di-

or tri-block copolymers. Most preferably oil mixable polymers are rubber based thermoplastic block polymers available from Shell Chemical Company under the tradename of Kraton®.

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In general, the oil drop should be 5 times or greater (i.e., at least 5 times as great) as the size of the suspended particles.

Typically the well structured oil comprises 60% to 99%, preferably 70% to 95% of the particle-in-oil dispersion and the solid particle comprises 1% to 40%, preferably 5% to 30% of the dispersion.

# 15 OTHER INGREDIENTS

In addition to ingredients noted above, the composition (i.e., surfactant composition) may comprise any one of various ingredients which may be found in personal wash compositions, including organic solvents (e.g., polyols such as C1 to C4 alkanols), auxiliary thickeners, sequestering agents (e.g., tetrasodium ethylenediaminetetraacetate), coloring agents, opacifiers, and pearlizers.

The compositions may also include antimicrobial agents, alkanolamides, antioxidants (e.g., butylated hydroxytoluene), exfoliants (polyoxyethylene beads) etc.

Except in the operating and comparative examples, or 30 where otherwise explicitly indicated, all numbers in this

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description indicating amounts or ratios of materials or conditions or reaction, physical properties of materials and/or use are to be understood as modified by the word "about".

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Where used in the specification, the term "comprising" is intended to include the presence of stated features, integers, steps, components, but not to preclude the presence or addition of one or more features, integers, steps, components or groups thereof.

The following examples are intended to further illustrate the invention and are not intended to limit the invention in any way.

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Unless indicated otherwise, all percentages are intended to be percentages by weight.

# EXAMPLE 1: DEPOSITION OF SOLID PARTICLES

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The following example shows how deposition of large solid particles from liquid cleanser can be dramatically enhanced using stable particle-in-oil dispersions of this invention. The "particles" used for these examples were two glitters — visible Spectratek Geometic Silver Pigments of  $100 \times 100$  microns and  $100 \times 50$  microns.

First, applicants prepared a liquid cleansing base with composition as noted below in Table 1 for incorporation of the stable glitter-in-oil dispersions:

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Table 1: Composition of Liquid Cleansing Base

Lauric Acid	2.88 wt.%
Triethanol Amine	2.0 wt.%
Na Lauroamphoacetate	5.8 wt.%
Na Lauryl Sulfosuccinate	5.8 wt.%
Xanthan Gum	1.2 wt.%
Glycerin	10.0 wt.%
DMDM HYDANTOIN	0.20 wt.%
Perfume	0.5 wt.%
Water	To 100 wt.%

It should be noted that the liquid base is a thickened base, i.e., thickened with xanthan gum.

Applicants next prepared glitter-in-oil dispersions noted below in Table 2:

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Table 2: Composition of Glitter-in-Oil Dispersion

	Particle-in-Oil Composition				
Example 1A	10% Spectratek 100 x 100 microns Geometric				
Silver (e.g., "particle") 90% petrolatum/polybutene (Panalane H300E)					
Example 1B	10% Spectratek 100 x 50 microns Geometric Silver				
	90% petrolatum/polybutene (Panalane H300E) 2 to				
	1 ratio				

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Example 1A and 1B were prepared by first injecting 8 parts of the prepared glitter-in-oil dispersion shown in Table 2 into 92 parts of xanthan-thickened liquid cleanser using a syringe pump to form big noodle-like particles. The mixture was then passed through a screen with 500 micron openings once to make glitter-in-oil dispersions with particle size larger than 300 microns. Examined under microscope, these glitters are well entrapped inside the petrolatum/polymer oil droplet for both examples.

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Two comparative examples were also prepared for comparison. Comparative Example 1 was prepared by mixing 0.8 parts of glitter, Spectratek 100 x 100 Geometric Silver pigment, with 99.2 parts xanthan-thickened liquid base at 15 room temperature (i.e., it was not glitter-in-oil dispersion, only glitter in general composition). Comparative Example 2 was prepared by first injecting 7 parts of petrolatum/Panalene H300 (2/1 ratio) into 93 parts Comparative Example 1. The cleanser was then passed through a screen with 500 micron opening once to make the Comparative Example 2. In Comparative 2, the oil and glitter were separate dispersions in the liquid.

The protocol for applying particles personal wash is as follows:

- (1) Wet arm (e.g., forearm) with tap water;
- (2) Apply sample (e.g., 0.5 g) on forearm and rub for 10 seconds;
- (3) Rinse the forearm for 15 seconds (about 15 ml/sec flow rate) under warm tap water (about 30°-32°); and

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(4) Pat dry skin lightly with a paper towel.

Using such protocol, arm washed with either Example 1A or 1B showed much higher amount of visible shining particles retained on the skin than the arm washed with either Comparative Example 1 or Comparative Example 2. This example has clearly showed that deposition of solid particles from a liquid personal washing product can be achieved using stable particle-in-oil dispersion as claimed in the invention. To achieve such deposition, the solid particle should be stably entrapped inside the oil droplet.

### EXAMPLE 2

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15 Four examples were prepared to show the effect of oil suspension properties on stability of particle-in-oil dispersion in a liquid cleanser as noted below:

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Table 3: Composition of Particle-in-Oil Dispersion

Example 2A	5% Timiron MP 1005, less than 15 microns mica					
	from Rona					
	95% petrolatum					
Example 2B 5% Timiron MP 1005, less than 15 microns r						
	from Rona					
	95% Geahlene 500 from Penreco (mineral oil					
	thickened with oil mixable polymer)					
Comparative 1 5% Timiron MP 1005, less than 15 microns						
	from Rona					
	95% mineral oil					
Comparative 2	5% Timiron MP 1005, less than 15 microns mica					
	from Rona					
	95% DC 200, 5,000 cps silicone oil					

Two of the oils, petrolatum and Geahlene 500 (which is mineral oil thickened with a block copolymer), have good suspension properties suitable for the invention. The other two oils, mineral oil and silicone oil, do not have good suspension properties to stably suspend large particles in the oil and are not suitable for the application of this invention. The particle-in-oil dispersion was prepared by mixing 5 parts Timiron with 95 parts oil at 50°C for 10 to 20 minutes and cooled to room temperature before added into a liquid cleanser. The prepared particle-in-oil dispersions are stable at room temperature to both petrolatum and Geahlene 500 for over 3 months, but not stable for either silicone oil or mineral oil.

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5 parts of the above prepared particle-in-oil dispersion was then added into 95 parts liquid cleansing base using the xanthan structured liquid and the method described in Example 1. Mica was retained well inside the oil phase for Example 2A and 2B. For the comparative examples, most of the micas were released into the cleanser. These examples show that oil suspension properties is very critical in the invention so that liquid cleansers containing particle-in-oil dispersion may deposit the entrapped particles on skin surface for improved sensory or skin care properties. To be useful for the application, the particle-in-oil dispersion should be stable at room temperature for at least 3 months without visible physical separation.

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# EXAMPLES 3-6

In order to show that solid grade particles (not just glitter) could be used, applicants prepared (i) micron size talc, (ii) mica (iii) TiO2, and (iv) silicone powder as particles. Talc, mica, and silicone powder (micron size) are used to manipulate tactile properties; and TiO2 (in form of submicron, water-dispersible TiO2) is for UV protection. Compositions of the dispersion are set forth in Table 4 below:

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Table 4: Composition of Particle-in-Oil Dispersion

Example 3	30% Cimpact 710, 1.6 microns talc from Luznac				
	America (particle)				
}	70% petrolatum (oil)				
Example 4	10% Timiron MP 1005, less than 15 microns mica				
	from Rona (particle)				
	90% petrolatum (oil)				
Example 5	10% Titan M212, submicron water-dispersible TiO2				
	ex. Preserpse Inc. (particle)				
	90% petrolatum (oil)				
Example 6	10% 9506 silicone powder from Dow Corning				
	(particle)				
	90% petrolatum/polybutene H300E of 2 to 1 ratio				
	(oil)				
	<b>i</b>				

All the above particle-in-oil dispersions are stable at R.T. for 3 months. Examples 3-6 were prepared by adding 8 parts of the oil dispersion into 92 parts of the xanthan-structured liquid using the method and base of Example 1.

This example shows dry skin, after wash skin feel can be manipulated by using various particles. Particles as noted were stable in liquid for over 3 months.

# EXAMPLE 7 - Perfume Capsules-in-Oil

Liquid compositions with perfume-in-oil dispersion are set forth in Table 5 below:

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Table 5: Liquid Composition Containing Perfume Capsule-in-Oil Dispersion

Example 5A	92 parts: xanthan-thickened liquid			
	8 parts: 15% 180 micrometers Limonene			
	Capsule (ex. 3M) (particle) dispersed in			
	85% petrolatum (oil)			
Example 5B 92 parts xanthan-thickened liquid				
	8 parts: 30 micrometers Limonene Capsule			
	(ex 3M) (particle) dispersed in 85%			
	petrolatum (oil)			

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Same procedure described in Example 1 was used to prepare Examples 5A and 5B with composition shown in Table 4. A xanthan-thickened liquid with the composition same as that shown in Table 1 without perfume was prepared for the incorporation of perfume capsule-in-oil dispersion. A control sample prepared by mixing 1 part of free limonene fragrance with 99 parts of xanthan-thickened liquid at room temperature was used for comparison. Sample 5A showed much better perfume retention than the comparative example. Arms washed with Sample 5A have much stronger fresh limolene smell than the comparative example containing free limonene fragrance.

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# **CLAIMS**

1.	A	personal	liquid	composition	comprising:
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- (a) a surfactant system comprising:
  - (i) 5% to 30% by wt. of a surfactant selected from anionic surfactant, amphoteric surfactant, non-ionic surfactant, cationic surfactant and mixtures thereof;
  - (ii) 0.1% to 10% by wt. water soluble thickener within said surfactant system;
- (b) 2% to 20% by wt. of a particle-in-oil dispersion comprising:
  - (i) 60 to 99% by wt. of petrolatum or thickened emollient oil which contains oil mixable polymers, which petrolatum or thickened oil have a viscosity higher than 10,000 cps at 0.1 S<sup>-1</sup> and 25°C and can suspend solids having a density greater than 1.05 at room temperature over 3 months, without visible physical separation; and
  - (ii) 1 to 40% of a particle selected from:
    - (a) solids having particle size 0.5 to 150 microns; and
    - (b) capsules in size range of 1 to 200
      microns;

wherein the personal liquid composition is prepared by first mixing the particle of item (b)(ii) with the petrolatum or thickened oil of (b)(i) to form particle-in-oil dispersion(b) and

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subsequently adding said particle-in-oil dispersion (b) into surfactant system (a) to form a particle-in-oil surfactant solution dispersion having a drop size of 50 to 5000 microns; wherein the size of the particle-in-oil droplet is at least 5 times larger than the suspended particles; and wherein said personal liquid composition is stable over 3 months at room temperature.

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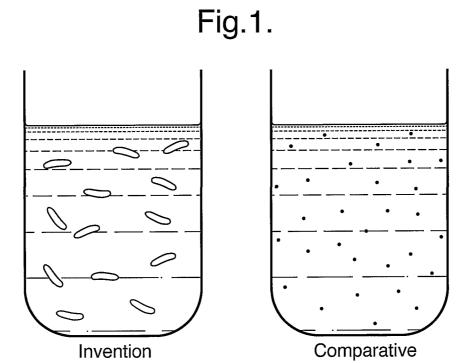
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- 2. A composition according to claim 1, wherein thickener (a)(ii) is xanthan gum.
- 3. A composition according to claim 1, wherein oil

  (b)(i) is a hydrocarbon oil containing an oil

  miscible polymer.

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# INTERNATIONAL SEARCH REPORT

Intertional Application No PCT/EP 02/01245

			101/11 01/01243
A. CLASSIF IPC 7	FICATION OF SUBJECT MATTER A61K7/48 A61K7/50		
According to	International Patent Classification (IPC) or to both national classifica	ation and IPC	
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	cumentation searched (classification system followed by classification	on symbols)	
Documentati	ion searched other than minimum documentation to the extent that s	uch documents are inclu	ded in the fields searched
Electronic da	ata base consulted during the international search (name of data bas	se and, where practical,	search terms used)
EPO-Int	ternal, WPI Data, PAJ, CHEM ABS Data		
C. DOCUME	ENTS CONSIDERED TO BE RELEVANT		
Category °	Citation of document, with indication, where appropriate, of the rele	evant passages	Relevant to claim No.
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А	WO 96 02224 A (UNILEVER PLC ;UNIL (NL)) 1 February 1996 (1996-02-01		
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Furth	ner documents are listed in the continuation of box C.	χ Patent family i	nembers are listed in annex.
"A" documer conside "E" earlier diffling de "L" documer which is citation "O" docume other m	nt defining the general state of the art which is not ered to be of particular relevance locument but published on or after the international ate in the properties of the pro	or priority date and cited to understand invention  "X" document of particulation cannot be conside involve an inventive and the consided document of particulation cannot be consided document is combinents, such combinents, such combinents act.  "8" document member	ished after the international filing date I not in conflict with the application but I the principle or theory underlying the I ar relevance; the claimed invention red novel or cannot be considered to e step when the document is taken alone I ar relevance; the claimed invention red to involve an inventive step when the ned with one or more other such docunination being obvious to a person skilled
	actual completion of the international search	_	he international search report
9	July 2002	29/07/2	UUZ
Name and m	nailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL – 2280 HV Rijswijk Tel. (+31–70) 340–2040, Tx. 31 651 epo nl,	Authorized officer Werner,	S
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\_ Information on patent family members

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