

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

(19) World Intellectual Property
Organization
International Bureau



(10) International Publication Number
WO 2018/044576 A1

(43) International Publication Date
08 March 2018 (08.03.2018)

(51) International Patent Classification:

A61K 8/73 (2006.01) A61K 8/44 (2006.01)
A61Q 19/10 (2006.01)

TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW,
KM, ML, MR, NE, SN, TD, TG).

(21) International Application Number:

PCT/US2017/047279

Published:

— with international search report (Art. 21(3))

(22) International Filing Date:

17 August 2017 (17.08.2017)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

62/381,184 30 August 2016 (30.08.2016) US

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(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, BN, BR, BW, BY, BZ,
CA, CH, CL, CN, CO, CR, CU, CZ, DE, DJ, DK, DM, DO,
DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN,
HR, HU, ID, IL, IN, IR, IS, JO, JP, KE, KG, KH, KN, KP,
KR, KW, KZ, LA, LC, LK, LR, LS, LU, LY, MA, MD, ME,
MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ,
OM, PA, PE, PG, PH, PL, PT, QA, RO, RS, RU, RW, SA,
SC, SD, SE, SG, SK, SL, SM, ST, SV, SY, TH, 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, LR, LS, MW, MZ, NA, RW, SD, SL, ST, SZ, TZ,
UG, ZM, ZW), Eurasian (AM, AZ, BY, KG, KZ, RU, TJ,
TM), European (AL, 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, RS, SE, SI, SK, SM,

(54) Title: PERSONAL CARE COMPOSITION

(57) Abstract: A personal care composition is provided, comprising: a vehicle; a surfactant; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 15 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and, wherein the body wash formulation contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.



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PERSONAL CARE COMPOSITION

[0001] The present invention relates to a personal care composition. In particular, the present invention relates to a personal care composition containing: a vehicle; a surfactant; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 15 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and, wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.

[0002] Conventional personal care systems such as shaving formulations (e.g., shaving creams and shaving gels), shampoos, hair conditioners, facial cleansing products, hair coloring systems, skin creams, lotions, under arm products (e.g., deodorants, antiperspirants), personal lubricating gels, oral care formulations (e.g., mouth washes, mouth moisturizers), hair styling agents (e.g., hair gels, mousses), hand soaps, shower gels, body washes, make-up products, sun screen systems have used commercially available polysaccharides such as nonionic water-soluble polysaccharide ethers (e.g., methyl cellulose (MC), hydroxypropyl methylcellulose (HPMC), hydroxyethyl cellulose (HEC), hydroxypropyl cellulose (HPC), ethyl hydroxyethyl cellulose (EHEC)), hydroxypropyl (HP) guar, hydroxyethyl guar, and hydroxypropyl starch and other nonionic starch and guar derivatives. Some hydrophobically modified polysaccharides have also been used in personal care products. The use of polysaccharides in personal care products face certain processing hurdles including incompatibility with other desirable ingredients, insolubility with certain other desirable ingredients, turbidity (when clarity is desired) and formulation stability.

[0003] Body washes (sometimes also referred to as shower gels) are a general term used to describe liquid surfactant containing formulations used to clean the body. While sometimes considered generically as "soap", such body wash formulations frequently do not contain soaps such as sodium or potassium salts of fatty acids. Conventional body wash formulations, typically contain one of a variety of components including one or more surfactants, various emollients, fragrances and other personal care ingredients. Conventional body washes offer less skin irritation, improved lather in hard water conditions and leave less residues on the skin and bathroom fixtures when compared with common soaps.

[0004] Conventional body wash formulations use a surfactant system consisting of a mixture of sodium laureth sulfate (SLES) (an anionic surfactant) and cocamidopropyl betaine (a zwitterionic surfactant). This surfactant mixture is frequently referred to as a

SLES/betaine surfactant mixture. While relatively inexpensive and effective, there is pressure from consumers to find a replacement for SLES in personal care compositions. There exists a belief by some that SLES may be a skin irritant. There also exists a belief by some that SLES may potentially contain low concentrations of 1,4-dioxane. Accordingly, some brand owners are seeking to provide "sulfate-free" formulations (i.e., personal care compositions (e.g., body washes) that do not contain SLES). There is also a perceived consumer demand for benign surfactants that are derived from biorenewable sources. Two types of such biorenewable surfactants include alkyl polyglucosides (APG) which are derivable from glucose and other monosaccharides and glycinate surfactants such as sodium cocoyl glycinate which may be derived from amino acids such as glycine.

[0005] Consumers expect that personal care compositions such as body wash formulations will exhibit a suitable viscosity. This viscosity serves at least two purposes. First, it improves handling and spreading of the composition. Second, it acts as a sensory cue that consumers tend to associate with product efficacy. Conventional SLES/betaine systems are easily thickened to a suitable viscosity using sodium chloride, a cheap and non-toxic material. Personal care compositions formulated with either APG or glycinate surfactants in substitution for SLES/betaine tend to be watery (non-viscous) in nature and are not subject to thickening through the addition of sodium chloride or other commonly used thickening agents. That is, many common polymers used to thicken conventional personal care compositions formulated with SLES/betaine such as hydroxyethyl cellulose (HEC) are incompatible with APG or glycinate surfactant compositions.

[0006] Accordingly, there remains a need for personal care compositions that are sulfate free but nevertheless exhibit a suitable viscosity. In particular, there remains a need for sulfate free body wash compositions.

[0007] The present invention provides a personal care composition, comprising: a vehicle; a surfactant; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 15 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and, wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.

[0008] The present invention provides a personal care composition, comprising: a vehicle; a surfactant; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and

wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.

[0009] The present invention provides a personal care composition, comprising: a vehicle; a surfactant; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose; wherein the hydrophobic group is a C8-16 linear or branched alkyl group bonded to the water-soluble cellulose ether base material; wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate; and wherein the personal care composition is a body wash formulation.

[0010] The present invention provides a personal care composition, comprising: a vehicle; a surfactant, wherein the surfactant includes at least one glycinate; a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of 900,000 to 2,500,000 Daltons; wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose; wherein the hydrophobic group is a C8-16 linear or branched alkyl group bonded to the water-soluble cellulose ether base material; wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate; and wherein the personal care composition is a body wash formulation.

DETAILED DESCRIPTION

[0011] We have surprisingly found that sulfate free personal care compositions can be appropriately thickened using a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms; wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and, wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.

[0012] Unless otherwise indicated, ratios, percentages, parts, and the like are by weight.

[0013] As used herein, unless otherwise indicated, the phrase "molecular weight" or Mw refers to the weight average molecular weight as measured in a conventional manner with gel permeation chromatography (GPC) and poly(ethylene oxide) standards. GPC techniques are discussed in detail in *Modern Size Exclusion Chromatography*, W. W. Yau, J. J. Kirkland, D. D. Bly; Wiley-Interscience, 1979, and in *A Guide to Materials*

Characterization and Chemical Analysis, J. P. Sibia; VCH, 1988, p.81-84. Molecular weights are reported herein in units of Daltons.

[0014] The term "cosmetically acceptable" as used herein and in the appended refers to ingredients typically used in personal care compositions, and is intended to underscore that materials that are toxic when present in the amounts typically found in personal care compositions are not contemplated as part of the present invention.

[0015] Preferably, the personal care composition of the present invention, comprises: a vehicle (preferably, wherein the vehicle is selected from the group consisting of water and aqueous C1-4 alcohol mixtures); a surfactant (preferably, wherein the surfactant is selected from the group consisting of alkyl polyglucosides (e.g., lauryl glucoside, coco-glucoside, decyl glucoside), glycines (e.g., sodium cocoyl glycinate), betaines (e.g., alkyl betaines such as cetyl betaine and amido betaines such as cocamidopropyl betaine), taurates (e.g., sodium methyl cocoyl taurate), glutamates (e.g., sodium cocoyl glutamate), sarcosinates (e.g., sodium lauroyl sarcosinate), isethionates (e.g., sodium cocoyl isethionate, sodium lauroyl methyl isethionate), sulfoacetates (e.g., sodium lauryl sulfoacetate), alaninates (e.g., sodium cocoyl alaninate), amphoacetates (e.g., sodium cocoamphoacetate), sulfonates (e.g., sodium C14-16 olefin sulfonate), succinates (e.g., disodium lauryl sulfosuccinate) and mixtures thereof); a water-soluble cellulose ether base material (preferably, wherein the water-soluble cellulose ether base material is selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose, ethyl hydroxyethyl cellulose, methyl cellulose, hydroxypropyl methyl cellulose, hydroxyethyl methyl cellulose) substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms (preferably, wherein the hydrophobic group contains a linear or branched carbon chain with 8 to 16 carbon atoms; more preferably, wherein the hydrophobic group is a C8-16 linear or branched alkyl group bonded to the water-soluble cellulose ether base material through at least one of an ether linkage (e.g., an ether linkage alone or an ether linkage and a 2-hydroxypropyl group), an ester linkage, an amide linkage and a urethane linkage; most preferably, wherein the hydrophobic group is a C8-12 alkyl group bonded to the water-soluble cellulose ether base material via either an ether linkage alone or an ether linkage and a 2-hydroxypropyl group); wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons (preferably, 875,000 to 5,000,000 Daltons; more preferably, 900,000 to 2,500,000 Daltons; most preferably, 1,250,000 to 1,750,000 Daltons); and, wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate (preferably, < 0.001 wt% alkyl sulfate and < 0.001 wt% alkyl

ether sulfate; more preferably, < 0.0001 wt% alkyl sulfate and < 0.0001 wt% alkyl ether sulfate; most preferably, < a detectable limit of alkyl sulfate and < a detectable limit of alkyl ether sulfate).

[0016] Preferably, the personal care composition of the present invention contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate. More preferably, the personal care composition of the present invention contains < 0.001 wt% alkyl sulfate and < 0.001 wt% alkyl ether sulfate. Still more preferably, the personal care composition of the present invention contains < 0.0001 wt% alkyl sulfate and < 0.0001 wt% alkyl ether sulfate. Most preferably, the personal care composition of the present invention contains < a detectable limit of alkyl sulfate and < a detectable limit of alkyl ether sulfate.

[0017] Preferably, the personal care composition of the present invention, comprises a vehicle, wherein the vehicle is selected from the group consisting of water, water and C1-4 alcohol mixture. More preferably, the personal care composition of the present invention, comprises: a vehicle, wherein the vehicle comprises water. Most preferably, the personal care composition of the present invention, comprises: a vehicle, wherein the vehicle is water.

[0018] Preferably, the personal care composition of the present invention, contains 50 to 99 wt% vehicle. More preferably, the personal care composition contains 70 to 95 wt% vehicle. Most preferably, the personal care composition contains 75 to 90 wt% vehicle.

[0019] Preferably, the personal care composition of the present invention, contains 50 to 99 wt% water. More preferably, the personal care composition contains 70 to 95 wt% water. Most preferably, the personal care composition contains 75 to 90 wt% water.

[0020] Preferably, the personal care composition of the present invention, comprises a surfactant, wherein the surfactant is selected from the group consisting of alkyl polyglucosides (e.g., lauryl glucoside, cocoyl-glucoside, decyl glucoside), glycinates (e.g., sodium cocoyl glycinate), betaines (e.g., alkyl betaines such as cetyl betaine and amido betaines such as cocamidopropyl betaine), taurates (e.g., sodium methyl cocoyl taurate), glutamates (e.g., sodium cocoyl glutamate), sarcosinates (e.g., sodium lauroyl sarcosinate), isethionates (e.g., sodium cocoyl isethionate, sodium lauroyl methyl isethionate), sulfoacetates (e.g., sodium lauryl sulfoacetate), alaninates (e.g., sodium cocoyl alaninate), amphotoacetates (e.g., sodium cocoamphoacetate), sulfonates (e.g., sodium C14-16 olefin sulfonate), succinates (e.g., disodium lauryl sulfosuccinate) and mixtures thereof. More preferably, the personal care composition of the present invention, comprises a surfactant, wherein the surfactant includes at least one of a betaine, a glycinate, a glucoside and a

succinate. Still more preferably, the personal care composition of the present invention, comprises a surfactant, wherein the surfactant includes at least one glycinate. Most preferably, the personal care composition of the present invention, comprises a surfactant, wherein the surfactant includes at least one glycinate, wherein the at least one glycinate is sodium cocoyl glycinate.

[0021] Preferably, the personal care composition of the present invention, comprises 0.01 to 35 wt% (more preferably, 1 to 30 wt%; still more preferably, 4 to 25 wt%, most preferably, 10 to 20 wt%) of a surfactant. More preferably, the personal care composition of the present invention, comprises 0.01 to 35 wt% (more preferably, 1 to 30 wt%; still more preferably, 4 to 25 wt%, most preferably, 10 to 20 wt%) of a surfactant, wherein the surfactant is selected from the group consisting of alkyl polyglucosides (e.g., lauryl glucoside, cocoyl-glucoside, decyl glucoside), glycinate (e.g., sodium cocoyl glycinate), betaines (e.g., alkyl betaines such as cetyl betaine and amido betaines such as cocamidopropyl betaine), taurates (e.g., sodium methyl cocoyl taurate), glutamates (e.g., sodium cocoyl glutamate), sarcosinates (e.g., sodium lauroyl sarcosinate), isethionates (e.g., sodium cocoyl isethionate, sodium lauroyl methyl isethionate), sulfoacetates (e.g., sodium lauryl sulfoacetate), alaninates (e.g., sodium cocoyl alaninate), amphotoacetates (e.g., sodium cocoamphoacetate), sulfonates (e.g., sodium C14-16 olefin sulfonate), succinates (e.g., disodium lauryl sulfosuccinate) and mixtures thereof. More preferably, the personal care composition of the present invention, comprises 0.01 to 35 wt% (more preferably, 1 to 30 wt%; still more preferably, 4 to 25 wt%, most preferably, 10 to 20 wt%) of a surfactant, wherein the surfactant includes at least one of a betaine, a glycinate, a glucoside and a succinate. Still more preferably, the personal care composition of the present invention, comprises 0.01 to 35 wt% (more preferably, 1 to 30 wt%; still more preferably, 4 to 25 wt%, most preferably, 10 to 20 wt%) of a surfactant, wherein the surfactant includes at least one glycinate. Most preferably, the personal care composition of the present invention, comprises 0.01 to 35 wt% (more preferably, 1 to 30 wt%; still more preferably, 4 to 25 wt%, most preferably, 10 to 20 wt%) of a surfactant, wherein the surfactant includes at least one glycinate, wherein the at least one glycinate is sodium cocoyl glycinate.

[0022] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material. More preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose, ethyl hydroxyethyl cellulose, methyl

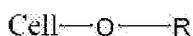
cellulose, hydroxypropyl methyl cellulose, hydroxyethyl methyl cellulose and mixtures thereof. Still more preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose and mixtures thereof. Most preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose.

[0023] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons. More preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of 875,000 to 5,000,000 Daltons. Still more preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of 900,000 to 2,500,000 Daltons. Most preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of 1,250,000 to 1,750,000 Daltons).

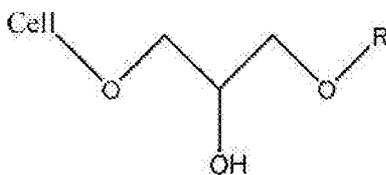
[0024] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons (preferably, 875,000 to 5,000,000 Daltons; more preferably, 900,000 to 2,500,000 Daltons; most preferably, 1,250,000 to 1,750,000 Daltons); and wherein the water-soluble cellulose ether base material is selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose, ethyl hydroxyethyl cellulose, methyl cellulose, hydroxypropyl methyl cellulose, hydroxyethyl methyl cellulose and mixtures thereof. Still more preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons (preferably, 875,000 to 5,000,000 Daltons; more preferably, 900,000 to 2,500,000 Daltons; most preferably, 1,250,000 to 1,750,000 Daltons); and wherein the water-soluble cellulose ether base material is selected from the group consisting of hydroxyethyl cellulose, hydroxypropyl cellulose and mixtures thereof. Most preferably, the personal care composition of the present invention, comprises

a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons (preferably, 875,000 to 5,000,000 Daltons; more preferably, 900,000 to 2,500,000 Daltons; most preferably, 1,250,000 to 1,750,000 Daltons); and wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose.

[0025] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms. More preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group having a linear or branched carbon chain with 8 to 16 carbon atoms bonded to the water-soluble cellulose ether base material through at least one of an ether linkage (e.g., an ether linkage alone or an ether linkage and a 2-hydroxypropyl group), an ester linkage, an amide linkage and a urethane linkage. Still more preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group, wherein the hydrophobic group is a C8-16 linear or branched alkyl group bonded to the water-soluble cellulose ether base material through at least one of an ether linkage (e.g., an ether linkage alone or an ether linkage and a 2-hydroxypropyl group), an ester linkage, an amide linkage and a urethane linkage. Most preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group, wherein the hydrophobic group is a C8-12 linear or branched alkyl group bonded to the water-soluble cellulose ether base material via an ether linkage alone (as in Formula I) or an ether linkage and a 2-hydroxypropyl group (as in Formula II)



(Formula I)



(Formula II)

wherein Cell—O is the water-soluble cellulose ether base material and wherein R is the C8-12 linear or branched alkyl group.

[0026] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms; wherein the degree of substitution, DS, of the hydrophobic group onto the water-soluble cellulose ether base material is 0.01 to 1 (more preferably, 0.02 to 0.2; most preferably, 0.025 to 0.1).

[0027] Preferably, the personal care composition of the present invention, comprises a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms (preferably, 8 to 16 carbon atoms; most preferably 8 to 12 carbon atoms). More preferably, the personal care composition of the present invention, comprises 0.1 to 15 wt% of a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms (preferably, 8 to 16 carbon atoms; most preferably 8 to 12 carbon atoms). Still more preferably, the personal care composition of the present invention, comprises 0.25 to 10 wt% of a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms (preferably, 8 to 16 carbon atoms; most preferably 8 to 12 carbon atoms). Yet more preferably, the personal care composition of the present invention, comprises 0.5 to 5 wt% of a water-soluble cellulose ether base material substituted with a hydrophobic group having carbon chain with 8 to 18 carbon atoms (preferably, 8 to 16 carbon atoms; most preferably 8 to 12 carbon atoms). Most preferably, the personal care composition of the present invention, comprises 0.75 to 2 wt% of a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms (preferably, 8 to 16 carbon atoms; most preferably 8 to 12 carbon atoms).

[0028] Preferably, the personal care composition of the present invention, further comprises at least one personal care ingredient. More preferably, the personal care composition of the present invention, further comprises at least one personal care ingredient, wherein the personal care ingredient is selected from the group consisting of emollients (e.g., hydrocarbon oils, esters, natural oils), cosmetically acceptable silicones (e.g., amodimethicone, cyclomethicone, dimethicone, dimethiconol, hexadecyl methicone, hexamethyldisiloxane, methicone, phenyl dimethicone, stearoxy dimethicone), waxes, soaps, sensory modifiers, lubricants, preservatives (e.g., benzoic acid, sorbic acid,

phenoxyethanol), antioxidants (e.g., butylated hydroxytoluene), chelating agents, antimicrobials, pH adjusting agents/buffers/neutralizing agents, humectants (e.g., glycerin, sorbitol, monoglycerides, lecithins, glycolipids, fatty alcohols, fatty acids, polysaccharides, sorbitan esters, polysorbates (e.g., Polysorbate 20, Polysorbate 40, Polysorbate 60, and Polysorbate 80), diols (e.g., propylene glycol), diol analogs, triols, triol analogs, polymeric polyols), sunscreen actives, vitamins, proteins/amino acids, plant extracts, natural ingredients, bio-actives, fragrances/perfumes, penetrants, polymers/resins/hair fixatives/film formers, surfactants/detergents/emulsifiers/opacifying agents, volatiles/propellants/solvents/carriers, liquid vehicles/solvents/carriers, salts, anti-static agents, anti-frizz agents, antidandruff agents, hair waving/straightening agents, absorbents, colorants, hard particles, and conditioning agents.

[0029] Preferably, the personal care composition of the present invention is a personal care composition selected from the group consisting of shampoos, leave-on hair conditioners, rinse-off hair conditioners, hair coloring agents, hair styling gels, soaps, body wash formulations, sunscreen agents and the like. More preferably, the personal care composition of the present invention is a personal care composition selected from the group consisting of shampoos, leave-on hair conditioners, rinse-off hair conditioners, hair coloring agents, hair styling gels, soaps, body wash formulations, sunscreen agents and the like; wherein the personal care composition contains < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl sulfate and < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl ether sulfate. More preferably, the personal care composition of the present invention is selected from the group consisting of shampoos, conditioners, hair styling agents and body wash formulations; wherein the personal care composition contains < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl sulfate and < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl ether sulfate. Most preferably, the personal care composition of the present invention is a body wash formulation; wherein the body wash formulation contains < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl sulfate and < 0.01 wt% (preferably < 0.001 wt%; more preferably, < 0.0001 wt%; most preferably, < a detectable limit) of alkyl ether sulfate.

[0030] Preferably, the personal care composition of the present invention is a body wash formulation. More preferably, the personal care composition of the present invention is a

body wash formulation, wherein the surfactant is selected from the group consisting of alkyl polyglucosides (e.g., lauryl glucoside, cocoyl-glucoside, decyl glucoside), glycinates (e.g., sodium cocoyl glycinate), betaines (e.g., alkyl betaines such as cetyl betaine and amido betaines such as cocamidopropyl betaine), taurates (e.g., sodium methyl cocoyl taurate), glutamates (e.g., sodium cocoyl glutamate), sarcosinates (e.g., sodium lauroyl sarcosinate), isethionates (e.g., sodium cocoyl isethionate, sodium lauroyl methyl isethionate), sulfoacetates (e.g., sodium lauryl sulfoacetate), alaninates (e.g., sodium cocoyl alaninate), amphoacetates (e.g., sodium cocoamphoacetate), sulfonates (e.g., sodium C14-16 olefin sulfonate), succinates (e.g., disodium lauryl sulfosuccinate) and mixtures thereof; more preferably, wherein the surfactant includes at least one of a betaine, a glycinate, a glucoside and a succinate; yet more preferably, wherein the surfactant includes at least one glycinate; most preferably, wherein the at least one glycinate is sodium cocoyl glycinate); wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose (preferably, wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons (preferably, 875,000 to 5,000,000 Daltons; more preferably, 900,000 to 2,500,000 Daltons; most preferably, 1,250,000 to 1,750,000 Daltons); and wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group having a linear or branched carbon chain with 8 to 12 carbon atoms bonded to the water-soluble cellulose ether base material through at least one of an ether linkage (e.g., an ether linkage alone or an ether linkage with a 2-hydroxypropyl group), an ester linkage, an amide linkage and a urethane linkage (more preferably, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group, wherein the hydrophobic group is a C8-12 linear or branched alkyl group bonded to the water-soluble cellulose ether base material through at least one of an ether linkage (e.g., an ether linkage alone or an ether linkage with a 2-hydroxypropyl group), an ester linkage, an amide linkage and a urethane linkage; most preferably, wherein the water-soluble cellulose ether base material is substituted with a hydrophobic group, wherein the hydrophobic group is a C8-12 linear or branched alkyl group bonded to the water-soluble cellulose ether base material through either an ether linkage alone or an ether linkage and a 2-hydroxypropyl group).

[0031] Preferably, the personal care composition of the present invention is a body wash formulation, wherein the body wash formulation has a viscosity of $\geq 3,000$ mPa·s as determined according to the method used herein in the Examples. More preferably, the personal care composition of the present invention is a body wash formulation, wherein the body wash formulation has a viscosity of 3,000 to 15,000 mPa·s as determined according to

the method used herein in the Examples. Still more preferably, the personal care composition of the present invention is a body wash formulation, wherein the body wash formulation has a viscosity of 4,000 to 12,000 mPa·s as determined according to the method used herein in the Examples. Most preferably, the personal care composition of the present invention is a body wash formulation, wherein the body wash formulation has a viscosity of 5,000 to 9,000 mPa·s as determined according to the method used herein in the Examples.

[0032] Preferably, the personal care composition of the present invention further comprises a pH adjusting agent. More preferably, the personal care composition of the present invention, further comprises a pH adjusting agent, wherein the personal care composition is a body wash formulation. Most preferably, the personal care composition of the present invention, further comprises a pH adjusting agent, wherein the personal care composition is a body wash formulation and wherein the body wash formulation has a pH of 5 to 9 (preferably, 6 to 8; most preferably, 6.25 to 7.75).

[0033] Preferably, the pH adjusting agent is selected from the group consisting of citric acid, lactic acid, hydrochloric acid, aminoethyl propanediol, triethanolamine, monoethanolamine, sodium hydroxide, potassium hydroxide, amino-2-methyl-1-propanol. More preferably, the pH adjusting agent is selected from the group consisting of citric acid, lactic acid, sodium hydroxide, potassium hydroxide, triethanolamine, amino-2-methyl-1-propanol. Most preferably, the pH adjusting agent is selected from the group consisting of citric acid and sodium hydroxide.

[0034] Preferably, the personal care composition of the present invention further comprises a biocide. More preferably, the personal care composition of the present invention further comprises a biocide, wherein the biocide is selected from the group consisting of phenoxyethanol, benzoic acid, benzyl alcohol, sodium benzoate, DMDM hydantoin, 2-ethylhexyl glyceryl ether and isothiazolinone (e.g., methylchloroisothiazolinone, methylisothiazolinone). Still more preferably, the personal care composition of the present invention, further comprises a biocide, wherein the biocide is an isothiazolinone (more preferably, wherein the biocide is selected from the group consisting of methylisothiazolinone, methylchloroisothiazolinone and mixtures thereof; most preferably, wherein the biocide is methylisothiazolinone). Most preferably, the personal care composition of the present invention, further comprises a biocide, wherein the biocide is an isothiazolinone (more preferably, wherein the biocide is selected from the group consisting of methylisothiazolinone, methylchloroisothiazolinone and mixtures thereof; most

preferably, wherein the biocide is methylisothiazolinone); and wherein the personal care composition is a body wash formulation.

[0035] Preferably, the personal care composition of the present invention further comprises a soap. More preferably, the personal care composition of the present invention, further comprises a soap, wherein the soap is selected from the group consisting of sodium stearate, sodium laurate, sodium tallowate, sodium palmitate, potassium stearate, potassium laurate, potassium tallowate, potassium palmitate and mixtures thereof (more preferably, wherein the soap is selected from the group consisting of sodium stearate, sodium laurate, potassium stearate, potassium laurate and mixtures thereof; still more preferably, wherein the soap is selected from the group consisting of sodium stearate, potassium stearate and mixtures thereof; most preferably, wherein the soap is sodium stearate). Most preferably, the personal care composition of the present invention, further comprises a soap, wherein the soap is selected from the group consisting of sodium stearate, sodium laurate, sodium tallowate, sodium palmitate, potassium stearate, potassium laurate, potassium tallowate, potassium palmitate and mixtures thereof (more preferably, wherein the soap is selected from the group consisting of sodium stearate, sodium laurate, potassium stearate, potassium laurate and mixtures thereof; still more preferably, wherein the soap is selected from the group consisting of sodium stearate, potassium stearate and mixtures thereof; most preferably, wherein the soap is sodium stearate); and wherein the personal care composition is a body wash formulation.

[0036] Preferably, the personal care composition of the present invention further comprises a colorant. More preferably, the personal care composition of the present invention, further comprises a colorant, wherein the personal care composition is a body wash formulation.

[0037] Some embodiments of the present invention will now be described in detail in the following **Examples**.

Comparative Examples C1-C9 and Examples 1-11

[0038] In each of **Comparative Examples C1-C9** and **Examples 1-11**, a 500 mL, four-necked, round-bottomed flask was charged with the Initial Components (as identified in **TABLE 1** in the quantities noted in **TABLE 1**). The flask was then fitted with a nitrogen inlet connected to a 60 mL pressure equalizing addition funnel, rubber septum cap, a stirring paddle connected to an electric motor, and a Claisen adaptor connected to a Friedrich condenser with a mineral oil bubbler outlet.

[0039] To the addition funnel was then charged a mixture of glycidyl ether (or alkyl bromide) (type and amount noted in **TABLE 1**) and isopropyl alcohol (in amount noted in

TABLE 1). Then, while stirring its contents, the head space of the flask was purged with a slow, steady flow of nitrogen for one hour to remove any entrained oxygen.

[0040] With continued stirring, a 50% aqueous sodium hydroxide solution was then added (in the quantity noted in **TABLE 1**) drop wise to the flask contents using a plastic syringe. Following the addition of the sodium hydroxide solution, the flask contents were allowed to stir for one hour, after which the solution of glycidyl ether (or alkyl bromide) in isopropyl alcohol in the addition funnel was added drop wise into the contents of the flask. The contents of the flask were then stirred under nitrogen for 20 minutes. The contents of the flask were then heated under nitrogen using a heating mantle and allowed to reflux for 4.5 hours.

[0041] The contents of the flask were then cooled by placing the flask in an ice water bath while maintaining a positive nitrogen pressure on the flask contents. The contents of the flask were then neutralized via the addition thereto of glacial acetic acid (5.0 g) using a syringe. The contents of the flask were then stirred for 10 minutes under nitrogen. The contents of the flask were then vacuum filtered through a large fritted metal Buchner funnel. The resulting filter cake was then washed three consecutive times in the Buchner funnel. First the filter cake was washed by adding a mixture of water (36 g) and isopropyl alcohol (164 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Then the filter cake was washed by adding a mixture of water (20 g) and isopropyl alcohol (180 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Finally, the filter cake was washed by adding a mixture of isopropyl alcohol (200 g), 40% aqueous glyoxal (0.44 g) and acetic acid (0.14 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. The filter cake was then briefly air-dried before being dried overnight under vacuum at 50 °C. The filter cake was then manually ground using a mortar and pestle and then sieved through a #30 U.S. sieve mesh to provide the product.

[0042] The mass of product recovered along with the volatiles and ash content of the product are provided in **TABLE 2**.

TABLE I

Ex. #	Initial Components			50% aq. NaOH (g)	Mole Ratio		Addition Funnel		
	Hydroxyethyl Cellulose (HEC) Type	Isopropyl alcohol (IPA) (g)	Deionized water (g)		HYD/HEC	NaOH/HEC	Glycidyl ether (except as indicated) (HYD)		
							Type	(g)	IPA (g)
C1	A	23.36	121	19	0.20	0.63	n-butyl-	2.2	6.0
C2	A	22.93	122	19	0.10	0.64	n-butyl-	1.1	6.1
C3	A	23.10	118	18	0.30	0.63	n-butyl-	3.4	6.0
C4	A	23.01	118	18	0.25	0.63	n-butyl-	2.7	6.1
C5	A	23.68	119	18	0.20	0.60	isobutyl-	2.2	6.0
C6	A	23.16	118	19	0.10	0.64	isobutyl-	1.1	6.0
C7	A	23.08	120	20	0.25	0.63	isobutyl-	2.7	6.0
C8	A	23.25	118	18	0.30	0.62	isobutyl-	3.4	6.0
C9	B	23.01	119	20	0.26	0.63	n-octyl-	4.1	6.0
1	A	23.34	119	19	0.29	0.62	n-octyl-	4.5	8.3
2	A	23.07	119	18	0.20	0.65	n-octyl-	3.1	6.1
3	A	23.08	118	18	0.10	0.64	n-octyl-	1.6	6.1
4	A	23.02	118	18	0.26	0.63	n-octyl-	4.0	6.0
5	A	23.61	120	19	0.29	0.61	2-ethyl hexyl-	4.5	8.1
6	A	23.24	119	19	0.19	0.63	2-ethyl hexyl-	3.0	7.3
7	A	23.30	118	18	0.10	0.64	2-ethyl hexyl-	1.6	6.3
8	A	23.02	118	18	0.26	0.64	2-ethyl hexyl-	4.0	8.0
9	C	23.03	120	19	0.26	0.64	n-octyl-	3.7	6.0
10	A	23.22	120	19	0.27	0.60	D	4.3	6.0
11	A	23.69	119	21	0.26	0.58	E	4.3	6.1

A – CELLOSE[™] QP-100MH hydroxyethyl cellulose with a weight average molecular weight, Mw, of 1,600,000 available from The Dow Chemical Company

B – CELLOSE[™] AM-103 hydroxyethyl cellulose with a Mw of 380,000 available from The Dow Chemical Company

C – CELLOSE[™] AP-4400MH hydroxyethyl cellulose with a Mw of 900,000 available from The Dow Chemical Company

D – n-octyl bromide used in place of glycidyl ether in this Example in quantity listed

E – 2-ethyl hexyl bromide used in place of glycidyl ether in this Example in quantity listed

Example 12

[0043] In **Example 12**, a 500 mL, four-necked, round-bottomed flask was charged with the CELLOSIZETM HEC QP-52,000H (33.89 g) and a mixture of isopropyl alcohol (174.7 g) and distilled water (27.5 g). The flask was then fitted with a stirring paddle and electric motor, a rubber serum cap, a nitrogen inlet and a Claisen adaptor fitted with a subsurface thermocouple and a Friedrich condenser connected to a mineral oil bubbler. The thermocouple was connected to a J-KEM controller and to a heating mantle.

[0044] While stirring the slurry, the head space in the flask was purged with nitrogen for one hour. Then a 50% aqueous sodium hydroxide solution (4.37 g) was added dropwise to the flask contents using a plastic syringe. The flask contents were left to stir under nitrogen for 30 minutes. Then 1-bromododecane (3.32 g) was added dropwise to the flask over a period of a minute. The flask contents were left to stir under nitrogen for 10 minutes. The flask contents were then heated to reflux for 4.5 hours with a 100 °C set point temperature on the J-KEM controller.

[0045] The contents of the flask were then cooled to room temperature by placing the flask in an ice water bath while maintaining a positive nitrogen pressure on the flask contents. The contents of the flask were then neutralized via the addition thereto of glacial acetic acid (6.43 g) using a syringe. The contents of the flask were then stirred for 10 minutes under nitrogen. The contents of the flask were then vacuum filtered through a large fritted metal Buchner funnel. The resulting filter cake was then washed four consecutive times in the Buchner funnel. First the filter cake was washed by adding a mixture of water (49 g) and isopropyl alcohol (221 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Then the filter cake was washed by adding a mixture of water (20 g) and isopropyl alcohol (180 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Then, the filter cake was washed by adding isopropyl alcohol (180 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Finally, the filter cake was washed by adding isopropyl alcohol (180 g), 40% aqueous glyoxal (0.60 g) and acetic acid (0.20 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. The filter cake was then briefly air-dried before being dried overnight under vacuum at 50 °C. The filter cake was then manually

ground using a mortar and pestle and sieved through a #30 U.S. sieve mesh plate to provide the product.

[0046] The mass of product recovered along with the volatiles and ash content of the product are provided in **TABLE 2**.

Example 13

[0047] In **Example 13**, a 500 mL, four-necked, round-bottomed flask was charged with the CELLOSIZETM HEC QP-52,000H (33.78 g) and a mixture of isopropyl alcohol (174.9 g) and distilled water (27.3 g). The flask was then fitted with a stirring paddle and electric motor, a rubber serum cap, a nitrogen inlet and a Claisen adaptor fitted with a subsurface thermocouple and a Friedrich condenser connected to a mineral oil bubbler. The thermocouple was connected to a J-KEM controller and to a heating mantle.

[0048] While stirring the slurry, the head space in the flask was purged with nitrogen for one hour. Then a 50% aqueous sodium hydroxide solution (5.43 g) was added dropwise to the flask contents using a plastic syringe. The flask contents were left to stir under nitrogen for 30 minutes. Then 1-bromododecane (6.67 g) was added dropwise to the flask over a period of a minute. The flask contents were left to stir under nitrogen for 10 minutes. The flask contents were then heated to reflux for 4.5 hours with a 100 °C set point temperature on the J-KEM controller.

[0049] The contents of the flask were then cooled to room temperature by placing the flask in an ice water bath while maintaining a positive nitrogen pressure on the flask contents. The contents of the flask were then neutralized via the addition thereto of glacial acetic acid (6.29 g) using a syringe. The contents of the flask were then stirred for 10 minutes under nitrogen. The contents of the flask were then vacuum filtered through a large fritted metal Buchner funnel. The resulting filter cake was then washed four consecutive times in the Buchner funnel. First the filter cake was washed by adding a mixture of water (49 g) and isopropyl alcohol (221 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Then the filter cake was washed by adding a mixture of water (20 g) and isopropyl alcohol (180 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Then, the filter cake was washed by adding isopropyl alcohol (180 g) to the filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. Finally, the filter cake was washed by adding isopropyl alcohol (180 g), 40% aqueous glyoxal (0.60 g) and acetic acid (0.20 g) to the

filter cake in the Buchner funnel and stirring the contents for five minutes followed by vacuum removal of the wash liquor through the Buchner funnel. The filter cake was then briefly air-dried before being dried overnight under vacuum at 50 °C. The filter cake was then manually ground using a mortar and pestle and sieved through a #30 U.S. sieve mesh plate to provide the product.

[0050] The mass of product recovered along with the volatiles and ash content of the product are provided in **TABLE 2**.

Volatiles Content

[0051] The volatile content (in wt%) in the product reported in **TABLE 2** was determined according to ASTM D-2364.

Ash Content

[0052] The ash content (in wt%) in the product reported in **TABLE 2** was determined according to ASTM D-2364, with the ash content reported as sodium acetate.

Viscosity

[0053] The viscosity of a 1 wt% aqueous solution of the product (corrected for ash and volatiles) was determined using a TA Instruments DHR-3 rheometer at 25 °C, equipped with a stainless steel 60 mm, 0.5° cone and plate sensor, a gap set at 17 microns and at a shear rate of 6.31 s⁻¹. The results are provided in **TABLE 2**.

TABLE 2

Product		Volatiles (wt%)	Ash (wt%)	Viscosity (mPa·s)
Ex. #	(g)			
C1	22.65	1.44	5.35	4,098
C2	22.41	2.44	5.88	4,560
C3	23.05	2.53	6.47	3,442
C4	23.10	2.47	7.02	3,651
C5	24.10	2.35	7.48	2,287
C6	22.76	2.74	6.36	4,219
C7	23.10	3.03	7.23	1,031
C8	24.78	2.91	12.67	3,719
C9	22.07	3.60	8.10	1,399
1	25.01	2.03	13.44	F
2	22.55	1.46	7.73	4,851
3	22.15	1.72	6.20	8,970
4	22.34	1.26	7.26	2,155
5	22.91	1.59	6.09	6,113
6	22.68	1.32	8.04	7,496
7	22.18	1.16	7.49	4,404
8	22.90	1.81	5.92	4,114
9	22.09	2.90	6.70	4,280
10	22.80	2.60	7.10	9,022
11	22.92	2.60	7.00	3,982
12	30.59	1.43	1.87	5,380
13	31.40	1.88	3.63	1,642
F – product insoluble in water				

Degree of Substitution

[0054] The degree of substitution of the glycidyl ether (hydrophobic group) substituent on the water-soluble cellulose ether base material for the product polymers produced according to **Examples 1-3** and **5-7** was determined through analysis of ^1H NMR spectra taken with a Varian Inova 600 MHz spectrometer using the following acquisition parameters: 10 seconds relaxation delay, 2 seconds acquisition time, 90 degree pulse of 7.25 μs , 128-256 scans. All measurements were taken without sample spinning at 10 °C and calibrating with ethylene glycol. The ^1H NMR spectra were referenced at 4.9 ppm for the peak of HOD at the noted temperature. The degree of substitution determined from the analysis are provided in **TABLE 3**.

Table 3

<u>Ex. #</u>	<u>Degree of Substitution</u>
1	0.094
2	0.083
3	0.034
5	0.093
6	0.054
7	0.028

Comparative Example C10

[0055] The hexadecyl-modified CELLOSIZTM QP-100MH used herein was prepared according to Example 22 of U.S. Patent No. 9,266,971.

Comparative Examples F1-F5 and Examples 14-19: Body wash formulation

[0056] Deionized water was added to a beaker. A heat source having a set point temperature adjusted to 60°C was brought into contact with the beaker. While the contents of the beaker were heating, cocamidopropyl betaine (Amphosol[®] CA available from Stepan Company) and decyl glucoside (EcoSenseTM 3000 available from The Dow Chemical Company) were added to the beaker. Once the beaker contents reached 60 °C, disodium lauryl sulfosuccinate (Mackanate[®] LO available from Solvay Novacare) was added to the beaker. The contents of the beaker were left to stir for 15 minutes before removing the heat source from contact with the beaker. Once the beaker contents cooled to 35 °C, the additive noted in **TABLE 5** was added to the contents of the beaker. The pH of the beaker contents was then adjusted to 6.5 with citric acid and methylisothiazolinone (Neolone 950 available from The Dow Chemical Company) was added to provide a body wash formulation having the composition noted in **TABLE 4**. The resulting body wash formulations were allowed to stand for two (2) days and observed for phase separation. The observations are provided in **TABLE 5**.

TABLE 4

Component	Conc. in Body Wash Formulation (wt%)
Cocamidopropyl betaine	3.85
Decyl glucoside	6.00
Disodium lauryl sulfosuccinate	4.80
Additive noted in TABLE 5	1.00*
Citric acid	0.16
Methylisothiazolinone	0.05
* -- unless otherwise noted in TABLE 5	

TABLE 5

Body Wash Formulation	Additive	Observation
Comp. Ex. F1	sodium hydroxypropyl starch*	Homogeneous
Comp. Ex. F2	CELLOSIZETM QP-100MH	Phase separated
Comp. Ex. F3	Prod. Comp. Ex. C4	Phase separated
Comp. Ex. F4	Prod. Comp. Ex. C9	Homogeneous
Comp. Ex. F5	Prod. Comp. Ex. C10	Phase separated
Ex. 14	Prod. Ex. 4	Homogeneous
Ex. 15	Prod. Ex. 8	Homogeneous
Ex. 16	Prod. Ex. 9	Homogeneous
Ex. 17	Prod. Ex. 10	Homogeneous
Ex. 18	Prod. Ex. 12	Homogeneous
Ex. 19	Prod. Ex. 13	Homogeneous
* -- sodium hydroxypropyl starch was loaded in formulation at 10 wt%		

Comparative Examples F6-F10 and Examples 20-25: Body wash formulation

[0057] Deionized water and cocamidopropyl betaine (Amphosol® CA available from Stepan Company) were added to a beaker. A heat source having a set point temperature adjusted to 50°C was brought into contact with the beaker. Once the beaker contents reached 50 °C, the sodium cocoyl glycinate (Amilite GCS-11 available from Ajinomoto OmniChem) was added to the beaker. The pH of the beaker contents was then adjusted to 7.5 with sodium hydroxide. The heat source was then removed from contact with the beaker. Once the beaker contents cooled to 35 °C, the additive noted in TABLE 7 was added to the contents of the beaker. Then the methylisothiazolinone (Neolone 950 available from The Dow Chemical Company) was added to the beaker to provide a body wash formulation having the composition noted in TABLE 6. The resulting body wash formulations were allowed to stand for two (2) days and observed for phase separation. The observations are provided in TABLE 7.

TABLE 6

Component	Conc. in Body Wash Formulation (wt%)
Cocamidopropyl betaine	7.00
Sodium cocoyl glycinate	3.00
Additive noted in TABLE 5	1.00*
sodium hydroxide	q.s.
Methylisothiazolinone	0.05
* -- unless otherwise noted in TABLE 5	

TABLE 7

Body Wash Formulation	Additive	Observation
Comp. Ex. F6	sodium hydroxypropyl starch*	Homogeneous
Comp. Ex. F7	CELLOSIZETM QP-100MH	Phase separated
Comp. Ex. F8	Prod. Comp. Ex. C4	Homogeneous
Comp. Ex. F9	Prod. Comp. Ex. C9	Homogeneous
Comp. Ex. F10	Prod. Comp. Ex. C10	Homogeneous
Ex. 20	Prod. Ex. 4	Homogeneous
Ex. 21	Prod. Ex. 8	Homogeneous
Ex. 22	Prod. Ex. 9	Homogeneous
Ex. 23	Prod. Ex. 10	Homogeneous
Ex. 24	Prod. Ex. 12	Homogeneous
Ex. 25	Prod. Ex. 13	Homogeneous
* -- sodium hydroxypropyl starch was loaded in formulation at 10 wt%		

Flash Foam

[0058] Flash foam measurements were obtained using an Oster® 16-Speed Blender Model No. 6878-042 and a 1,000 mL graduated cylinder according to the following procedure. A sample (5 g) of the body wash formulation noted in TABLE 8 and water (145 g) were added to the blender. The blender contents were subjected to blending at the "Grate" setting for ten seconds. The blender contents were then poured into the 1,000 mL graduated cylinder. The initial height of the foam, H0, was recorded. After two minutes, the initial liquid level, L0, was recorded. The flash foam volume reported in TABLE 8 was determined using the following equation

$$\text{Flash Foam (in mL)} = H0 - L0.$$

[0059] The viscosity of the body wash formulation was measured using a TA Instruments DHR-3 rheometer at 25 °C, equipped with a stainless steel 60 mm, 0.5° cone and plate sensor, and a gap set at 17 microns. The results are provided in TABLE 8.

TABLE 8

Body Wash Formulation	Base HEC Weight Average Molecular Weight, MW	Flash Foam (mL)	Viscosity at 6.31 s⁻¹ (mPa·s)
Comp. Ex. F1	--	540 ± 31	7,186
Comp. Ex. F4	380,000	--	1,432
Comp. Ex. F6	--	211 ± 48	3,550
Comp. Ex. F8	1,600,000	--	1,823
Comp. Ex. F9	380,000	--	231
Comp. Ex. F10	--	262 ± 35	2,927
Control¹	--	430 ± 26	7,692
Ex. 14	1,600,000	442 ± 55	13,600
Ex. 15	1,600,000	417 ± 89	5,844
Ex. 16	900,000	--	6,927
Ex. 17	1,600,000	--	4,290
Ex. 18	1,400,000	--	523
Ex. 19	1,400,000	--	8,098
Ex. 20	1,600,000	257 ± 30	5,828
Ex. 21	1,600,000	216 ± 18	5,863
Ex. 22	900,000	--	2,170
Ex. 23	1,600,000	--	5,309
Ex. 24	1,400,000	--	2,772
Ex. 25	1,400,000	--	5,037
1 -- Dove Deep Moisture body wash commercially available from Unilever.			

We Claim:

1. A personal care composition, comprising:
 - a vehicle;
 - a surfactant;
 - a water-soluble cellulose ether base material substituted with a hydrophobic group having a carbon chain with 8 to 18 carbon atoms;
 - wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of > 800,000 Daltons; and,
 - wherein the personal care composition contains < 0.01 wt% alkyl sulfate and < 0.01 wt% alkyl ether sulfate.
2. The personal care composition of claim 1, wherein the personal care composition is a body wash formulation; wherein the water-soluble cellulose ether base material is hydroxyethyl cellulose; and wherein the hydrophobic group is a C8-16 linear or branched alkyl group bonded to the water-soluble cellulose ether base material.
3. The body wash formulation of claim 2, wherein the hydrophobic group is bonded to the water-soluble cellulose ether base material through an ether linkage or an ether linkage and a 2-hydroxypropyl group.
4. The body wash formulation of claim 3, further comprising a pH adjusting agent.
5. The body wash formulation of claim 3, further comprising a biocide.
6. The body wash formulation of claim 3, further comprising a fragrance.
7. The body wash formulation of claim 3, further comprising a colorant.
8. The body wash formulation of claim 3, further comprising a soap.
9. The body wash formulation of claim 3, wherein the surfactant includes at least one glycinate; and wherein the water-soluble cellulose ether base material has a weight average molecular weight, MW, of 900,000 to 2,500,000 Daltons.
10. The body wash formulation of claim 9, further comprising at least one of: a pH adjusting agent, a biocide, a fragrance, a colorant and a soap.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/047279

A. CLASSIFICATION OF SUBJECT MATTER
INV. A61K8/73 A61Q19/10 A61K8/44
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)
A61K A61Q

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 practicable, search terms used)
EPO-Internal, WPI Data, CHEM ABS Data

C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	EP 1 191 039 A1 (KAO CORP [JP]) 27 March 2002 (2002-03-27) paragraphs [0001], [0005] - paragraphs [0028], [0103]; claims; examples 1,6,7,8,11; table 2 -----	1-8
X	DE 10 2007 015757 A1 (BEIERSDORF AG [DE]) 2 October 2008 (2008-10-02) paragraph [0030] - paragraph [0049]; claims; examples D-E,K paragraph [0108] -----	1-8
X	WO 2013/048778 A1 (ROHM & HAAS CHEMICALS LLC [US]; UNION CARBIDE CHEM PLASTIC [US]; WANG) 4 April 2013 (2013-04-04) page 2, line 21 - page 3, line 14; claims; table 3 ----- -/--	1-5

Further documents are listed in the continuation of Box C.

See patent family annex.

* Special categories of cited documents :

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Date of the actual completion of the international search 12 October 2017	Date of mailing of the international search report 23/10/2017
Name and mailing address of the ISA/ European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Fax: (+31-70) 340-3016	Authorized officer Loloiu, Teodora

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2017/047279

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT		
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A	JEN-TING LO ET AL: "Interaction between Hydrophobically Modified 2-Hydroxyethyl Cellulose and Sodium Dodecyl Sulfate Studied by Viscometry and Two-Dimensional NOE NMR Spectroscopy", JOURNAL OF PHYSICAL CHEMISTRY PART B: CONDENSED MATTER, MATERIALS, SURFACES, INTERFACES & BIOPHYSICAL, vol. 118, no. 24, 19 June 2014 (2014-06-19), pages 6922-6930, XP055413661, US ISSN: 1520-6106, DOI: 10.1021/jp500355n the whole document -----	1

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Information on patent family members

International application No

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