(19) World Intellectual Property Organization

International Bureau





PCT

(43) International Publication Date 8 January 2009 (08.01.2009)

(51) International Patent Classification:

 A61K 8/02 (2006.01)
 A61Q 5/06 (2006.01)

 A61K 8/73 (2006.01)
 A61Q 5/12 (2006.01)

(21) International Application Number:

PCT/US2008/068369

(22) International Filing Date: 26 June 2008 (26.06.2008)

(25) Filing Language: English

(26) Publication Language: English

(30) Priority Data:

60/947,128 29 June 2007 (29.06.2007) US

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(10) International Publication Number

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

Published:

with international search report



HAIR STYLING AND CONDITIONING PERSONAL CARE FILMS

Field

5 The present invention relates to personal care films for use in hair styling or conditioning.

Background

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Personal care films are an exciting development in the hair care industry. In theory, such films allow the styling power of an array of cans and bottles to be conveyed to a use site in a pocket-size package. At the use site, the film can be wetted with an aqueous liquid to form a wet product that can then be applied to the hair. Unfortunately, this unparalleled portability has never been optimized.

Currently available styling films demonstrate poor "in hands" properties. For example, they can be tacky, as a result of a combination of polymers and plasticizers present. Similarly, some currently available styling films do not dissolve fast enough and therefore can feel grainy, lumpy, or stringy as a result of relatively long disintegration times.

The success of a cosmetic, including personal care films, depends in great measure on the way it feels to a user at the time of use. Remedying the current drawbacks to currently available films is not straightforward. For example, merely removing tack-causing ingredients is not an option as it may destroy the film's styling/fixative performance and/or negatively impact its mechanical properties. Similarly, lowering disintegration times can result in a wet product that runs through the fingers instead of being appropriately viscous and may actually increase tackiness.

Another challenge in the styling industry is to create products that do not flake off the hair upon drying. Combing, or in some cases just touching, the hair can result in the appearance of flakes that can look like dandruff and are unsightly. For dissolvable styling films, the concentrated product form results in gels with relatively higher content of film-forming materials which exacerbates flaking.

Furthermore, to optimize in-use properties and dissolution time, formulators of dissolvable films often have to leave out or reduce the amount of the "traditional" conditioning agents, such as, for example, polymers, cationic surfactants, and/or silicones. This approach yields styling products that do not offer *in-situ* conditioning, exhibited as, for example, improved hair feel, comb, and/or mitigation of electrostatic fly-away. Conditioning is highly desirable to the consumer and greatly contributes to the overall grooming experience.

Thus, what is needed are new types of personal care films with better in hands properties and improved multifunctional performance on hair.

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Summary

In one embodiment, the present invention provides personal care dissolvable films, comprising greater than about 30 weight percent water-soluble chitosan derivative and a cosmetically acceptable plasticizer.

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Detailed Description

In one embodiment, the present invention provides a personal care dissolvable film, comprising greater than about 30 weight percent water-soluble chitosan derivative, and a cosmetically acceptable plasticizer.

"Weight percent" refers to the weight of the component in a theoretical

completely dried film, in other words, as if the film had been dried until only nonvolatile components remained. Thus, for this application, 30 weight percent is independent of humidity.

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The water-soluble chitosan derivative includes anionic, cationic, amphoteric or nonionic chitosan polymers. In one embodiment, the water-soluble chitosan derivative is a chitosan salt of pyrrolidone carboxylic acid, which is, for example available under the trade name of KYTAMER PC from The Dow Chemical Company. Chitosan PCA salt is known for its moisturization properties, in fact, it has been described by some as a film plasticizer. Applicants have unexpectedly found that films of this invention, i.e., those comprising greater than about 30 weight percent water-soluble chitosan derivative, actually exhibit superior overall performance as opposed to currently available films, as will be discussed in the Examples section. Thus, in one embodiment, the present invention provides a personal care dissolvable film wherein the water-soluble chitosan derivative is the main ingredient.

In one embodiment, the water-soluble chitosan derivative is present in the personal care dissolvable film in a range from about 30 weight percent to about 99.99 weight percent. In one embodiment, the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 60 weight percent. In one embodiment, the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 50 weight percent. In one embodiment, the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 40 weight percent. The recited ranges are given to amply illustrate certain features of the invention; however, additional ranges are understood to be contemplated. Indeed, all novel combinations and subcombinations found within the above ranges are contemplated and may be placed in the appended claims.

The plasticizers include any of the plasticizers listed in McCutcheon's Functional Materials (1992). Preferably, the plasticizer is lipid, a polyol, an acid, a polyester, or water-soluble organopolysiloxane.

Examples of lipid plasticizers include waxes (such as ethoxylated jojoba or beeswax), mineral oils, paraffin derivatives, vegetable oils, triglycerides, lanolins, unsaturated fatty acids, and their derivatives.

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Examples of polyol plasticizers include glycerin, ethylene glycol, propylene glycol, sugar alcohols (such as sorbitol, SORBETH-30, manitol, maltitol, lactitol), saccharides (such as fructose, glucose, sucrose, maltose, lactose, and high fructose corn syrup), polysaccharides, ascorbic acid, decyl glucoside, propylene glycol, polyethylene glycol, PEG derivatives (ether, ester), and dimethicone copolyols (such as PEG-12 dimethicone, PEG/PPG-18/18 dimethicone, and PPG-12 dimethicone).

Examples of acid plasticizers include carboxylic acids (such as citric acid, maleic acid, succinic acid, adipic acid, azelaic acid, benzoic acid, dimer acids, fumaric acid, isobutyric acid, isophthalic acid, lauric acid, linoleic acid, maleic acid, maleic anyhydride, melissic acid, myristic acid, oleic acid, palmitic acid, phosphoric acid, phthalic acid, ricinoleic acid, sebacic acid, stearic acid, succinic acid, 1,2-benzenedicarboxylic acidpolyacrylic acid, and polymaleic acid), alpha and beta hydroxy acids (such as glycolic acid, lactic acid (including sodium, ammonium, and potassium salts), and salicylic acid), and sulfonic acid derivatives.

Examples of polyester plasticizers include glycerol triacetate, acetylated-monoglyceride, diethyl phthalate, triethylcitratetriethyl citrate, tributyl citrate, acetyl triethyl citrate, acetyl triethylcitrate, diisobutyl adipate, butyl stearate, and phtalates.

In one embodiment, the plasticizer includes a mixture of at least two plasticizers.

In one embodiment, the plasticizer includes at least two of small molecule polyol, polyethylene glycol derivative of dimethicone, and alkyl glucoside.

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The plasticizer is present in an amount from about 0.01 to about 30 weight percent, that is, by the weight of the plasticizer in a theoretical completely dried film (as if the film had been dried until only nonvolatile components remained). In a preferred embodiment, the plasticizer is present in an amount from about 5 to about 25 weight percent. In one embodiment, the plasticizer is present in an amount from about 10 to about 20 weight percent. In one embodiment, the plasticizer is present in an amount of about 16 weight percent.

In some embodiments, personal care dissolvable films of the present invention further comprise a water soluble film forming agent. In one embodiment, the water soluble film forming agent is a pullulan. In one embodiment, the water soluble film forming agent is a cellulose ether based polymer. In one embodiment, the water soluble film forming agent is at least one of methylcellulose, hydroxypropyl methylcellulose, hydroxyethyl cellulose, cationic hydroxyethyl cellulose, hydrophobically modified hydroxyethyl cellulose, or cationic hydrophobically modified hydroxyethyl cellulose. In a preferred embodiment, the water soluble film forming agent is hydroxypropyl methylcellulose.

When present, the water soluble film forming agent is present in an amount from about 0.1 weight percent to about 69.99 weight percent of the dry film. In one embodiment, the water soluble film forming agent is present in an amount from about 5 weight percent to about 60 weight percent. In one embodiment, the water soluble film forming agent is present in an amount from about 1 weight percent to about 10 weight percent.

In one embodiment, the personal care dissolvable film contains less than 0.5% of

modified starch.

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Composition of the present invention can further incorporate other ingredients known in the art of hair care formulations and dissolvable films. Other optional ingredients for personal care compositions of the present invention include at least one of the following: additional film forming agents, cosmetically acceptable emollients, moisturizers, conditioners, oils, sunscreens, surfactants, emulsifiers, preservatives, rheology modifiers, colorants, preservatives, pH adjustors, propellants, reducing agents, fragrances, foaming or de-foaming agents, tanning agents, depilatory agents, flavors, astringents, antiseptics, deodorants, antiperspirants, insect repellants, bleaches, lighteners, anti-dandruff agents, adhesives, polishes, strengtheners, fillers, barrier materials, or biocides.

The moisturizers include 2-pyrrolidone-5-carboxylic acid and its salts and esters, alkyl glucose alkoxylates or their esters, fatty alcohols, fatty esters, glycols and, in particular, methyl glucose ethoxylates or propoxylates and their stearate esters, isopropyl myristate, lanolin or cetyl alcohols, aloe, silicones, and polyols, such as, for example, propylene glycol, glycerol and sorbitol.

Conditioners include stearalkonium chloride, dicetyldimonium chloride, lauryl methyl gluceth-10 hydroxypropyldimonium chloride, and natural and synthetic conditioning polymers such as polyquaternium-4, polyquaternium-7, polyquaternium-10, polyquaternium-24, polyquaternium-67 and the like, chitosan and derivatives thereof.

Examples of oils include hydrocarbon-based oils of animal origin, such as squalene, hydrocarbon-based oils of plant origin, such as liquid triglycerides of fatty acids comprising from 4 to 10 carbon atoms, for instance heptanoic or octanoic acid triglycerides, or alternatively, oils of plant origin, for example sunflower oil, corn oil, soybean oil, marrow oil, grapeseed oil, sesame seed oil, hazelnut oil, apricot oil,

macadamia oil, arara oil, coriander oil, castor oil, avocado oil, jojoba oil, shea butter oil, or caprylic/capric acid triglycerides, MIGLYOL 810, 812 and 818 (from Dynamit Nobel), synthetic esters and ethers, especially of fatty acids, for instance the oils of formulae R¹COOR² and R¹OR² in which R¹ represents a fatty acid residue comprising from 8 to 29 carbon atoms and R² represents a branched or unbranched hydrocarbon-5 based chain comprising from 3 to 30 carbon atoms, for instance purcellin oil, isononyl isononanoate, isopropyl myristate, 2-ethylhexyl palmitate, 2-octyldodecyl stearate, 2octyldodecyl erucate or isostearyl isostearate, hydroxylated esters, for instance isostearyl lactate, octyl hydroxystearate, octyldodecyl hydroxystearate, diisostearyl malate, 10 triisocetyl citrate and fatty alcohol heptanoates, octanoates and decanoates, polyol esters, for instance propylene glycol dioctanoate, neopentyl glycol diheptanoate and diethylene glycol diisononanoate, pentaerythritol esters, for instance pentaerythrityl tetraisostearate, lipophilic derivatives of amino acids, such as isopropyl lauroyl sarcosinate, such as is sold under the name ELDEW SL 205 (from Ajinomoto), linear or branched 15 hydrocarbons of mineral or synthetic origin, such as mineral oils (mixtures of petroleumderived hydrocarbon-based oils), volatile or non-volatile liquid paraffins, and derivatives thereof, petroleum jelly, polydecenes, isohexadecane, isododecane, hydrogenated isoparaffin (or polyisobutene), silicone oils, for instance volatile or non-volatile polymethylsiloxanes (PDMS) comprising a linear or cyclic silicone chain, which are liquid or pasty at room temperature, especially cyclopolydimethylsiloxanes (cyclomethicones) such as cyclopentasiloxane and cyclohexadimethylsiloxane, polydimethylsiloxanes comprising alkyl, alkoxy or phenyl groups, which are pendent or at the end of a silicone chain, these groups comprising from 2 to 24 carbon atoms, phenyl silicones, for instance phenyl trimethicones, phenyl dimethicones,

25 phenyltrimethylsiloxydiphenylsiloxanes, diphenyl dimethicones,

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diphenylmethyldiphenyltrisiloxanes 2-phenylethyltrimethyl siloxysilicates and polymethylphenylsiloxanes, fluoro oils such as partially hydrocarbon-based and/or partially silicone-based fluoro oils, ethers such as dicaprylyl ether (CTFA name: dicaprylyl ether), and C_{12} - C_{15} fatty alcohol benzoates (FINSOLV TN from Finetex), mixtures thereof.

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Oils include mineral oil, lanolin oil, coconut oil and derivatives thereof, cocoa butter, olive oil, almond oil, macadamia nut oil, aloe extracts such as aloe vera lipoquinone, jojoba oils, safflower oil, corn oil, liquid lanolin, cottonseed oil, peanut oil, hydrogenated vegetable oil, squalane, castor oil, polybutene, sweet almond oil, avocado oil, calophyllum oil, ricin oil, vitamin E acetate, olive oil, silicone oils such as dimethylopolysiloxane and cyclomethicone, linolenic alcohol, oleyl alcohol, and the oil of cereal germs.

Other suitable emollients include dicaprylyl ether, C₁₂₋₁₅ alkyl benzoate, DC 200 FLUID 350 silicone fluid (from Dow Corning Corp.), isopropyl palmitate, octyl palmitate, isopropyl myristate, hexadecyl stearate, butyl stearate, decyl oleate, acetyl glycerides, the octanoates and benzoates of C₁₂₋₁₅ alcohols, the octanoates and decanoates of alcohols and polyalcohols such as those of glycol and glyceryl, ricinoleates esters such as isopropyl adipate, hexyl laurate and octyl dodecanoate, dicaprylyl maleate, phenyltrimethicone, and aloe vera extract. Solid or semi-solid cosmetic emollients include glyceryl dilaurate, hydrogenated lanolin, hydroxylated lanolin, acetylated lanolin, petrolatum, isopropyl lanolate, butyl myristate, cetyl myristate, myristyl myristate, myristyl lactate, cetyl alcohol, isostearyl alcohol and isocetyl lanolate.

Dyes include water-soluble dyes such as copper sulfate, iron sulfate, watersoluble sulfopolyesters, rhodamines, natural dyes, for instance carotene and beetroot juice, methylene blue, caramel, the disodium salt of tartrazine and the disodium salt of

fuschin, and mixtures thereof. Liposoluble dyes from the list above may also optionally be used.

Preservatives include alcohols, aldehydes, methylchloroisothiazolinone and methylisothiazolinone, p-hydroxybenzoates, and in particular methylparaben, propylparaben, glutaraldehyde and ethyl alcohol.

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The pH adjustors, include inorganic and organic acids and bases and in particular aqueous ammonia, citric acid, phosphoric acid, acetic acid, and sodium hydroxide.

Reducing agents include ammonium thioglycolate, hydroquinone and sodium thioglycolate.

Fragrances may be aldehydes, ketones, or oils obtained by extraction of natural substances or synthetically produced as described above. Often, fragrances are accompanied by auxiliary materials, such as fixatives, extenders, stabilizers and solvents.

Biocides include antimicrobials, bactericides, fungicides, algaecides, mildicides, disinfectants, antiseptics, and insecticides.

The amount of optional ingredients effective for achieving the desired property provided by such ingredients can be readily determined by one skilled in the art.

Examples

The following examples are for illustrative purposes only and are not intended to limit the scope of the present invention. All percentages are by weight unless otherwise specified.

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Example 1

Exemplary personal care compositions contain the components recited in TABLE

1.

TABLE 1

#	Component	Batch 1	Batch 2	Batch 3	Batch 4	Batch 5
1	METHOCEL E3 hydroxypropyl methylcellulose		1.80	2.40	3.50	4.80
2	KYTAMER PC chitosan/pyrrolidone carboxylic acid salt (5% soln)	97.56	84.00	72.00	70.00	64.00
	(Wt. % (in final dry film))	(82 wt %)	(56 wt %)	(48 wt %)	(40 wt %)	(32 wt %)
3	Glycerin	0.24	0.30	0.30	0.35	0.40
4	DC-193 PEG-12 Dimethicone	0.24	0.30	0.30	0.35	0.40
5	Citric Acid (10% soln)	0.98	2.40	2.40	2.80	3.50
6	PLANTAREN 2000 Decyl Glucoside (50% solution)	0.98	1.20	1.20	1.40	1.60
7	Deionized Water		10.00	21.40	21.60	25.30

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All numerals without parentheses are in grams.

The parentheticals are provided to illustrate that the films in TABLE 1 all have more than 30 weight percent chitosan/pyrrolidone carboxylic acid salt in the resulting dry film. Though provided for convenience, the weight percent can also be calculated by

dividing the weight of the dry KYTAMER PC component (a) by the total weight of the remaining non-volatile ingredients in the film pre-mix (b) where:

(a) = 5% of the total weight of the KYTAMER PC aqueous solution used, and

(b) = Weight (1) + (a) + Weight (3) + Weight (4) +
$$0.1*$$
 Weight (5) + $0.5*$

5 Weight (6)

The ingredients are combined into a liquid pre-mix formulation for each batch.

Three drops of GLYDANT preservative were added at the end to each formulation. The liquid pre-mix formulations are then cast by drawing down and drying overnight at room temperature to afford 2 mils thick films.

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Example 2 (Comparative)

Comparative compositions contain the components recited in TABLE 2.

TABLE 2

Component	Comparative Sample A	Comparative Sample B
METHOCEL E3 hydroxypropyl methylcellulose	10.00	
CELLOSIZE QP 09L hydroxyethyl cellulose		10.00
KYTAMER PC chitosan/pyrrolidone carboxylic acid salt (5% soln)		-
Glycerin	0.50	0.50
DC-193 PEG-12 Dimethicone	0.50	0.50
Citric Acid (10% soln)	4.0	4.0
PLANTAREN 2000 Decyl Glucoside (50% solution)	2.00	2.00
Deionized Water	83.0	83.0

All numerals are in grams. The ingredients are combined into a liquid pre-mix formulation for each sample. Three drops of GLYDANT preservative were added at the

end to each formulation. The liquid pre-mix formulations are then cast by drawing down and drying overnight at room temperature to afford 2 mils thick films.

Example 3

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In-situ formulations were prepared by dissolving 0.3 g of dry film (made substantially according to the protocol of Example 1 and Example 2, and representing Batch 1, Batch 4, Comparative Sample A, and Comparative Sample B) in 12 g of water and stirring until complete dissolution was achieved. Then, 0.4 g of the *in situ* gel was applied to pre-wetted, pre-combed, eight inch long, ~ 4.5 g tresses of European virgin brown hair using a pipette in small portions, evenly from top (swatched end) to bottom (hair tips). The gel was then worked into the hair with fingers going from top to bottom of each tress five times. The tress was then reversed and the same procedure was repeated five more times. At the end, the tress was combed to eliminate knots, smoothed with fingers, and hung to dry overnight.

The next day, the tresses were visually inspected and felt for stiffness. Tresses treated with Comparative Sample A and Comparative Sample B were flexible with little-to-no stiffness, for example, when their central portions were placed on a support beam, the unsupported ends drooped down. In stark contrast, tresses treated with gels corresponding Batch 1 and Batch 4 felt rigid, and did not bend at all when their central portions were placed on a support beam, indicating excellent hair stiffening and styling performance.

The tresses were also visually inspected for flake. Each tress was held at the swatched end with one (left) hand, and a fingernail was forcefully run down the length of the tress (right hand). After inspection, tresses treated with film formulations Batch 1, Batch 4, and Comparative Sample A did not reveal any flakes, while the tress treated

with Comparative Sample B showed excessive flaking (dandruff-like specs resulted from formulation coming off from hair).

Example 4

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A film made substantially according to the protocol of Example 1 representing Batch 5 was compared to a commercially available styling film product, OSIS SHOCKFROSTER hair styling strips (modified corn starch, PVP, water, propylene glycol, octylacrylamide/acrylates/butylaminoethyl/methacrylate copolymer, aminomethyl propanol, aluminum starch octenylsuccinate, fragrance, benzyl salicylate, limonene, butylphenyl methylproprional, linalool, and Red 40).

In-situ formulations were prepared by dissolving 0.3 g of dry film in 12 g of water and stirring until complete dissolution was achieved. Then, 0.4 g of the *in situ* gel was applied to pre-wetted, pre-combed, eight inch long, ~ 4.5 g tresses of European virgin brown hair using a pipette in small portions, evenly from top (swatched end) to bottom (hair tips). The gel was then worked into the hair with fingers going from top to bottom of each tress five times. The tress was then reversed and the same procedure was repeated five more times. At the end, the tress was combed to eliminate knots, smoothed with fingers, and hung to dry overnight.

The treated hair tresses were distributed in pairs to six expert panelists trained to evaluate performance of cosmetic products on hair. Each panelist evaluated two pairs of tresses, one tress treated with Batch 5 versus one OSIS SHOCKFROSTER hair styling strip control in each pair. The panelists were asked to pick one tress that was more rigid/stiff, combed easier, showed more flaking, felt softer/smoother, combed easier, and had more static flyaways.

The evaluation procedures for each of these properties are as follows:

Stiffness: Tresses were gently handled and "felt" for differences in stiffness. Using two fingers, the middles of the swatches were held in a horizontal position to determine which one was bending more than the other. The more rigid one was noted.

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<u>Dry comb:</u> The ease of combing was evaluated. The one tress that combed more easily was noted.

<u>Flake</u>: The tress was held at the bound end with one (left) hand, and a fingernail was forcefully run down the length of the tress (right hand). After inspection of both tresses, the one with more flaking was noted.

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Feel: The tress that felt silkier/softer/smoother was noted.

Static flyaway's: Each tress was combed at least 5 times and the amounts of flyaway's generated each time were compared. The tress that generated more flyaway's was noted.

The subjective evaluations were statistically analyzed to identify differences at above 85 % confidence level. The findings showed that that Batch 5 was superior to the OSIS SHOCKFROSTER hair styling strip control. For example, Batch 5 significantly outperformed the OSIS SHOCKFROSTER hair styling strips based on dry combability (12/12), flaking (2/12 (indicating less)), and feel (12/12). Stiffness and static flyaway properties were statistically no different between Batch 5 and the OSIS

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SHOCKFROSTER hair styling strips at the chosen confidence level.

Example 5

A film made substantially according to the protocol of Example 1 representing Batch 5 was compared to a commercially available styling film product, AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strips (pullulan, modified

corn starch, glycerin, camellia oleifera leaf extract, aloe barbadensis leaf extract, linseed extract, hydrolyzed wheat protein, hydrolyzed wheat starch, caprylic/capric triglyceride, fragrance, limonene, linalool, geraniol, eugenol, citronellol, amyl cinnamal, benzyl benzoate, citral, benzyl salicylate, and farnesol).

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In-situ formulations were prepared by dissolving 0.3 g of dry film in 12 g of water and stirring until complete dissolution was achieved. Then, 0.4 g of the *in situ* gel was applied to pre-wetted, pre-combed, eight inch long, ~ 4.5 g tresses of European virgin brown hair using a pipette in small portions, evenly from top (swatched end) to bottom (hair tips). The gel was then worked into the hair with fingers going from top to bottom of each tress five times. The tress was then reversed and the same procedure was repeated five more times. At the end, the tress was combed to eliminate knots, smoothed with fingers, and hung to dry overnight.

The treated hair tresses were distributed in pairs to six expert panelists trained to evaluate performance of cosmetic products on hair. Each panelist evaluated two pairs of tresses, one tress treated with Batch 5 versus one AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strip control in each pair. The panelists were asked to pick one tress that was more rigid/stiff, combed easier, showed more flaking, felt softer/smoother, combed easier, and had more static flyaways, using the evaluation procedures recited in Example 4.

The subjective evaluations were statistically analyzed to identify differences at above 85 % confidence level. The findings showed that Batch 5 was superior to the AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strip control in all respects. For example, Batch 5 significantly outperformed the AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strips based on stiffness (11/12), dry

combability (12/12), flaking (1/12 (indicating less)), feel (12/12), and static flyaway (2/12 (indicating less)).

Example 6

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Tactile properties are fundamentally important to consumer preferences. A subjective in-hands study was conducted to compare in-hands properties of films made substantially according to the protocol of Example 1 representing Batch 5 to commercially available OSIS SHOCKFROSTER hair styling strips (see Example 4 for ingredients). Ten panelists participated in this study, with each panelist being asked to compare the ease of dissolution and in-hands tackiness of the respective films and choose one that dissolved faster/easier and felt less tacky. Seven out of ten panelists concluded that the film of the present invention (Batch 5) was easier to dissolve, and eight of ten believed that Batch 5 felt less tacky than the commercial control.

15 Example 7

Films made substantially according to the protocol of Example 1 representing Batch 1, Batch 3, and Batch 5 were compared to commercially available OSIS SHOCKFROSTER hair styling strips (see Example 4 for ingredients) for dissolution rates.

Dissolution rates were measured using the Hand Rubbed Dissolution Test that simulates real-life usage conditions. A 2cm x 3cm piece of dissolvable film is placed in the palm of the operator's left hand. 2 ml of water are added and the operator rubs the film with the water using two fingers of the right hand in a circular motion (each circle taking approximately one second) until the film is completely dissolved. The dissolution times (average of two measurements) are determined.

Using the Hand Rubbed Dissolution Test, Batch 3 dissolved in 6 (±2) seconds; Batch 5 dissolved in 7 (±2) seconds; Batch 1 dissolved in 10 (±2) seconds; and the OSIS SHOCKFROSTER hair styling strips dissolved in 15 (±2) seconds. Thus, films of the present invention performed significantly better. Any improvement in dissolution time is important, as a relatively faster dissolution time relates to positive consumer experience.

Example 8

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Viscosity is yet another important tactile property to consumers. Viscosities were determined for films made substantially according to the protocol of Example 1 (representing Batch 1 and Batch 3), and Example 2 (representing Comparative Sample A and Comparative Sample B), as well as commercially available OSIS SHOCKFROSTER hair styling strips (see Example 4 for ingredients), AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strips (see Example 5 for ingredients), and SMART H₂O STYLING STRIPS hair styling strips (PVP, modified corn starch, fragrance phenoxyethanol, dimethicone, amodimethicone, methylparaben, C12-14 SEC Pareth 7, C12-14 SEC Pareth 5, ethylparaben, butylparaben, laureth-4, laureth-23, and isobutylparaben).

Viscosities of the *in situ* gels (prepared by dissolving 0.3 g of dry film in 12 g of water and stirring until complete dissolution) were measured. The comparative formulations (Sample A and Sample B) and all three commercial products gave water-thin *in situ* gels, which is inconvenient to the consumer and may lead to the loss of some product while still in hands before it gets applied to the hair. In contrast, films of the invention (Batch 1 and Batch 3) resulted in thicker creamier gels, which are closer to the conventional non-film styling gel products and easier to handle and apply. The

viscosities measured using a Brookfield viscometer, model LVDVII+, spindle # 60 at 60 rpm and 22°C are listed in TABLE 3.

TABLE 3

Film	Viscosity (cps)
Batch 1	117
Batch 3	71
Sample A (comparative)	4
Sample B (comparative)	9.5
OSIS SHOCKFROSTER hair styling strips (comparative)	7
AVEDA CONTROL TAPE EXTREME STYLE STRIPS hair styling strips (comparative)	4.5
SMART H ₂ O STYLING STRIPS hair styling strips (comparative)	8.5

It is understood that the present invention is not limited to the embodiments specifically disclosed and exemplified herein. Various modifications of the invention will be apparent to those skilled in the art. Such changes and modifications may be made without departing from the scope of the appended claims.

Moreover, each recited range includes all combinations and subcombinations of ranges, as well as specific numerals contained therein. Additionally, the disclosures of each patent, patent application, and publication cited or described in this document are hereby incorporated herein by reference, in their entireties.

Claims:

1. A personal care dissolvable film, comprising:

greater than about 30 weight percent water-soluble chitosan derivative; and a cosmetically acceptable plasticizer.

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- 2. The personal care dissolvable film of claim 1, further comprising a water soluble film forming agent.
- 3. The personal care dissolvable film of claim 2, wherein the water soluble film formingagent is a pullulan.
 - 4. The personal care dissolvable film of claim 2, wherein the water soluble film forming agent is a cellulose ether based polymer.
- 5. The personal care dissolvable film of claim 2, wherein the water soluble film forming agent is at least one of methylcellulose, hydroxypropyl methylcellulose, hydroxyethyl cellulose, cationic hydroxyethyl cellulose, hydrophobically modified hydroxyethyl cellulose, or cationic hydrophobically modified hydroxyethyl cellulose.
- 20 6. The personal care dissolvable film of claim 2, wherein the water soluble film forming agent is hydroxypropyl methylcellulose.
 - 7. The personal care dissolvable film of claim 2, wherein the water soluble film forming agent is present in an amount from about 0.1% to about 69.99% by weight of the dry
- 25 film.

8. The personal care dissolvable film of claim 1, wherein the water-soluble chitosan derivative is a chitosan salt of pyrrolidone carboxylic acid.

- 9. The personal care dissolvable film of claim 1, wherein the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 99.99 weight percent.
- 10. The personal care dissolvable film of claim 1, wherein the water-soluble chitosanderivative is present in a range from about 30 weight percent to about 60 weight percent.
 - 11. The personal care dissolvable film of claim 1, wherein the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 50 weight percent.
- 15 12. The personal care dissolvable film of claim 1, wherein the water-soluble chitosan derivative is present in a range from about 30 weight percent to about 40 weight percent.
 - 13. The personal care dissolvable film of claim 1, wherein the plasticizer includes at least one of a lipid, a polyol, an acid, a polyester, or water-soluble organopolysiloxane.
 - 14. The personal care dissolvable film of claim 1, wherein the plasticizer is a polyol plasticizer.

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15. The personal care dissolvable film of claim 1, wherein the plasticizer is present in anamount from about 0.01 to about 30 weight percent.

16. The personal care dissolvable film of claim 1, wherein the film contains less than 0.5% modified starch.

- 17. The personal care dissolvable film of claim 1, further comprising at least one of cosmetically acceptable additional film forming agents, emollients, moisturizers, conditioners, oils, sunscreens, surfactants, emulsifiers, preservatives, rheology modifiers, colorants, preservatives, pH adjustors, propellants, reducing agents, fragrances, foaming or de-foaming agents, tanning agents, depilatory agents, flavors, astringents, antiseptics, deodorants, antiperspirants, insect repellants, bleaches, lighteners, anti-dandruff agents, adhesives, polishes, strengtheners, fillers, barrier materials, or biocides, or an active ingredient selected from skin care actives, nail care actives, or hair care actives.
 - 18. A gel, comprising:
- the personal care dissolvable film of claim 1; and an aqueous component present in an amount sufficient to dissolve the film.
 - 19. The gel of claim 18, wherein the gel has ratio of dry film to water in a range from about 1:10 to about 1:50.

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- 20. The gel of claim 18, wherein the gel has ratio of dry film to water of about 1:40.
- 21. The gel of claim 18, wherein the gel as a Brookfield viscosity of greater than 15 cps at 25°C.

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22. The gel of claim 18, wherein the gel has a Brookfield viscosity of greater than 50 at 25°C.

INTERNATIONAL SEARCH REPORT

International application No PCT/US2008/068369

A. CLASSIFICATION OF SUBJECT MATTER INV. A61K8/02 A61K8/73

1K8/73 A610

A61Q5/06

A61Q5/12

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) $A61K \quad A61Q$

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 practical, search terms used)

EPO-Internal, WPI Data

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•	claims 1,28,31,32; examples 2,4,7,8	
X	WO 2005/070376 A (WELLA AG ET AL.) 4 August 2005 (2005-08-04)	1-3,7-18
	page 1, line 5 - line 10 page 4, line 16 - line 31	
	page 12, line 6 - line 7 page 12, line 16 - line 19	•
	page 13, line 5 - line 17 page 14, line 13 - line 17	
	page 18, line 1 - line 31	
	page 20, line 12 - line 20 examples 2-5	
γ	claims 17,18,21,25,26	4.6
		4–6
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X Further documents are listed in the continuation of Box C.	X See patent family annex.
* Special categories of cited documents: "A" document defining the general state of the art which is not considered to be of particular relevance "E" earlier document but published on or after the international filing date "L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified) "O" document referring to an oral disclosure, use, exhibition or other means "P" document published prior to the international filing date but later than the priority date claimed	 "T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention "X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone "Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art. "&" document member of the same patent family
Date of the actual completion of the international search 15 October 2008	Date of mailing of the international search report 23/10/2008
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 Alvarez Alvarez, C

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2008/068369

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INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No
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