



- (51) **International Patent Classification:**
A61K 8/49 (2006.01) *A61Q 5/06* (2006.01)
- (21) **International Application Number:**
PCT/US2016/037959
- (22) **International Filing Date:**
17 June 2016 (17.06.2016)
- (25) **Filing Language:** English
- (26) **Publication Language:** English
- (30) **Priority Data:**
62/181,488 18 June 2015 (18.06.2015) US
- (71) **Applicant:** THE PROCTER & GAMBLE COMPANY
[US/US]; One Procter & Gamble Plaza, Cincinnati, Ohio
45202 (US).
- (72) **Inventors:** WASHINGTON, Randy, Purnell; 5062 Hunt-
ington Circle, West Chester, Ohio 45069 (US). FLOHR,
Andreas; Procter & Gamble Service GmbH, Berliner Allee
65, 64274 Darmstadt (DE).
- (74) **Agent:** KREBS, Jay A.; c/o THE PROCTER & GAMBLE
COMPANY, Global Patent Services, One Procter &
Gamble Plaza, C8-229, Cincinnati, Ohio 45202 (US).

- (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, DK, DM, DO, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IR, IS, JP, KE, KG, KN, KP, KR, 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, TR), OAPI (BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, KM, ML, MR, NE, SN, TD, TG).

Published:

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

(54) **Title:** SHAPING KERATIN FIBRES USING 2-OXAZOLIDINONE COMPOUNDS

(57) **Abstract:** Described herein is a method for shaping keratin fibres including (i) providing a crosslinking composition, wherein the crosslinking composition includes a 2-oxazolidinone compound, a photocatalyst being a hydroxy-substituted aromatic compound, and a cosmetically acceptable carrier; (ii) applying the crosslinking composition to keratin fibres; and (iii) mechanically shaping the keratin fibres with an appliance at a temperature of from about 80C to about 180C while exposing the crosslinking composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm.



WO 2016/205578 A1

SHAPING KERATIN FIBRES USING 2-OXAZOLIDINONE COMPOUNDS

FIELD OF THE INVENTION

Described herein is a method for shaping keratin fibres including providing a crosslinking
5 composition, wherein the crosslinking composition includes a 2-oxazolidinone compound and a
photocatalyst, applying the crosslinking composition to the hair, and exposing the crosslinking
composition to electromagnetic radiation.

BACKGROUND OF THE INVENTION

10 Consumers are constantly demanding products that meet their daily styling and
conditioning needs without damaging the hair. The perceived and sometimes real impact of
various treatments and the implements such as blow drying and flat iron can have detrimental
effects on the acute and chronic nature of hair.

Permanent methods – or relaxers – usually comprise the steps of applying onto hair a
15 composition comprising a high pH solution (or combination of components to generate high pH),
leaving on for a protracted time and then applying a neutralizing composition. A relaxer is a
treatment predominately used by people of African-descent to permanently straighten hair. The
treatment relies on either the one-step sodium hydroxide (lye) or a two step (e.g. guanidine
carbonate and calcium hydroxide) to achieve very high pH (pH 12-14).

20 Semi-permanent benefits can be achieved using redox chemistry such as thioglycolic acid
(TGA) and hydrogen peroxide. Here, the curly hair is transformed into the straight hair because
the disulfide bonds are broken by the reaction with TGA. The straighter style is locked in during
the oxidation step with hydrogen peroxide.

Non-permanent methods usually comprise the step of heating the hair with a flat-iron or
25 heating element. Methods using such devices in combination with chemically-modifying the
internal hair fibres can obtain long-lasting effects e.g. over several months. The Brazilian Keratin
Treatments (BKTs) enable the achievement of a straight hairstyle that lasts several months via a
shampoo treatment. The key active in BKTs is formaldehyde. The most efficacious treatments
(used mainly in salons) often rely on high temperature – usually 232°C (450°F) – with
30 formaldehyde. Hair treated with products with high concentration of formaldehyde such as
Brazilian Blowout delivers semi- permanent straight hair. Over time and following shampooing,
the hair reverts back to a curly configuration.

The known methods for straightening hair all have drawbacks. The permanent methods are typically time-consuming and may damage hair. In addition, such methods show little flexibility so that any need and/or wish for changing the hairstyle would require conducting again a "permanent" wave onto hair, which is time-consuming and further damages the hair.

5 Along with the high potential skin irritation during application, relaxers tend to permanently change the hair by breaking the natural disulfide bonds in the hair. This leaves the hair weaker and more prone for further breakage. Over-processing can also increase hair damage and skin irritation. Consumer products using redox chemistry to achieve semi-permanent benefits, but overprocessing the hair and the strong sulphur smell are concerns of technologies
10 based on reducing chemistry.

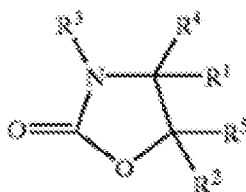
 According to the US National Toxicology Program, formaldehyde is known to be a human carcinogen. Therefore, providing a semi-permanent style with carcinogen-free formulation is paramount. Given the safety concern of formaldehyde and the potential damaging effect of relaxers and reducing chemistry, there is a need for a safe alternative to durable
15 straightening that does not break disulfide bonds.

 None of the above methods allow achieving a hairstyle that may be retained and/or recovered after at least one shampoo treatment without severely damaging the hair or using a carcinogenic active. There is a need for the provision of a method for achieving a hairstyle that lasts at least five shampoo treatments so that the user would not need to re-shape hair after each
20 shampoo but would still have the opportunity to change hairstyle after some time and without needing to use stringent or harsh hair treatment, such as the permanent technologies outlined above.

 Accordingly, there is a need therefore for providing a method for achieving and semi-permanently retaining and/or recovering hairstyle using actives that do not break disulfide bonds
25 (reducing agents) or that are considered carcinogenic. There is also the need for providing a method for obtaining a hairstyle exhibiting resistance to shampoo treatments. Particularly, there is a need for providing a method for retaining and/or recovering hairstyle after at least one shampoo treatment, particularly after five shampoo treatments, more particularly after ten shampoo treatments. In addition, there is a need for providing a method for achieving and
30 retaining and/or recovering hairstyle, without damaging hair. Also, there is a need for providing more economic semi-permanent hair straightening or hair relaxing treatments.

SUMMARY OF THE INVENTION

Described herein is a method for shaping keratin fibres comprising (a) providing a crosslinking composition, wherein the crosslinking composition comprises (i) a 2-oxazolidinone compound having the structure:



wherein:

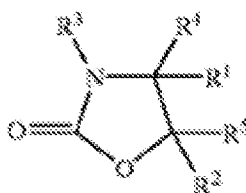
R^1 and R^2 independently of one another are selected from hydrogen, hydroxyl, phenyl, and C_1 - C_6 -alkyl;

R^3 is selected from hydrogen, C_1 - C_{12} -alkyl, N-hydroxy- (C_2-C_6) -alkyl, C_1 - C_{12} -alkenyl, and C_6 - C_{12} -aryl; and

R^4 and R^5 independently of one another are selected from C_1 - C_{12} -alkyl, C_1 - C_{12} -alkenyl, C_6 - C_{12} -aryl, C_1 - C_{12} -alkoxy, and hydrogen;

(ii) a photocatalyst being a hydroxy-substituted aromatic compound; (iii) a cosmetically acceptable carrier; wherein the crosslinking composition has not been exposed to electromagnetic radiation having a wavelength of about 750 nm or less for about 30 min or more; (c) applying the crosslinking composition to keratin fibres; and (d) mechanically shaping the keratin fibres with an appliance at a temperature of from about 80°C to about 180°C while exposing the crosslinking composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm.

Also described herein is a crosslinking composition for shaping keratin fibres comprising (a) a 2-oxazolidinone compound having the structure:



wherein:

R^1 and R^2 independently of one another are selected from hydrogen, hydroxyl, phenyl, and C_1 - C_6 -alkyl;

R³ is selected from, C₁-C₁₂-alkyl, N-hydroxy-(C₂-C₆)-alkyl, C₁-C₁₂-alkenyl, and C₆-C₁₂-aryl; and

R⁴ and R⁵ independently of one another are selected from C₁-C₁₂-alkyl, C₁-C₁₂-alkenyl, C₆-C₁₂-aryl, C₁-C₁₂-alkoxy, and hydrogen;

- 5 (b) a photocatalyst being a hydroxy-substituted aromatic compound; and (c) a cosmetically acceptable carrier; wherein the crosslinking composition has not been exposed to electromagnetic radiation having a wavelength of 750 nm or less for 30 min or more.

10 These and other features, aspects, and advantages of the invention will become evident to those of ordinary skill in the art from a reading of the following disclosure.

DETAILED DESCRIPTION OF THE INVENTION

In this document, including in all embodiments of all aspects, the following definitions apply unless specifically stated otherwise. All percentages are by weight of the total composition.

15 All ratios are weight ratios. References to 'parts' e.g. a mixture of 1 part X and 3 parts Y, is a ratio by weight. "QS" or "QSP" means sufficient quantity for 100% or for 100g. +/- indicates the standard deviation. All ranges are inclusive and combinable. The number of significant digits conveys neither a limitation on the indicated amounts nor on the accuracy of the measurements. All numerical amounts are understood to be modified by the word "about". All measurements are

20 understood to be made at 25°C and at ambient conditions, where "ambient conditions" means at 1 atmosphere (atm) of pressure and at 50% relative humidity. "Relative humidity" refers to the ratio (stated as a percent) of the moisture content of air compared to the saturated moisture level at the same temperature and pressure. Relative humidity can be measured with a hygrometer, in particular with a probe hygrometer from VWR® International. Herein: "min" means "minute" or

25 "minutes"; "mol" means mole; "nanometers" is abbreviated "nm"; "g" following a number means "gram" or "grams". All weights as they pertain to listed ingredients are based on the active level and do not include carriers or by-products that may be included in commercially available materials. Herein, "comprising" means that other steps and other ingredients can be in addition. "Comprising" encompasses the terms "consisting of" and "consisting essentially of".

30 The compositions, formulations, methods, uses, kits, and processes described herein can comprise, consist of, and consist essentially of the elements and limitations described herein, as well as any of the additional or optional ingredients, components, steps, or limitations described herein. Embodiments and aspects described herein may comprise or be combinable with

elements, features or components of other embodiments and/or aspects despite not being expressly exemplified in combination, unless an incompatibility is stated. "In at least one embodiment" means that one or more embodiments, optionally all embodiments or a large subset of embodiments, has/have the subsequently described feature. Where amount ranges are given, these are to be understood as being the total amount of said ingredient in the composition, or where more than one species fall within the scope of the ingredient definition, the total amount of all ingredients fitting that definition, in the composition. For example, if the composition comprises from about 1% to about 5% fatty alcohol, then a composition comprising 2% stearyl alcohol and 1% cetyl alcohol, would fall within the scope.

"Molecular weight" or "M.Wt." or "MW" and grammatical equivalents mean the number average molecular weight.

"Viscosity" is measured at 25°C using a HAAKE Rotation Viscometer VT 550 with cooling/heating vessel and sensor systems according to DIN 53019 at a shear rate of 12.9 s⁻¹.

"Water-soluble" refers to any material that is sufficiently soluble in water to form a clear solution to the naked eye at a concentration of 0.1% by weight of the material in water at 25°C. The term "water-insoluble" refers to any material that is not "water-soluble".

"Substantially free from" or "substantially free of" means less than about 1%, or less than 0.8%, or less than 0.5%, or less than 0.3%, or about 0%, by total weight of the composition or formulation.

"Keratin fibres" means fibrous material composed of keratin. "Hair" means mammalian keratin fibres including scalp hair, facial hair, eyelashes, and body hair. It includes such hair still being attached to a living subject and also hair that has been removed therefrom such as hair swatches and hair on a doll/mannequin. In at least one embodiment, "hair" means human hair. "Hair shaft" or "hair fibre" means an individual hair strand and may be used interchangeably with the term "hair."

"Cosmetically acceptable" means that the compositions, formulations or components described are suitable for use in contact with human keratinous tissue without undue toxicity, incompatibility, instability, allergic response, and the like. All compositions and formulations described herein which have the purpose of being directly applied to keratinous tissue are limited to those being cosmetically acceptable.

“Derivatives” includes but is not limited to, amide, ether, ester, amino, carboxyl, acetyl, acid, salt and/or alcohol derivatives of a given compound. In at least one embodiment, “derivatives thereof” means the amide, ether, ester, amino, carboxyl, acetyl, acid, salt and alcohol derivatives.

5 “Monomer” means a discrete, non-polymerised chemical moiety capable of undergoing polymerisation in the presence of an initiator or any suitable reaction that creates a macromolecule e.g. such as polycondensation, polyaddition, anionic or cationic polymerization. “Unit” means a monomer that has already been polymerised i.e. is part of a polymer.

“Polymer” means a chemical formed from the polymerisation of two or more monomers.
10 The term “polymer” shall include all materials made by the polymerisation of monomers as well as natural polymers. Polymers made from only one type of monomer are called homopolymers. Herein, a polymer comprises at least two monomers. Polymers made from two or more different types of monomers are called copolymers. The distribution of the different monomers can be random, alternating or block-wise (i.e. block copolymer). The term “polymer” used herein
15 includes any type of polymer including homopolymers and copolymers.

“Kit” means a package comprising a plurality of components. “Kit” may be referred to as “kit-of-parts”. An example of a kit is, for example, a first composition and a separately packaged second composition and optionally application instructions.

20 Description

Described herein is a method for shaping keratin fibres. The method allows the achievement of a semi-permanent hairstyle i.e. a durable hairstyle, and semi-permanent shaping of eyelashes. This semi-permanent hairstyle is retained after at least one shampoo treatment, particularly after five shampoo treatments, more particularly after 10 shampoo treatments.
25 Besides the increased durability of the hairstyle, this method prevents clumping of hair and/or improves post-shampoo detangling of hair and feel. In addition, the inventors have found that this method increases the water- and humidity-resistance of the shape, increases the ease of style and/or increases the manageability of the shape after shampooing.

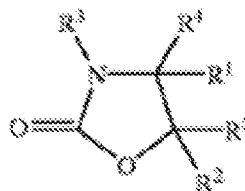
Without wishing to be bound by any theory, it is believed that the above benefits are due
30 to the steps conducted, their sequence, as well as the specific components used including the active agent. It is believed that the selected active agent diffuses into the shaft of the keratin fibre, reacts with amino groups in the keratin polypeptide and crosslinks these functional groups in the keratin protein structure, providing sufficient crosslinks to overcome the innate restoring force of

the keratin fibre structure. This results in a durable shaping of keratin fibres, for example a durable hairstyle or durable eyelash shape.

The details of the different aspects of the invention are described hereinafter.

Crosslinking composition

- 5 The crosslinking composition comprises a 2-oxazolidinone compound as an active agent which is useful for reacting with and providing crosslinks in keratin. The 2-oxazolidinone compounds have the structure:



wherein:

- 10 R^1 and R^2 independently of one another are selected from the group consisting of hydrogen, hydroxyl, phenyl, and C_1 - C_6 -alkyl;
 R^3 is selected from the group consisting of hydrogen, C_1 - C_{12} -alkyl, N-hydroxy-
 $(C_2$ - $C_6)$ -alkyl, C_1 - C_{12} -alkenyl, and C_6 - C_{12} -aryl; and
 R^4 and R^5 independently of one another are selected from the group consisting of
 C_1 - C_{12} -alkyl, C_1 - C_{12} -alkenyl, C_6 - C_{12} -aryl, C_1 - C_{12} -alkoxy, and hydrogen.

- 15 Examples of suitable crosslinkers of this type are 2-oxazolidinone, N-methyl-2-oxazolidinone, N-acetyl-2-oxazolidinone, and N-hydroxyethyl-2-oxazolidinone.

- The crosslinking composition may comprise a second active agent, such as a sugar. The reducing sugar may be selected from the group consisting of arabinose, xylose, ribose, and combinations thereof. Sugars such as arabinose are useful because they are naturally-derived.
 20 This is not only for perceived health and lack of sensitisation reasons, but also for sustainability and environmental reasons – sugars break down naturally and quickly and do not require special disposal methods. Furthermore, sugars are also easy to source and relatively inexpensive.

- Without wishing to be bound by theory, the second active agent may be useful for reacting with and providing further crosslinks in keratin. The hair comprises polypeptides of
 25 keratin having functional groups $-COOH$, OH , and NH_2 , and SH . Different active agents may react preferentially with each functional group. For instance, arabinose may react with amino groups whereas 2-oxazolidinone compounds may react with $-COOH$.

The crosslinking composition may comprise from about 0.1% to about 40%, or from about 0.5% to about 20%, or from about 1% to about 15%, or from about 7% to about 20%, or from about 8% to about 19%, or from about 10% to about 18% of a 2-oxazolidinone compound, by weight of the crosslinking composition.

5 In an embodiment, the crosslinking composition comprises from about 7% to about 20%, or from about 8% to about 19%, or from about 10% to about 18% of a 2-oxazolidinone compound, by weight of the crosslinking composition. In at least one embodiment, the crosslinking composition comprises from about 12% to about 18% of a 2-oxazolidinone compound weight of the crosslinking composition.

10 In at least one embodiment, the crosslinking composition comprises a cosmetically acceptable carrier. In at least one embodiment, the cosmetically acceptable carrier is any carrier suitable for formulating the active agent into a crosslinking composition being suitable for application onto hair. In at least one embodiment, the cosmetically acceptable carrier is selected from either an aqueous medium or an aqueous-alcoholic medium. In at least one embodiment,
15 when the carrier is an aqueous-alcoholic carrier, this carrier comprises water and an alcohol. In at least one embodiment, the alcohol is selected from the group consisting of: ethanol, isopropanol, propanol, and mixtures thereof. In at least one embodiment, when the carrier is an aqueous carrier, this carrier consists essentially of water and is substantially free of alcohol. In at least one embodiment, the crosslinking composition comprises a safe and effective amount of cosmetically
20 acceptable carrier. In at least one embodiment, the crosslinking composition comprises from about 0.1% to about 99%, or from about 1% to about 98%, or from about 10% to about 97%, or from about 30% to about 95% water.

The crosslinking composition comprises a photocatalyst comprises a hydroxy-substituted aromatic compound. A photocatalyst is an acid or base (or conjugate thereof) having a pKa (or
25 pKb) value that decreases (or increases) upon exposure to electromagnetic radiation, particularly light. Photoacids are mentioned for example in Domcke and Sobolewski (2003), *Unraveling the Molecular Mechanisms of Photoacidity*, 302, p.1693 and in Kowalewska (2005), *Photoacid catalyzed sol-gel process*, J.Mater. Chem. 15, p.4997, which are both incorporated herein by reference. In at least one embodiment, the electromagnetic radiation is selected from the group
30 consisting of: ambient light, sunlight, incandescent light, fluorescent light, LED light, laser light, and combinations thereof. In at least one embodiment, the electromagnetic radiation is selected from the group consisting of: visible light, near or far ultraviolet light, or near or far infrared light, and combinations thereof. In at least one embodiment, the photocatalyst can be activated to

a photo-excited state by excitation with incident radiation with a wavelength from about 300 nm to about 750 nm and wherein the appliance emits radiation with a wavelength from about 300 nm to about 750 nm.

In at least one embodiment, the hydroxy-substituted aromatic compound is a quinoline compound or a naphthol compound. In at least one embodiment, the hydroxy-substituted aromatic compound is a fluorescein or a derivative thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is a halogen-substituted fluorescein. In at least one embodiment, the hydroxy-substituted aromatic compound is bromo- or iodo-substituted fluorescein. In at least one embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: diiodofluorescein, 4,5,6,7-tetrachloro-2',4',5',7'-tetraiodofluorescein (rose Bengal), a salt of 2,4,5,7-tetraiodofluorescein (erythrosine), Eosin Y, Eosin B, 4,5-dibromofluorescein, and mixtures thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is a hydroxyflavone or a derivatives thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is a dihydroxyflavone or a trihydroxyflavone or a tetrahydroxyflavone or a mixture thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: 3 -hydroxy flavones, 7-hydroxy flavones, 5,7-hydroxy flavones, 4',5,7-trihydroxyflavone, 2-(3,4-dihydroxyphenyl)-3,5,7-trihydroxy-4*H*-chromen-4-one (quercitin), and mixtures thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is a hydroxyltriarylmethane, for example FD&C Green 3. In at least one embodiment, the hydroxy-substituted aromatic compound is an anthocyanidins or an anthocyanin. In at least one embodiment, the hydroxy-substituted aromatic compound is cyanidin (2-(3,4-dihydroxyphenyl) chromenylium-3,5,7-triol), malvidin, palargonidin or extracts containing anthocyanins such as elderberry, blueberry, cranberry, bilberry, red cabbage, sorghums, blackberry, black current, cherry red and black raspberry, and mixtures thereof.

In an embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: 8-hydroxyquinoline, 8-hydroxyquinoline sulfate, 8-quinolinol-1-oxide, 5-hydroxyquinoline, 6-hydroxyquinoline, 7-hydroxyquinoline, 5-iodo-7-sulfo-8-hydroxyquinoline, 5-fluoro-8-hydroxyquinoline, 5-fluoro-7-chloro-8-hydroxyquinoline, 5-fluoro-7-bromo-8-hydroxyquinoline, 5-fluoro-7-iodo-8-hydroxyquinoline, 7-fluoro-8-hydroxyquinoline, 5-chloro-8-hydroxyquinoline, 5,7-dichloro-8-hydroxyquinoline, 5-chloro-7-brono-8-hydroxyquinoline, 5-chloro-7-iodo-8-hydroxyquinoline, 7-chloro-8-hydroxyquinoline, 5-bromo-8-hydroxyquinoline, 5-bromo-7-chloro-8-hydroxyquinoline, 5,7-dibromo-8-hydroxyquinoline, 5-bromo-7-iodo-8-hydroxyquinoline, 7-bromo-8-hydroxyquinoline, 5-iodo-8-hydroxyquinoline, 5-iodo-7-chloro-8-

hydroxyquinoline, 5,7-diiodo-8-hydroxyquinoline, 7-iodo-8-hydroxyquinoline, 5-sulfonic acid-8-hydroxyquinoline, 7-sulfonic acid-8-hydroxyquinoline, 5-sulfonic acid-7-iodo-8-hydroxyquinoline, 5-thiocyano-8-hydroxyquinoline, 5-chloro-8-hydroxyquinoline, 5-bromo-8-hydroxyquinoline, 5,7-dibromo-8-hydroxyquinoline, 5-iodo-8-hydroxyquinoline, 5,7-diiodo-8-hydroxyquinoline, 7-azaindole, 7-cyano-2-naphthol, 8-cyano-2-naphthol, 5-cyano-2-naphthol, 1-hydroxy-3,6,8-pyrenetrisulfonic acid, Trans-3-hydroxystilbene, 2-hydroxymethylphenol, Pelargonidin, and mixtures thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: 8-quinolinol-1-oxide, 8-hydroxyquinoline, 7-cyano-2-naphthol, 8-cyano-2-naphthol, 5-cyano-2-naphthol, and mixtures thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: 8-quinolinol-1-oxide, 8-hydroxyquinoline, and mixtures thereof. In at least one embodiment, the hydroxy-substituted aromatic compound is 8-hydroxyquinoline. 8-hydroxyquinoline may act as a photoacid catalyst in lower pH solutions or as a photobase catalyst in higher pH solutions. 8-hydroxyquinoline has the CAS Number 148-24-3 and is available from Sigma-Aldrich. In at least one embodiment, the hydroxy-substituted aromatic compound is multi-cyclic. 8-hydroxyquinoline has the advantage of being easily available and characterised for use in cosmetic compositions such as hair dye compositions.

In an embodiment, the crosslinking composition comprises from about 10 ppm to about 500 ppm photocatalyst being a hydroxy-substituted aromatic compound. The concentration of photocatalyst may dependent, in part, on a variety of factors including, for example, the chemical structure of the photocatalyst, the reaction medium, the reaction type, and the substrate. In at least one embodiment, the crosslinking composition comprises from about 20 ppm to about 500 ppm, or from about 30 ppm to about 450 ppm, or from about 30 ppm to about 400 ppm, or from about 50 ppm to about 350 ppm, or from about 70 ppm to about 330 ppm, or from about 80 ppm to about 310 ppm, or from about 90 ppm to about 300 ppm, or from about 100 ppm to about 290 ppm, or to about 260 ppm, or to about 250 ppm, or to about 240 ppm, or to about 220 ppm, or to about 210 ppm, or to about 200 ppm photocatalyst being a hydroxy-substituted aromatic compound. To note: 1 ppm = 1 parts per million = $1 \times 10^{-4}\%$ = 0.0001% and 10 ppm = $1 \times 10^{-3}\%$ = 0.001% and 100 ppm = $1 \times 10^{-2}\%$ = 0.01%.

The crosslinking composition has not been exposed to electromagnetic radiation having a wavelength of 750 nm or less and for 30 min or more. The exposure of the crosslinking composition is important in view of the efficacy of the photocatalyst since when it is exposed to electromagnetic radiation within a certain wavelength range it will react with surround molecules

if they are available, including compounds in the packaging wall. In at least one embodiment, the crosslinking composition is packaged in a container where electromagnetic radiation having a wavelength of 750 nm or less is not able to contact the crosslinking composition. In at least one embodiment, the crosslinking composition has not been exposed to electromagnetic radiation
5 having a wavelength of 750 nm or less, or from about 300 nm to about 750 nm, or visible light, or UV light, and for 20 min or more, or 10 min or more, or 2 min or more. In at least one embodiment, the crosslinking composition is packaged in an opaque container. In at least one embodiment, the crosslinking composition is packaged in an amber- or brown-coloured container.

10 Other ingredients may be present in the crosslinking composition. In an embodiment, the crosslinking composition comprises antioxidant. Antioxidant is useful in view of providing longer-term stability for the crosslinking composition. In at least one embodiment, the crosslinking composition comprises a safe and effective amount of an antioxidant. In at least one
15 embodiment, the crosslinking composition comprises from about 0.001% to about 5%, or from about 0.5% to about 1.0% antioxidant. In at least one embodiment, the antioxidant is selected from the group consisting of: ascorbic acid (vitamin C), ascorbyl esters of fatty acids, ascorbic acid derivatives (e.g., magnesium ascorbyl phosphate, sodium ascorbyl phosphate, ascorbyl sorbate), tocopherol (vitamin E), tocopherol sorbate, tocopherol acetate, other esters of tocopherol, butylated hydroxy benzoic acids and their salts, peroxides including hydrogen
20 peroxide, perborate, thioglycolates, persulfate salts, 6-hydroxy-2,5,7,8-tetramethylchroman-2-carboxylic acid (commercially available under the tradename Trolox™), gallic acid and its alkyl esters, especially propyl gallate, uric acid and its salts and alkyl esters, ferulic acid and its salts and esters, sorbic acid and its salts, lipoic acid, amines (e.g., N,N-diethylhydroxylamine, amino-guanidine), sulfhydryl compounds (e.g., glutathione), dihydroxy fumaric acid and its salts, lysine
25 pidolate, arginine pilolate, nordihydroguaiaretic acid, bioflavonoids, curcumin, lysine, 1-methionine, proline, superoxide dismutase, silymarin, tea extracts, grape skin and/or grape seed extracts, melanin, rosemary extracts, and mixtures thereof. In at least one embodiment, the antioxidant is tocopherol sorbate or an ester of tocopherol. In at least one embodiment, the antioxidant is sodium benzoate. In at least one embodiment, the antioxidant is ascorbic acid.
30 Ascorbic acid has the benefit of enhancing the oxidative stability of the formulation. In at least one embodiment, the crosslinking composition comprises a safe and effective amount of ferulic acid. In at least one embodiment, the crosslinking composition comprises from about 0.001% to about 5%, or from about 0.5% to about 1.0% ferulic acid.

In at least one embodiment, the crosslinking composition comprises a chelator or chelating agent. As used herein, "chelator" or "chelating agent" means an active agent capable of removing a metal ion from a system by forming a complex so that the metal ion cannot readily participate in or catalyze chemical reactions. The inclusion of a chelating agent is especially useful for providing protection against UV radiation which can contribute to excessive scaling or skin texture changes and against other environmental agents which can cause skin damage, in order to decrease the local iron load, which generates, as indicated above, a pro-oxidant situation and pigmentation. A chelating agent is useful in view of providing longer-term stability for the crosslinking composition. In at least one embodiment, the crosslinking composition comprises a safe and effective amount of a chelator or chelating agent. In at least one embodiment, the crosslinking composition comprises a chelating agent, and wherein the chelating agent is selected from the group consisting of: N-hydroxysuccinimide, EDTA, NTA, deferoxamine, hydroxamic acids and their salts, phytic acid, phytate, gluconic acid and its salts, transferrine, lactoferrin, and mixtures thereof. In at least one embodiment, the crosslinking composition comprises a safe and effective amount of chelating agent. In at least one embodiment, the crosslinking composition comprises from about 0.001% to about 10%, or from about 0.01% to about 5%, or from about 0.1% to about 5%, or from about 0.5% to about 1.0% chelating agent. Exemplary chelators that are useful herein are disclosed in U.S. Pat. No. 5,487,884, issued Jan. 30, 1996 to Bissett *et al.*; International Publication No. 91/16035, Bush *et al.*, published Oct. 31, 1995; and International Publication No. 91/16034, Bush *et al.*, published Oct. 31, 1995. In at least one embodiment, the chelating agent is selected from the group consisting of: N-hydroxysuccinimide deferoxamine, lactoferrin, hydroxamic acids, gluconic acid, phytic acid, derivatives thereof, and mixtures thereof.

In at least one embodiment, the crosslinking composition is in a form suitable for application onto hair. In at least one embodiment, the crosslinking composition is in the form of an emulsion, a solution, or a dispersion. In at least one embodiment, the crosslinking composition comprises a surfactant. The surfactant can be useful in providing an emulsion. In at least one embodiment, when being in the form of an emulsion, said emulsion may be a water-in-oil emulsion, an oil-in-water emulsion, or a multiple emulsion. An emulsion has the benefit of providing an easy-to-apply composition for the consumer to apply to the hair and has aesthetic advantages. The crosslinking composition may be a leave-in composition or a rinse-off composition. The crosslinking composition may be in a form selected from: a shampoo; a hair conditioning composition; a hairstyling composition; or combinations thereof. When being a

hairstyling composition, said composition may be a gel composition; a spray gel composition, optionally dispensed using a mechanical spray device and/or at least one propellant; a non-aerosol hairspray, optionally dispensed using a suitable mechanically operated spraying device; a foamable composition, optionally dispensed using devices for foaming; hair wax composition; hair lotion composition; hair cream composition; or combinations thereof. In at least one embodiment, the crosslinking composition is a mascara composition.

The crosslinking composition may further comprise at least one cosmetic hair treatment agent selected from hairstyling polymers, conditioning agents, hair cleansing agents, or mixtures thereof. In at least one embodiment, the crosslinking composition comprises a hairstyling polymer. In at least one embodiment, the hairstyling polymer is selected from the group consisting of: non-ionic hairstyling polymer, anionic hairstyling polymer, zwitterionic and/or amphoteric hairstyling polymer, cationic hair styling polymer, or mixtures thereof. Suitable hairstyling polymers may be found in the CTFA International Cosmetics Ingredient Dictionary and Handbook, "Hair Fixatives", 12th edition (2008). Suitable hairstyling polymers are, for example, those materials disclosed from page 12, line 5 to page 19, line 1 of the European patent application 08151246.9 filed on 11 February 2008, which is incorporated herein by reference.

In at least one embodiment, the crosslinking composition comprises from about 0.01% to about 10% by weight, or from about 0.1% to about 8%, or from about 0.1% to about 5% hairstyling polymer.

In at least one embodiment, the crosslinking composition comprises a non-ionic hairstyling polymer. In at least one embodiment, the non-ionic hairstyling polymer is a natural or synthetic polymer. In at least one embodiment, the non-ionic hair styling polymers is a polymer obtained from the polymerisation of at least one type of monomer selected from: vinylpyrrolidone; vinylcaprolactam; vinyl esters; vinyl alcohol; vinyl acetate; (meth)acrylamide, and/or its derivatives; (meth)acrylic acid, its salts, and/or its derivatives; propylene and/or ethylene glycol acid; crotonic acid; or mixtures thereof. For example, such polymers are available under the trade names Luviskol® or Luviset Clear®.

In at least one embodiment, the crosslinking composition comprises an anionic hairstyling polymer. In at least one embodiment, the anionic hairstyling polymer is selected from the group consisting of: acrylic acid/alkyl acrylate/Nalkylacrylamide terpolymer; vinyl acetate/crotonic acid copolymer; C1-C5-alkyl acrylate/(meth)acrylic acid copolymer; sodium polystyrenesulfonate; vinyl acetate/crotonic acid/vinyl alkanoate copolymer; vinyl acetate/crotonic acid/vinyl neodecanoate copolymer; aminomethylpropanol acrylate copolymer;

vinylpyrrolidone/(meth)acrylic copolymer; methyl vinyl ether/maleic monoalkyl esters copolymer; aminomethylpropanol salts of allyl methacrylate/(meth)acrylate copolymer; ethyl acrylate/methacrylic acid copolymer; vinyl acetate/mono-nbutyl maleate/isobornyl acrylate copolymer; octylacrylamid/(meth)acrylic acid copolymer; polyesters of diglycol, 5 cyclohexanedimethanol, isophthalic acid and sulfoisophthalic acid; and mixtures thereof.

In at least one embodiment, the crosslinking composition comprises a zwitterionic or amphoteric hairstyling polymer. In at least one embodiment, the zwitterionic or amphoteric hairstyling polymer is selected from the group consisting of: alkylacrylamide/alkylaminoalkyl methacrylate/(meth)acrylic acid copolymers; copolymers which are formed from at least one first 10 monomer type which has quaternary amine groups, and at least one second monomer type which has acid groups; copolymers of fatty alcohol acrylates, of alkylamine oxide methacrylate and at least one monomer chosen from acrylic acid and methacrylic acid; methacryloylethylbetaine/methacrylic acid and/or esters copolymers; polyquaternium-47; polyquaternium-43; oligomers or polymers, preparable from quaternary croton betaines or 15 quaternary croton betaine esters; or mixtures thereof.

In at least one embodiment, the crosslinking composition comprises a cationic hairstyling polymer. In at least one embodiment, the cationic hairstyling polymer is selected from the group consisting of homopolymers or copolymers where a quaternary nitrogen groups are present either in the polymer chain or as substituent on one or more of the cationic monomers. The monomers 20 containing ammonium groups may be copolymerized with non-cationic monomers. Suitable cationic monomers may be unsaturated, free-radically polymerizable compounds which carry at least one cationic group, in particular ammonium-substituted vinyl monomers, such as, for example, trialkylmethacryloxyalkylammonium, trialkylacryloxyalkylammonium, dialkyldiallylammonium and quaternary vinylammonium monomers with cyclic, cationic 25 nitrogen-containing groups, such as pyridinium, imidazolium or quaternary pyrrolidones, e.g. alkylvinylimidazolium, alkylvinylpyridinium, or alkylvinylpyrrolidone salts. The alkyl groups of these monomers may be lower alkyl groups, for example, C1 to C7-alkyl groups, or C1 to C3-alkyl groups. Suitable non-cationic monomers may be selected from (meth)acrylamide, derivatives thereof; acrylate, its derivative thereof; vinylcaprolactone, vinylcaprolactam, 30 vinylpyrrolidone, vinyl esters, vinyl alcohol, propylene glycol or ethylene glycol. For example, suitable cationic hairstyling polymers are available under the tradenames Gafquat 755 N; Gafquat 734; Gafquat HS 100; Luviquat HM 550; Merquat Plus 3300; Gaffix VC 713; Aquaflex SF 40. In at least one embodiment, the crosslinking composition comprises a cationic hairstyling

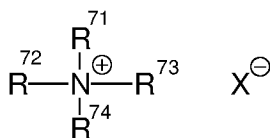
polymer derived from a natural polymer. In at least one embodiment, the cationic hairstyling polymer derived from a natural polymer is derived from a natural polymer selected from the group consisting of: cationic derivatives polysaccharides such as cellulose, starch and/or guar; chitosan, its salts, and/or its derivatives; or mixtures thereof. In at least one embodiment, the cationic hairstyling polymers are selected from the group consisting of: polyquaternium-4; polyquaternium-10; polyquaternium-24; guar hydroxypropyltrimonium chloride; chitosonium pyrrolidonecarboxylate; and mixtures thereof.

In at least one embodiment, the crosslinking composition comprises a conditioning agent, or a hair conditioning agent. The crosslinking composition may comprise any suitable and conventional hair conditioning agents. The term “hair conditioning agent” herein means any cosmetically acceptable compound having a cosmetic effect on hair, such as providing gloss to hair, making hair more manageable, improving hair touch, improving combability and/or giving hair more volume. Suitable hair conditioning agents may be found in the CTFA International Cosmetics Ingredient Dictionary and Handbook, “Hair conditioning agents”, 12th edition (2008).

In at least one embodiment, the hair conditioning agent is selected from the group consisting of: cationic surfactants, non-ionic surfactants, silicone compounds, organic oily conditioning agents, and mixtures thereof. Suitable hair conditioning agents are, for example, those materials disclosed from page 19, line 3 to page 27, line 33 of the European patent application 08151246.9 filed on 11 February 2008, which is incorporated herein by reference.

In at least one embodiment, the conditioning agent is a cationic surfactant. In at least one embodiment, the cationic surfactant comprises amino or quaternary ammonium moieties. In at least one embodiment, the crosslinking composition comprises from about 0.05% to about 3.5%, or from about 0.1% to about 3.0%, or from about 0.5% to about 2.5%, or from about 1.0% to about 2.0% cationic surfactant. In at least one embodiment, cationic surfactant is according to

Formula II:



II

wherein at least one of R⁷¹, R⁷², R⁷³ and R⁷⁴ is selected from: an aliphatic group of from 8 to 30 carbon atoms; an aromatic, alkoxy, polyoxyalkylene, alkylamido, hydroxyalkyl, aryl; or an alkylaryl group having from 7 to 22 carbon atoms; wherein the remainder of R⁷¹, R⁷², R⁷³ and R⁷⁴ are independently selected from the group consisting of: an aliphatic group consisting of from 1 to 22 carbon atoms; and an aromatic, alkoxy, polyoxyalkylene, alkylamido, hydroxyalkyl, aryl or

alkylaryl group having up to 22 carbon atoms; wherein X is selected from the group consisting of: halogen, acetate, citrate, lactate, glycolate, phosphate, nitrate, sulfonate, sulfate, alkylsulfate, alkyl sulfonate radicals, and mixtures thereof. In at least one embodiment, cationic surfactant is according to Formula II (see above), wherein at least one of R^{71} , R^{72} , R^{73} and R^{74} is an aliphatic group having from 16 to 24 carbon atoms; wherein the remainder of R^{71} , R^{72} , R^{73} and R^{74} are independently selected from the group consisting of aliphatic groups having from 1 to 4 carbon atoms; wherein X is selected from the group consisting of: chloride or sulfate. In at least one embodiment, the cationic surfactant is selected from the group consisting of: behenyltrimethylammonium chloride, methyl sulfate or ethyl sulfate; stearyltrimethylammonium chloride, methyl sulfate or ethyl sulfate; and mixtures thereof. It is believed that a longer alkyl group provides improved smoothness and soft feeling on wet and dry hair, compared to cationic surfactants with a shorter alkyl group. It is also believed that such cationic surfactants can provide reduced scalp irritation, compared to those having a shorter alkyl group. In at least one embodiment, the cationic surfactant is a di-long alkyl quaternized ammonium salt selected from the group consisting of: dialkyl (14-18 carbons) dimethyl ammonium chloride, ditallow alkyl dimethyl ammonium chloride, dihydrogenated tallow alkyl dimethyl ammonium chloride, distearyl dimethyl ammonium chloride, dicetyl dimethyl ammonium chloride, and mixtures thereof. In at least one embodiment, the cationic surfactant is a tertiary amidoamine having an alkyl group of from about 12 to about 22 carbons. In at least one embodiment, the cationic surfactant is selected from the group consisting of: cetyl trimethyl ammonium salts; behenyl trimethyl ammonium salts; dimethyl ditallow ammonium salts; stearyl amidopropyl dimethylamine; (di)esterquats; quaternium 8, 14, 15, 18, 22, 24, 26, 27, 30, 33, 37, 53, 60, 61, 72, 78, 80, 81, 82, 83, 84, and/or 91; or mixtures thereof.

In at least one embodiment, the conditioning agent is a non-ionic surfactant. Suitable non-ionic surfactants may be surfactants having a HLB of less than 8. Suitable nonionic surfactants may be selected from glyceryl esters; sugar esters; alkylpolyglucoside ethers; oleyl- or isostearylpolyglucoside; polyoxyethylene (20) sorbitan monostearate; or mixtures thereof.

In at least one embodiment, the conditioning agent is a silicone compound. In at least one embodiment, the silicone compound is volatile or nonvolatile, and/or soluble or insoluble silicones. For example, suitable silicone conditioning agents are available under the tradenames SF 1075 methyl phenyl fluid (Electric company); DC200 Fluid, DC244, DC245, DC345, Dow 5-7113, DC556 Cosmetic Grade Fluid, DC1248 (Dow Corning). In at least one embodiment, the crosslinking composition comprises a conditioning agent being the reaction product of: (a) an

aminosilane; (b); polysiloxane; and optionally (c) a polyether. In at least one embodiment, the crosslinking composition comprises a conditioning agent being the reaction product of: (a) an aminosilane; (b); polysiloxane; and (c) a polyether. In at least one embodiment, the crosslinking composition comprises a conditioning agent, and wherein the conditioning agent is selected from the group consisting of: epoxyaminosilane copolymers, and polysiloxane/polyurea block copolymers, and mixtures thereof. In at least one embodiment, the crosslinking composition comprises a conditioning agent being the reaction product of: (a) an aminosilane; (b) polysiloxane; and (c) a polyether; and optionally (d) an amine. In at least one embodiment, the polysiloxane is an epoxy encapped polysiloxane. In at least one embodiment, the polysiloxane comprises at least two oxirane or oxetane groups. In at least one embodiment, the polysiloxane comprises from about 10 to about 450 silicon atoms, or from about 40 to about 400 silicon atoms, from about 75 to about 350 silicon atoms, from about 150 to about 250 silicon atoms. In at least one embodiment, the polysiloxane is an epoxy encapped polysiloxane. In at least one embodiment, the polyether has the average structure $\text{CH}_2(\text{O})\text{CHCH}_2\text{O}(\text{CH}_2(\text{CH}_3)\text{CH}_2\text{O})_n\text{CH}_2\text{CH}(\text{O})\text{CH}_2$ wherein n is an integer from 1 to 10. In at least one embodiment, the amine comprises from 1 to 10 carbon atoms, or from 2 to 5 carbon atoms. In at least one embodiment, the amine is an alkylamine that is substituted with at least one alkyl group. In at least one embodiment, the amine is selected from the group consisting of: methylamine, ethylamine, propylamine, ethanol amine, isopropylamine, butylamine, isobutylamine, hexylamine, dodecylamine, oleylamine, aniline aminopropyltrimethylsilane, aminopropyltriethylsilane, aminomorpholine, aminopropyldiethylamine benzylamine, naphthylamine 3-amino-9-ethylcarbazole, 1-aminoheptaphlorohexane, 2,2,3,3,4,4,5,5,6,6,7,7,8,8,8-pentadecafluoro-1-octanamine, and mixtures thereof. In at least one embodiment, the amine is selected from the group consisting of: methylethylamine, methylhexylamine, methyloctadecylamine, diethanolamine, dibenzylamine, dihexylamine dicyclohexylamine, piperidine, pyrrolidine phthalimide, and mixtures thereof. In at least one embodiment, the conditioning agent is an epoxyaminosilane copolymer. In at least one embodiment, the conditioning agent is conditioning agent being the reaction product of: (a) an aminosilane; (b) polysiloxane, wherein the polysiloxane comprises from about 10 to about 450 silicon atoms, or from about 40 to about 400 silicon atoms; and (c) a polyether; and (d) an amine, wherein the amine is an alkylamine that is substituted with at least one alkyl group. Epoxyaminosilane copolymers are described in EP2214633B1 (filing date 30th Oct 2008, which is incorporated herein by reference) and are available from Momentive™ Performance Materials

Inc., Columbus, Ohio, USA. Epoxyaminosilane copolymers have excellent durability benefits. Such an exemplary epoxyaminosilane copolymer may be synthesised as follows: aminopropyltriisopropoxy silane (40.77g), an epoxy encapped polysiloxane with the average structure

- 5 $\text{CH}_2(\text{O})\text{CHCH}_2\text{OCH}_2\text{CH}_2\text{Si}(\text{CH}_3)_2\text{O}[\text{Si}(\text{CH}_3)_2\text{O}]_{50}\text{Si}(\text{CH}_3)_2\text{CH}_2\text{CH}_2\text{CH}_2\text{OCH}_2\text{CH}(\text{O})\text{CH}_2$
(171.40g) and an epoxy endcapped polyether with the average structure $\text{CH}_2(\text{O})\text{CHCH}_2\text{O}(\text{CH}_2(\text{CH}_3)\text{CH}_2\text{O})_7\text{CH}_2\text{CH}(\text{O})\text{CH}_2$ (37.83g) and isopropanol (425.68g) is combined in a 500 mL flask. The material is brought to reflux and stirred with an overhead stirrer. The refluxing continued for 15.5 hr until all epoxy groups are consumed as determined by
- 10 titration. The material is transferred to a rotary evaporator and stripped at 70°C and 532 Pa (4 torr) for 2 hrs to remove the isopropanol. Another exemplary epoxyaminosilane copolymer may be synthesised as follows: aminopropyltriisopropoxy silane (14.27g), 3-(diethylamino)propylamine (7.05g), an epoxy encapped polysiloxane with the average structure $\text{CH}_2(\text{O})\text{CHCH}_2\text{OCH}_2\text{CH}_2\text{CH}_2\text{Si}(\text{CH}_3)_2\text{O}[\text{Si}(\text{CH}_3)_2\text{O}]_{200}\text{Si}(\text{CH}_3)_2\text{CH}_2\text{CH}_2\text{CH}_2\text{OCH}_2\text{CH}(\text{O})\text{CH}_2$
- 15 (447.87g) and an epoxy encapped polyether with the average structure $\text{CH}_2(\text{O})\text{CHCH}_2\text{O}(\text{CH}_2\text{CH}_2\text{O})_{14}\text{CH}_2\text{CH}(\text{O})\text{CH}_2$ (30.81 g) and isopropanol (500 g) is combined in a 2000 mL flask. The material is brought to reflux and stirred with an overhead stirrer. The refluxing continued for 24 hr until all epoxy groups are consumed as determined by titration. The material is transferred to a rotary evaporator and stripped at 70°C and 532 Pa (4 torr) for 2 hrs to
- 20 remove the isopropanol.

In an embodiment, the conditioning agent is selected from the group consisting of: epoxyaminosilane copolymers, and polysiloxane/polyurea block copolymers, and mixtures thereof. A polysiloxane/polyurea block copolymer is described in EP2074986B1 filed on 10th Dec 2008, which is incorporated herein by reference. In at least one embodiment, the

25 polysiloxane/polyurea block copolymer comprises at least one polysiloxane sequence (or block) and at least one polyurea sequence (block) in the backbone of the copolymer. In at least one embodiment, the amount of polysiloxane present in the copolymer is greater than 90% by weight relative to the total weight of the polysiloxane/polyurea block copolymer. In at least one embodiment, the polysiloxane/polyurea block copolymer of the does not comprise polyurethane.

30 By way of non-limiting example, the copolymer can be a non-ionic polysiloxane/polyurea copolymer, that is to say that it does not comprise an ionized or ionizable group. By way of example of a copolymer, non-limiting mention may be made of the dimethylpolysiloxane/polyurea block copolymer having the INCI name polyureadimethicone.

Such a dimethylpolysiloxane/polyurea block copolymer can be obtained, for instance, by copolymerization of an α,ω -aminosilicone with a diisocyanate. Polysiloxane/polyurea block copolymers corresponding to these characteristics are, for example, the products sold under the reference Wacker-Belsil® UD 60, Wacker-Belsil® UD 80, Wacker-Belsil® DU 140 and
5 Wacker-Belsil® UD 200 by Wacker. In at least one embodiment, the polysiloxane/polyurea copolymer is non-ionic. In at least one embodiment, the crosslinking composition comprises from about 0.05 to about 20%, for example from 0.1 to 15%, or from 0.5 to 10% polysiloxane/polyurea block copolymer. In at least one embodiment, the conditioning agent is a polydimethylsiloxane-derivative comprising aminoalkyl groups and having an amine number of
10 at least 0.1 meq/g of polydimethylsiloxane. Such polydimethylsiloxane-derivative can, for example, be methoxy-terminated or hydroxy-terminated, or mixtures thereof. The polydimethylsiloxane-derivative can, for example be a mixture of methoxy- and hydroxy-terminated Poly[3-((2-aminoethyl)amino)propyl]methyl(dimethyl)siloxane as commercially sold as Wacker-Belsil® ADM 1370.

15 In at least one embodiment, the conditioning agent is an organic oily conditioning agent. In at least one embodiment, the organic oily conditioning agent is non-volatile, water-insoluble, oily or fatty. Organic oily conditioning agents may be selected from hydrocarbon oils and fatty esters. In at least one embodiment, the conditioning agent is a fatty alcohol. In at least one embodiment, the fatty alcohol is a non-volatile low melting point fatty alcohol. In at least one
20 embodiment, the conditioning agent is a fatty alcohol and the fatty alcohol is selected from the group consisting of: capryl alcohol, lauryl alcohol, stearyl alcohol, cetyl alcohol, myristyl alcohol, palmitoleyl alcohol, and mixtures thereof.

The crosslinking composition may further comprise at least one direct hair dye. In at least one embodiment, the crosslinking composition comprises from about 0.01% to about 15%, or
25 from about 0.1% to about 10 %, or from about 0.5% to about 8% direct hair dye.

The crosslinking composition may further comprise at least one viscosity-modifying agent. In at least one embodiment, the crosslinking composition comprises from about 0.01% to about 20%, or from about 0.05% to about 10%, or from about 0.1% to about 5% viscosity-modifying agent.

The crosslinking composition may further comprise at least one emulsifier and/or surfactant. In at least one embodiment, the emulsifier and/or surfactant is selected from nonionic surfactants; anionic surfactants; amphoretic surfactants; or mixtures thereof. In at least one embodiment, the crosslinking composition comprises from about 0.01% to about 20%, or from
5 about 0.05% to about 10%, or from about 0.1% to about 5%, emulsifier and/or surfactant.

The crosslinking composition may further comprise at least one pigment. In at least one embodiment, the pigment is selected from natural pigments; synthetic pigments; or mixtures thereof. The pigments may be selected from organic pigment, inorganic pigment; or mixtures thereof. The pigments may be selected from coloured pigments; pearlescent pigments; or
10 mixtures thereof. Said crosslinking composition may comprise from about 0.01% to 10%, or from about 1% to about 2% pigment present in the product mass in undissolved form by weight of the total crosslinking composition. The crosslinking composition may comprise pigment materials such as inorganic, nitroso, monoazo, disazo, carotenoid, triphenyl methane, triaryl methane, xanthene, quinoline, oxazine, azine, anthraquinone, indigoid, thionindigoid,
15 quinacridone, phthalocianine, botanical, natural colors, including: water-soluble components such as those having C.I. Names.

In at least one embodiment, the crosslinking composition comprises at least one particulate substance. In at least one embodiment, the particulate substance is selected from silica; silicates; aluminates; clay earths; mica; insoluble salts, particularly insoluble inorganic
20 metal salts; metal oxides; minerals; insoluble polymer particles; or mixtures thereof. In at least one embodiment, the crosslinking composition comprises from about 0.01% to about 10%, or from about 0.05% to about 5% of at least one particulate substance. In at least one embodiment, the crosslinking composition is substantially free of a particulate substance such as clay.

In at least one embodiment, the crosslinking composition comprises at least one
25 preservative. In at least one embodiment, the crosslinking composition may comprise from about 0.01% to about 5% by weight, or from about 0.05% to about 1% preservative.

A variety of additional optional ingredients may be incorporated into the crosslinking composition described herein. Non-limiting examples of these additional ingredients may be selected from preservatives; antioxidants; sequestering agents; solvents; fragrances & perfumes;
30 fillers; screening agents; odour absorbers; colouring materials; lipid vesicles; deterative surfactants; thickening agents and suspending agents; viscosity modifiers; pearlescent aids; UV-filters and sunscreens; agents for combating free radicals; polyvinyl alcohol; pH adjusting agents; salts; colouring agents; polymer plasticizing agents; direct dyes; or mixtures thereof. The

crosslinking composition may comprise from about 0%, or from about 0.1% to about 5% antimicrobial agents. In at least one embodiment, the crosslinking composition comprises an organic acid selected from the group consisting of: glycine, L-methionine, L-arginine, biotin, creatine, and mixtures thereof. In at least one embodiment, the crosslinking composition
5 comprises an antidandruff agent. In at least one embodiment, the crosslinking composition comprises zinc pyrithione. In at least one embodiment, the crosslinking composition comprises panthenol. In at least one embodiment, the crosslinking composition comprises a wax compound. In at least one embodiment, the crosslinking composition comprises beeswax.

In at least one embodiment, the crosslinking composition has a viscosity, measured at
10 25°C, of from about 0.1 mPa·s to about 1,000,000 mPa·s, or from about 1 mPa·s to about 80,000 mPa·s, or from about 5 mPa·s to about 3,500 mPa·s. The viscosity is measured by HAAKE Rotation Viscometer VT 550 with cooling/heating vessel and sensor systems according to DIN 53019 (MV-DIN, SV-DIN), shear rate is 12.9 s⁻¹.

In at least one embodiment, the crosslinking composition is substantially free of:
15 formaldehyde, derivatives of formaldehyde, methylene glycol, formalin, and any compound that produces formaldehyde upon heating. "Heating" means raising the temperature of the compound above 25°C. In at least one embodiment, the crosslinking composition comprises 0% formaldehyde. In at least one embodiment, the derivatives of formaldehyde are 1,3,5-trioxane and paraformaldehyde. In at least one embodiment, the crosslinking composition is substantially
20 free of: formaldehyde, 1,3,5-trioxane, paraformaldehyde, methylene glycol, formalin. Formaldehyde may not be desired in view of its safety profile. Formalin is not advantageous because formalin is a derivative of formaldehyde. Formaldehyde exists in multiple forms. In water, formaldehyde becomes hydrated and forms methylene glycol. A saturated solution of formaldehyde (about 40% formaldehyde) in water is more commonly known as formalin.
25 Methanol and/or methylene diol can be used as a stabilizer in formalin and is therefore not advantageous. In at least one embodiment, the crosslinking composition is substantially free of a quaternary ammonium compound and/or a surfactant. In at least one embodiment, the crosslinking composition is substantially free of: ceramide compound, an alpha-hydroxy acid, a thioglycolate and/or thiolactate compound, a bisulfate compound, clay, a reducing agent. In at
30 least one embodiment, the crosslinking composition is substantially free of: ceramide compound, an alpha-hydroxy acid, a thioglycolate and/or thiolactate compound, a bisulfate compound. In at least one embodiment, the crosslinking composition is substantially free of a carbonate compound. In at least one embodiment, the crosslinking composition is substantially free of:

ceramide compound, an alpha-hydroxy acid, a thioglycolate or thiolactate compound, a bisulfate compound, clay, formaldehyde, 1,3,5-trioxane, paraformaldehyde, methylene glycol, quaternary ammonium compound, surfactant.

In at least one embodiment, the crosslinking composition has a pH of from about pH 6 to about pH 11. In at least one embodiment, the crosslinking composition has a pH of from about pH 6.5 to about pH 10.75, or from about pH 7.0 to about pH 10.5, or from about pH 7.5 to about pH 10.25, or from about pH 8.0 to about pH 10.0. A basic pH is useful for 2-oxazolidinone compounds in view of penetration into hair. In at least one embodiment, the crosslinking composition comprises a buffering agent. In at least one embodiment, the buffering agent is a phosphate buffer. In at least one embodiment, the buffering agent is a borate buffer or a carbonate buffer. In at least one embodiment, the buffering agent is selected from the group consisting of: glycine/sodium hydroxide; sodium carbonate/sodium hydrogen carbonate, sodium tetraborate/sodium hydroxide; sodium bicarbonate/sodium hydroxide; ammonium chloride/ammonia. The buffering agent has the advantage of controlling the pH, which aids the stability of the crosslinking composition. In at least one embodiment, the crosslinking composition comprises an alkalizing agent and/or an agent for adjusting the pH value. The crosslinking composition may further comprise a protonating agent. The protonating agent may be a monoprotic or polyprotic acid, water-soluble or water-insoluble acid, and/or an organic or inorganic acid. In at least one embodiment, the protonating agent is selected from formic acid, acetic acid, sulfuric acid, hydrochloric acid, citric acid, and mixtures thereof. In at least one embodiment, the protonating agent is citric acid. Citric acid is useful because it is naturally available from lemons.

Method

Also described herein is a method comprising (a) providing a crosslinking composition; and (b) applying the crosslinking composition to keratin fibers, and mechanically shaping the kerton fibers with an appliance at a temperature of from about 80°C to about 180°C, alternatively from about 80°C to about 175°C, and alternatively from about 80°C to about 170°C, and exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm.

In an one embodiment, the method does not comprise a rinsing step between step (a) and step (b). It may not be helpful for the efficacy of the crosslinking composition described herein if the active agent and photocatalyst are washed off the keratin fibres prior to exposing the composition to electromagnetic radiation. In at least one embodiment, the mechanical shaping of

keratin fibres is selected from the group consisting of: straightening keratin fibres, such as human scalp hair or human eyelashes, and curling keratin fibres, such as human scalp hair or human eyelashes.

The method comprises (b) applying the crosslinking composition to keratin fibres. Further
5 detail on embodiments and description of the crosslinking composition is provided above. In at least one embodiment, the crosslinking composition remains on the keratin fibres for a time X, wherein time X is from about 2 min to about 60 min, or from about 5 min to about 30 min.

The method uses an appliance or implement for mechanically shaping the keratin fibers. In at least one embodiment, the appliance is a hair straightening appliance comprising a heating
10 element. In at least one embodiment, the appliance is a hair curling appliance comprising a heating element. In at least one embodiment, the appliance is a hair straightening appliance comprising a heating element and a source of electromagnetic radiation for exposing hair to electromagnetic radiation. In at least one embodiment, the implement is a combing or brushing
15 means and the crosslinking composition is exposed to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm using a separate source of electromagnetic radiation such as a lamp. In at least one embodiment, the appliance comprises light-emitting diodes.

The crosslinking composition is exposed to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm. In at least one embodiment, the
20 electromagnetic radiation has a wavelength of from about 310 nm, or from about 320 nm, or from about 330 nm, or from about 340 nm, or from about 350 nm, or from about 360 nm, or from about 370 nm, or from about 380 nm, or from about 390 nm, or from about 400 nm, or from about 410 nm, to about 740 nm, or to about 730 nm, or to about 720 nm, or to about 710 nm, or to about 700 nm, or to about 690 nm, or to about 680 nm, or to about 670 nm, or to about 650
25 nm, or to about 640 nm. In at least one embodiment, the electromagnetic radiation has a wavelength of from 380 nm to about 550 nm.

Irradiance, that is the power of electromagnetic radiation in Watts per unit area, has the unit Watts per m² or W/m². Irradiance is thus a measurement of the intensity of electromagnetic radiation. Light intensity can also be measured in lux (lx), which is the unit of illuminance. 1 lx =
30 about 1.5×10^{-7} W/cm² (at 555 nm). An average laboratory or office space would have a light intensity of about 200 lx to about 1000 lx i.e. an irradiance of about 2.9×10^{-5} W/cm² to about 1.4×10^{-4} W/cm² (at 555 nm). In at least one embodiment, the electromagnetic radiation has an irradiance of at least about 1×10^{-3} W/cm², or at least about 5×10^{-3} W/cm², or at least about $1 \times$

10^{-2} W/cm², or at least about 5×10^{-2} W/cm², or at least about 1×10^{-1} W/cm², or at least about 5×10^{-1} W/cm². In at least one embodiment, the electromagnetic radiation has an illuminance of at least about 1000 lx, or at least about 2000 lx, or at least about 3000 lx, or at least about 4000 lx, or at least about 5000 lx, or at least about 6000 lx, or at least about 7000 lx, or at least about 8000 lx, or at least about 9000 lx, or at least about 10000 lx, or at least about 20000 lx, or at least about 30000 lx, or at least about 40000 lx, or at least about 50000 lx, or at least about 60000 lx, or at least about 70000 lx, or at least about 80000 lx.

In at least one embodiment, the electromagnetic radiation is selected from the group consisting of: ambient light, sunlight, incandescent light, fluorescent light, LED light, laser light, and combinations thereof. In at least one embodiment, the electromagnetic radiation is selected from the group consisting of: visible light, near or far ultraviolet light, or near or far infrared light, and combinations thereof. In at least one embodiment, the electromagnetic radiation is light. In at least one embodiment, the suitable light may be provided from any source capable of illuminating the substrate surface. In at least one embodiment, the light is selected from the group consisting of: ambient sunlight, incandescent light, and fluorescent light. In at least one embodiment, the light is provided by conventional sources such as lamps and portable or battery-powered lights. Specific devices may be developed or adapted for use with the crosslinking compositions and method described herein. In at least one embodiment, the appliance is a hair brush configured to incorporate LEDs. In at least one embodiment, the light is laser light. Laser may be used to provide precise targeting, for example. In at least one embodiment, the appliance is hybrid heat and light providing hair straightening irons.

In at least one embodiment, step (a) is carried out and then subsequently step (b). Step (b) comprises applying the crosslinking composition to keratin fibres, mechanically shaping the keratin fibres with an appliance or implement, and exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm. In at least one embodiment, step (b) comprises (i) applying the crosslinking composition to keratin fibres; and then subsequently (ii) mechanically shaping the keratin fibres with an appliance and exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm. In at least one embodiment, step (b) comprises (i) applying the crosslinking composition to keratin fibres; and then subsequently (ii) mechanically shaping the keratin fibres with an appliance and exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm. In at least one embodiment, step (b) comprises (i) applying the crosslinking composition to keratin fibres; (ii) drying the fibres; and then subsequently (iii)

mechanically shaping the keratin fibres with an appliance, wherein the temperature of the appliance is from about 50°C to about 250°C, or from about 80°C to about 180°C, or from about 80°C to about 170°C and then exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm.

5 In at least one embodiment (b) comprises drying the fibres. In an embodiment, the hair drying is carried out by a blow drier. In an embodiment, the hair drying is carried out for a duration of from about 1 min to about 45 min, or from about 2 min to 20 min, or from about 5 min to 15 min. In general, following the hair drying, the hair can still be damp, but needs to have reasonable e.g. 75% hair fibre separation of the head of hair. Some residual moisture in the hair is
10 acceptable.

In at least one embodiment, (b) comprises drying the hair fibre is carried out by a hood appliance. In an embodiment, the hair drying is carried out by towelling hair and/or by pressing hair with hands.

Hair dryer or blow dryer distances between device and head are typically down to about
15 10 cm. Blow dryers direct hot air through some sort of attachment for combing or otherwise treating the hair. A blow dryer is typically used such that the distance to the hair (for example at a distance of 20 or 30 or 40 centimetres) and often is used with the aid of a comb or a brush. In an embodiment, the hair drying is carried out by a blow drier at a temperature of from about from 50°C to about 100°C. In an embodiment, the hair drying is carried out by a blow drier at a
20 temperature of up to 130°C. In an embodiment, the hair drying (b) is carried out with a blow drier with brushing to help straighten the hair.

In at least one embodiment, (b) comprises elevating the temperature of the appliance to a temperature of from about 80°C to about 180°C. In at least one embodiment, (b) involves elevating the temperature of the appliance to a temperature of from about 50°C to about 250°C.

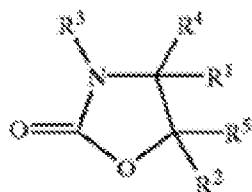
25 In at least one embodiment, (b) involves elevating the temperature of the appliance to a temperature of from about 50°C, or from about 60°C, or from about 70°C, or from about 80°C to about 220°C, or to about 200°C, or to about 180°C, or to about 170°C, or to about 160°C, or to about 150°C, or to about 140°C, or to about 130°C.

In at least one alternative embodiment, the crosslinking composition is formed on the
30 keratin fibres rather than provided prior to applying to keratin fibres. For example, a crosslinking formulation comprising the active agent and a cosmetically acceptable carrier is applied to the keratin fibres and then subsequently, a photocatalyst formulation comprising the photocatalyst is applied to the keratin fibres.

In an embodiment, a method for shaping keratin fibres comprising:

- (a) providing a crosslinking composition, wherein the crosslinking composition comprises:

(i) a 2-oxazolidinone compound of the general structure:



Where:

R^1 and R^2 independently of one another are hydrogen, hydroxyl, phenyl or C_1 - C_6 -alkyl, R^3 is hydrogen, C_1 - C_{12} -alkyl, N-hydroxy- $(C_2$ - $C_6)$ -alkyl, C_1 - C_{12} -alkenyl or C_6 - C_{12} -aryl, and R^4 and R^5 independently of one another are C_1 - C_{12} -alkyl, C_1 - C_{12} -alkenyl, C_6 - C_{12} -aryl, C_1 - C_{12} -alkoxy, or hydrogen.

(ii) a photocatalyst being a hydroxy-substituted aromatic compound;

(iii) a cosmetically acceptable carrier;

wherein the crosslinking composition has not been exposed to electromagnetic radiation having a wavelength of about 750 nm or less for 30 min or more;

- (b) applying the crosslinking composition to keratin fibres, drying the fibres, mechanically shaping the keratin fibres with an appliance, and exposing the composition to electromagnetic radiation having a wavelength of from about 300 nm to about 750 nm.

In at least one embodiment, the crosslinking composition further comprises a viscosity-modifying agent and a conditioning agent. In at least one embodiment, the crosslinking composition has a pH of from about pH 8 to about pH 11.

In an embodiment, a method for shaping keratin fibres comprising: (a) providing a solid composition, wherein the solid composition comprises 2-oxazolidinone; a photocatalyst being a hydroxy-substituted aromatic compound; and then mixing the solid composition with a cosmetically acceptable carrier to form a crosslinking composition. The crosslinking composition is described herein. The second aspect has the advantage that it is not a requirement that solid composition has not been exposed to electromagnetic radiation having a wavelength of 750 nm or less for 30 min or more. This is because in solid form, the photocatalyst, if exposed to

electromagnetic radiation having a wavelength of 750 nm or less, cannot react with undesired materials e.g. container packaging. The solid composition is solid at 25°C.

In an embodiment, the crosslinking composition may be used for shaping keratin fibres. In at least one embodiment, the use is for straightening keratin fibres. In at least one embodiment, the use is for mechanically shaping keratin fibres. In at least one embodiment, the mechanical shaping of keratin fibres is selected from the group consisting of: straightening keratin fibres, such as human scalp hair or human eyelashes, and curling keratin fibres, such as human scalp hair or human eyelashes.

In an embodiment, a kit comprises a crosslinking formulation comprising a first and second active agent, wherein the first active agent is a 2-oxazolidinone and/or N-hydroxyethyl-2-oxazolidinone, and the second active agent is arabinose. In at least one embodiment, the crosslinking formulation comprises a cosmetically acceptable carrier. In at least one embodiment, the crosslinking formulation is in liquid form. In at least one embodiment, the crosslinking formulation comprises from about 0.1% to about 99%, or from about 1% to about 98%, or from about 10% to about 97%, or from about 30% to about 95% water. In at least one embodiment, the crosslinking formulation comprises a conditioning agent. Conditioning agents has been described extensively herein and these features are fully compatible and combinable with the 5th aspect. In at least one embodiment, the crosslinking formulation has a pH of from about pH 6 to about pH 10. In at least one embodiment, the conditioning agent is selected from the group consisting of: epoxyaminosilane copolymers, and polysiloxane/polyurea block copolymers, and mixtures thereof. In at least one embodiment, the crosslinking formulation comprises from about 0.01% to about 10% by weight, or from about 0.1% to about 8%, or from about 0.1% to about 5% hairstyling polymer. In at least one embodiment, the crosslinking formulation is a solid.

The kit comprises a photocatalyst formulation comprising a photocatalyst, wherein the photocatalyst is a hydroxy-substituted aromatic compound. Photocatalysts agent have been described extensively herein and these features are fully compatible and combinable with the 5th aspect. In at least one embodiment, the hydroxy-substituted aromatic compound is multicyclic. In at least one embodiment, the hydroxy-substituted aromatic compound is selected from the group consisting of: 8-quinolinol-1-oxide, 8-hydroxyquinoline, Eosin Y, 4,5-dibromofluorescein, and mixtures thereof. In at least one embodiment, the photocatalyst formulation is in solid form.

In at least one embodiment, the kit comprises an appliance for mechanically shaping keratin fibres. In at least one embodiment, the appliance is a hair brush configured to incorporate

LEDs. In at least one embodiment, the appliance is hybrid heat- and light-providing hair straightening irons.

In an embodiment, a crosslinking composition comprises a first, second, and third crosslinking agent. The first crosslinking agent may be arabinose, the second may be ethylene carbonate, and the third may be 1,2-bis(2-maleimidoethoxy)ethane (CAS#115597-84-7). All the description in relation to the other aspects is suitable for and combinable with this aspect as well.

In an embodiment, the photocatalyst is selected from the group consisting of: sulfonated pyrene compounds, onium salts, diazomethane derivatives, bissulfone derivatives, disulfuno derivatives, nitrobenzyl sulfonate derivatives, sulfonic acid ester derivatives, sulfonic acid esters of an N-hydroxyimide, glyoxime derivatives, and mixtures thereof. In certain other embodiments, the photocatalyst is 8-hydroxy-1,3,6-pyrenetrisulfonic acid trisodium salt (D&C Green 8). In at least one embodiment of the alternative aspect, the photocatalyst is a photobase. Photobase catalysts may include derivatives of trityl alcohols such as, for example, Malachite green. Photobase catalysts may also include acridine derivatives such as, for example, 9-hydroxy-10-methyl-9-phenyl-9,10-dihydroacridine. Photobase catalysts may also include photoactive carbamate-containing compounds.

EXAMPLES

The following examples further describe and demonstrate embodiments within the scope of the method and composition described herein. The examples are given solely for the purpose of illustration, and are not to be construed as limitations since many variations thereof are possible without departing from its scope.

Table 1: Example crosslinking compositions

| | A | B | C | D | E(*) |
|--------------------------------|------|-------|----|------|------------|
| 2-oxazolidinone | 0.1 | - | - | 2 | 74.9 (15) |
| N-methyl-2-oxazolidinone | - | 40 | - | - | - |
| N-hydroxyethyl-2-oxazolidinone | - | - | 20 | 4 | - |
| 4',5'-dibromofluorescein | 0.01 | - | - | 0.01 | 0.5 (0.01) |
| 8-hydroxyquinoline | - | 0.005 | - | 0.01 | - |
| 7-cyano-2-naphthol | - | 0.005 | - | - | - |

| | | | | | |
|--|------|------|------|-----|------------|
| 8-quinolinol-1-oxide | - | 0.01 | 0.01 | - | - |
| Eosin Y | 0.01 | - | 0.01 | - | 0.5 (0.01) |
| Preservatives ¹ | 1 | 1 | 1 | 1 | 5 (1) |
| Fragrance | 2 | 2 | 2 | 2 | 10 (2) |
| Viscosity-modifying agent ² | 2 | 2 | 2 | 2 | 10 (2) |
| Conditioning agent ³ | 1 | 1 | 1 | 1 | - |
| Deionised water buffered at pH 10 ⁴ | QSP | QSP | QSP | QSP | - (QSP) |

KEY: ¹ = sodium benzoate; ² = Acusol 823 from Rohm and Haas; hydrophobically modified alkali soluble acrylic polymer emulsion (HASE); ³ = epoxyaminosilane copolymer as described in EP2214633B1 (filing date 30th Oct 2008) available from Momentive™ Performance Materials Inc., Columbus, Ohio, USA; ⁴ = pH 10 buffer is a standard pH calibration buffer available from suppliers such as VWR, alternatively one can use water and modify the pH. Crosslinking composition E is in solid form; * = wt% after mixing in deionised water.

Data

The hair straightening efficacy is tested for the crosslinking compositions described herein. Switches of low lift naturally curly hair are employed. These are shampooed with a Pantene clarifying shampoo to ensure the hair is in a clean state with no residues that could affect the end result. The switches are then rinsed. Excess water is removed from the hair by wringing out the switches. The switches are treated with a crosslinking composition which is pre-prepared in a dark room. The crosslinking composition comprises: active agent as listed in TABLE 2; 100 ppm 4,5-dibromofluorescein; 100 ppm Eosin Y; QSP water buffered at pH 10. These ingredients are mixed on a spinner plate for 30 mins. The crosslinking composition is stored in an amber bottle or a bottle covered in electrical tape to ensure no light access to the crosslinking composition. 0.5 g of crosslinking composition per 1 g hair is employed. The crosslinking composition is left on the hair for 30 minutes. After this time, the hair is blow dried, brushed, and is then mechanically straightened with hybrid light and heat emitting straightening irons (at 250°F [121°C] and emitting UV light at 380 nm) with 8 passes.

The switches are then imaged. To simulate durability, the switches are then given one wash-and-dry cycle. One wash-and-dry cycle involves shampooing with a Hairtrition shampoo (Hairtrition Color Protect sulfate-free shampoo from Zotos), rinsing and then drying in a hot box.

Once dry the switches are imaged again. The switches are then given 4 further wash-and-dry cycles. Once dry the switches are imaged again. An expert grader gives the images of the switches a score on a 0-10 scale. The scale is a standard scale set as curly hair having 4-5 nodes of curls is a score of 0 and very straight hair is a 10. Thus, the switches are compared to a normal state of hair.

Using an expert grader is reliable because the grader is trained on measuring/scaling the configuration of the hair from straight to curly in a consistent way. Numeric data obtained by this method is averaged, and statistical analysis is conducted using JMP statistical software available from SAS Institute at the 87% confidence level. Treatments not containing the same significance grouping letter are significantly different. For example, the 20% N-acetyl-2-oxazolidinone and control treatments contain the letter “B” for their significance grouping, and are not considered to be significantly different from each other.

Table 2: Straightening efficacy results after 5 wash cycles

| Treatment | Significance grouping | Straightening efficacy Mean grade |
|--------------------------------------|-----------------------|--------------------------------------|
| 20% N-(hydroxyethyl)-2-oxazolidinone | A | 4.3 |
| 20% 2-oxazolidinone | A B | 3.7 |
| 20% N-acetyl-2-oxazolidinone | A B | 3.0 |
| Control | B | 2.0 |

As can be seen in Table 2, the composition comprising N-(hydroxyethyl)-2-oxazolidinone delivered significantly better straightening efficacy than the control.

In relation to eyelashes as keratin fibres, the following procedure may be carried out: the eyelashes used are artificial eyelashes that are made from scalp hair glued to a bar and artificially shaped and dried to look like eyelashes. The eyelashes are bent out of shape. Firstly the eyelashes are washed with a clarifying shampoo to get them clean and remove any residue. The eyelashes are soaked in a crosslinking composition consisting of 15% N-hydroxyethyl-2-oxazolidinone, 100 ppm 4,5-dibromofluorescein; 100 ppm Eosin Y; QSP water buffered at pH 10. The eyelashes are submerged overnight in the crosslinking composition. Afterwards, the eyelashes are shaped, for example by pulling the lashes straight with tweezers or by curling them. The eyelashes are

then exposed to light (Visible light) for 90 seconds. A shaping benefit may be seen for the eyelases receiving a crosslinking treatment.

The dimensions and values disclosed herein are not to be understood as being strictly limited to the exact numerical values recited. Instead, unless otherwise specified, each such dimension is intended to mean both the recited value and a functionally equivalent range surrounding that value. For example, a dimension disclosed as “40 mm” is intended to mean “about 40 mm.”

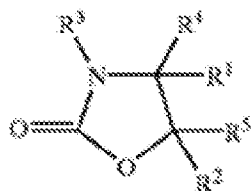
Every document cited herein, including any cross referenced or related patent or patent publication, is hereby incorporated herein by reference in its entirety unless expressly excluded or otherwise limited. The citation of any document is not an admission that it is prior art with respect to any document disclosed or claimed herein or that it alone, or in any combination with any other reference or references, teaches, suggests or discloses any such embodiment. Further, to the extent that any meaning or definition of a term in this document conflicts with any meaning or definition of the same term in a document incorporated by reference, the meaning or definition assigned to that term in this document shall govern.

While particular embodiments have been illustrated and described herein, it should be understood that various other changes and modifications may be made without departing from the spirit and scope of the claimed subject matter. Moreover, although various aspects of the claimed subject matter have been described herein, such aspects need not be utilized in combination. It is therefore intended that the appended claims cover all such changes and modifications that are within the scope of the claimed subject matter.

CLAIMS

What is claimed is:

1. A method for shaping keratin fibres comprising:
 - (a) providing a crosslinking composition, wherein the crosslinking composition comprises:
 - (i) a 2-oxazolidinone having the structure:



wherein:

R^1 and R^2 independently of one another are selected from hydrogen, hydroxyl, phenyl, and C_1 - C_6 -alkyl;

R^3 is selected from hydrogen, C_1 - C_{12} -alkyl, N-hydroxy- (C_2-C_6) -alkyl, C_1 - C_{12} -alkenyl, and C_6 - C_{12} -aryl; and

R^4 and R^5 independently of one another are selected from C_1 - C_{12} -alkyl, C_1 - C_{12} -alkenyl, C_6 - C_{12} -aryl, C_1 - C_{12} -alkoxy, and hydrogen;

(ii) a photocatalyst, wherein the photocatalyst is a hydroxy-substituted aromatic compound;

(iii) a cosmetically acceptable carrier;

wherein the crosslinking composition has not been exposed to electromagnetic radiation having a wavelength of 750 nm or less for 30 min or more;

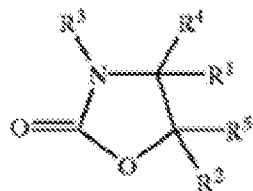
(b) applying the crosslinking composition to keratin fibres; and

(c) mechanically shaping the keratin fibres with an appliance at a temperature of from 80°C to 180°C while exposing the crosslinking composition to electromagnetic radiation having a wavelength of from 300 nm to 750 nm.

2. The method according to Claim 1, wherein the mechanical shaping of keratin fibres is straightening keratin fibres or curling keratin fibres.
3. The method according to Claims 1 or 2, wherein the temperature of the appliance is from 80°C to 170°C.

4. The method according to any preceding claim, wherein the appliance comprises light-emitting diodes.
5. The method according to any preceding claim, wherein the photocatalyst can be activated to a photo-excited state by excitation with incident radiation with a wavelength from 300 nm to 750 nm and wherein the appliance emits radiation having a wavelength from 300 nm to 750 nm.
6. The method according to any preceding claim, wherein the hydroxy-substituted aromatic compound is selected from the group consisting of 8-hydroxyquinoline, 4',5'-dibromofluorescein, Eosin Y, and combinations thereof.
7. The method according to any preceding claim, wherein the crosslinking composition is substantially free of formaldehyde, derivatives of formaldehyde, formalin, and any compound that produces formaldehyde upon heating.
8. The method according to any preceding claim, wherein the method does not comprise a rinsing step.
9. The method according to any preceding claim, wherein the crosslinking composition comprises from 1% to 30% N-hydroxyethyl-2-oxazolidinone, preferably from 10% to 25% N-hydroxyethyl-2-oxazolidinone, more preferably from 13% to 22% N-hydroxyethyl-2-oxazolidinone, by weight of the crosslinking composition.
10. The method according to any preceding claim, wherein the crosslinking composition comprises from 10% to 25% of the 2-oxazolidinone, by weight of the crosslinking composition.
11. The method according to any preceding claim, wherein the crosslinking composition comprises from 15% to 20% of the 2-oxazolidinone, by weight of the crosslinking composition.
12. The method according to any preceding claim, wherein the crosslinking composition comprises from 15% to 20% N-hydroxyethyl-2-oxazolidinone, by weight of the crosslinking composition.
13. The method according to any preceding claim, wherein the crosslinking composition further comprises from 2% to 18% of a reducing sugar selected from the group consisting of arabinose, xylose, ribose, and combinations thereof, by weight of the crosslinking composition.

14. The method according to any preceding claim, wherein the crosslinking composition further comprises 1,2-bis(2-maleimidoethoxy)ethane.
15. A crosslinking composition for shaping keratin fibres comprising:
- (i) a 2-oxazolidinone having the structure:



wherein:

R^1 and R^2 independently of one another are selected from hydrogen, hydroxyl, phenyl, and C_1 - C_6 -alkyl;

R^3 is selected from hydrogen, C_1 - C_{12} -alkyl, N-hydroxy-(C_2 - C_6)-alkyl, C_1 - C_{12} -alkenyl, and C_6 - C_{12} -aryl; and

R^4 and R^5 independently of one another are selected from C_1 - C_{12} -alkyl, C_1 - C_{12} -alkenyl, C_6 - C_{12} -aryl, C_1 - C_{12} -alkoxy, and hydrogen;

(ii) a photocatalyst, wherein the photocatalyst is a hydroxy-substituted aromatic compound; and

(iii) a cosmetically acceptable carrier.

INTERNATIONAL SEARCH REPORT

International application No
PCT/US2016/037959

A. CLASSIFICATION OF SUBJECT MATTER
INV. A61K8/49 A61Q5/06
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, 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,P | WO 2015/094758 A1 (PROCTER & GAMBLE [US]) 25 June 2015 (2015-06-25) page 32; claims 1-15; examples B-E page 30, lines 31-32 page 28, lines 17-32 page 9, line 26 - page 10, line 27 | 1-10,15 |
| X,P | WO 2015/094788 A1 (PROCTER & GAMBLE [US]) 25 June 2015 (2015-06-25) the whole document | 1 |
| X | US 4 382 765 A (MOELLER HINRICH [DE] ET AL) 10 May 1983 (1983-05-10) | 15 |
| Y | examples 3,8 | 1-14 |
| | ----- -/-- | |



Further documents are listed in the continuation of Box C.



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 application or patent 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

12 September 2016

Date of mailing of the international search report

19/09/2016

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

Skulj, Primoz

INTERNATIONAL SEARCH REPORT

International application No

PCT/US2016/037959

C(Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

| Category* | Citation of document, with indication, where appropriate, of the relevant passages | Relevant to claim No. |
|-----------|--|-----------------------|
| Y | WO 2014/123805 A1 (LUBRIZOL ADVANCED MAT INC [US]) 14 August 2014 (2014-08-14) paragraphs [0015], [0103] - [0108], [0111], [0142], [0083], [0094], [0095]; claims 1,13; table 6 ----- | 1-14 |
| X | WO 2009/140076 A1 (PROCTER & GAMBLE [US]; BAKER ELLEN SCHMIDT [US]; CONSTANTINIDES IOANNI) 19 November 2009 (2009-11-19) | 1-14 |
| Y | page 5; figure 5 page 13, lines 28-31 page 17, line 21 - page 20, line 8; figure 3; example 81 ----- | 1-14 |
| A | WO 99/42494 A1 (BASF AG [DE]; FUNK RUEDIGER [DE]; FRENZ VOLKER [DE]; RIEGEL ULRICH [DE]) 26 August 1999 (1999-08-26) ----- | 1 |

INTERNATIONAL SEARCH REPORT

Information on patent family members

International application No

PCT/US2016/037959

| Patent document cited in search report | Publication date | Patent family member(s) | Publication date |
|---|---------------------|--|--|
| WO 2015094758 A1 | 25-06-2015 | US 2015174028 A1 WO 2015094758 A1 | 25-06-2015 25-06-2015 |
| WO 2015094788 A1 | 25-06-2015 | CN 105873563 A US 2015174432 A1 WO 2015094788 A1 | 17-08-2016 25-06-2015 25-06-2015 |
| US 4382765 A | 10-05-1983 | AT 357681 B BE 863635 A1 DE 2704904 A1 FR 2379283 A1 GB 1574401 A IT 1092415 B JP S5399335 A NL 7800444 A US 4382765 A | 25-07-1980 03-08-1978 10-08-1978 01-09-1978 03-09-1980 12-07-1985 30-08-1978 08-08-1978 10-05-1983 |
| WO 2014123805 A1 | 14-08-2014 | AU 2014215569 A1 CA 2899597 A1 CN 105073090 A EP 2953610 A1 JP 2016507528 A KR 20150113206 A US 2015374604 A1 WO 2014123805 A1 | 13-08-2015 14-08-2014 18-11-2015 16-12-2015 10-03-2016 07-10-2015 31-12-2015 14-08-2014 |
| WO 2009140076 A1 | 19-11-2009 | CA 2724264 A1 CN 102026614 A EP 2273968 A1 EP 2727578 A2 JP 5592354 B2 JP 2011523413 A JP 2014196313 A US 2009285768 A1 WO 2009140076 A1 | 19-11-2009 20-04-2011 19-01-2011 07-05-2014 17-09-2014 11-08-2011 16-10-2014 19-11-2009 19-11-2009 |
| WO 9942494 A1 | 26-08-1999 | AU 2927199 A BR 9908006 A CA 2320778 A1 CN 1291202 A DE 19807502 A1 EP 1056787 A1 ES 2205788 T3 ID 25873 A JP 4210432 B2 JP 2002504566 A MY 124251 A US 6472478 B1 WO 9942494 A1 | 06-09-1999 24-10-2000 26-08-1999 11-04-2001 16-09-1999 06-12-2000 01-05-2004 09-11-2000 21-01-2009 12-02-2002 30-06-2006 29-10-2002 26-08-1999 |