

**TREATMENT AND/OR PREVENTION OF INFLAMMATION AND
CUTANEOUS PHOTODAMAGE AND PHOTOPROTECTION OF THE SKIN
WITH A WATER-SOLUBLE EXTRACT FROM PLANT OF *SOLANUM*
GENUS**

5 This application claims priority of Taiwanese application no. 100116659,
filed on May 12, 2011.

 The present invention relates to a pharmaceutical composition for
preventing and treating inflammation and cutaneous photodamage, which
contains a water-soluble extract from a plant of *Solanum* genus. The
10 composition also has a photoprotective effect, and can be used as a cosmetic
composition.

 In aerobic organisms, oxygen is utilized in aerobic respiration, thereby
producing reactive oxygen species (ROS) and free radicals. Ultraviolet (UV),
ionizing radiation, and certain medications or xenobiotics will also stimulate
15 production of ROS and free radicals. ROS and free radicals are likely to react
with components within the cell (e.g., DNA, protein and lipids etc.) due to
instability thereof, thereby resulting in oxidative damage to cells and tissues.

 In general, antioxidant enzymes in organisms form an interacting
network to protect cells and tissues from oxidative damage. Oxidative stress
20 forms when ROS and free radicals exceed the antioxidant capacity provided
by the interacting network of the cells or tissues. It has been reported that
oxidative stress plays an important role in the pathological process of
inflammation and photodamage (Simon R. *et al* (2010), *Free radical biology &*
Medicine, 49:1603-1616; Afaq F. *et al.* (2006), *Experimentla Dermatology*,
25 15:678-684).

 Inflammation is a protective response of cells or tissues to pathogen and
external stressful stimuli. During inflammation, the cells or tissues at a site of

injury will stimulate the expression of specific genes through NF- κ B, followed by an increased expression of chemokines, thereby leading to the accumulation of polynuclear leukocytes, monocytes, macrophages and mast cells at the site of injury (i.e., infiltration). The recruited macrophages will be
5 activated by lipopolysaccharides (LPS) that are expressed on a surface of a pathogen. The activated macrophages will induce the expression of proinflammatory genes (including cyclooxygenase-2 gene, COX-2, and inducible nitric oxide synthase gene, *iNOS*) to reinforce inflammatory response. In addition, the activated macrophages release ROS and free
10 radicals to kill pathogens. However, prolonged inflammatory response leads to oxidative stress and damage due to excess accumulation of ROS and free radicals, thereby resulting in chronic inflammation and ultimately potentiating the possibility of chronic illnesses or cancer.

Photodamage occurs when skin of an organism is exposed to ultraviolet
15 (especially ultraviolet-B, UV-B), leading to skin damage. Exposure to UV radiation will accelerate the accumulation of ROS and free radicals in skin cells, increase oxidative stress to the skin cells and induce expression of matrix metalloproteinases (MMP), thereby resulting in oxidative photodamage. Symptoms of oxidative photodamage include: telangiectasia,
20 thinning of epidermis, reduction in collagen fiber and elastic fiber, dryness, wrinkle formation, inflammatory cell infiltration, premature skin aging and skin pathologic change.

In recent years, phytochemicals or phytochemopreventive agents derived from plants have been proven to have antioxidant, anti-inflammatory
25 properties and improve photodamage. Examples of the phytochemical include: green tea polyphenols (GTPs), epigallocatechin gallate (EGCG), genistein, resveratrol, curcumin, apigenin, lycopene, etc. (Adhami V.M. *et al*

(2008), *Photochem. Photobiol.*, 84:489-500). The phytochemicals have been proven to be safer for clinical application without undesired side effects, and have caught attention in the field of medical research.

Plants of *Solanum* genus include *Solanum incanum* L., synonymous with
5 *Solanum undatum*, *Solanum incanum* Ruiz. & Pav., *Solanum coagulans*
Forsskal ; bitter apple in English), *Solanum indicum*, *Solanum nigrum* (Long
kui in Chinese ; black nightshade in English), *Solanum capsicastrum* (false
jerusalem cherry in English), *Solanum xanthocarpum*, *Solanum melongena*,
Solanum coagulans, *Solanum tuberosum*, *Solanum sodomium* (apple of
10 Sodom in Australia), *Solanum turburosum*, *Solanum aculeastrum*, *Solanum*
lycocalpum, *Solanum khasianum*, *Solanum suaveolens*, *Solanum uporo*,
Solanum abutiloides, *Solanum coccineum*, *Solanum unguiculatum*, *Solanum*
robustum, *Solanum anguivi*, *Solanum platanifolium*, *Solanum mammosum*,
etc. It is known that steroidal alkaloids can be extracted from the *Solanum*
15 genus, and commonly include solasonine and solamargine.

It is disclosed in JP 2010/2753593 a dry tomato powder from *Solanum*
Jycopersium having the ability to treat dermatitis, which includes chapped
skin, cracked skin, dry skin, rash or other inflammatory skin disease. The
active component of the tomato powder is identified as Esculeoside A and B.

20 It is disclosed in several publications that solasonine and solamargine
are able to treat skin diseases. For example, WO 2004/002497 A1 discloses
pharmaceutical compositions comprising at least two glycoalkaloids, such as
solamargine and solasonine. Such composition may be used in the treatment
of cancer and psoriasis. It is disclosed that solutions of solamargine and
25 solasonine were prepared by the same in 3% acetic acid, suggesting the
inability of the glycoalkaloids to dissolve in pure water or water with neutral
pH.

WO 03/029269 A1 discloses the extraction of solasonine and solamargine from the fruit of *Solanum* genus. It is clear from the extraction procedures that the glycoside extract obtained is insoluble in water since the precipitate of the extract obtained during the extraction can be washed with
5 water.

US 7,078,063 B2 issued to the inventors of the present invention discloses a water-soluble extract from a plant of *Solanum* genus, especially *Solanum* incanum L., which includes at least 60% of solasonine and solamargine., and a method for preparing the water-soluble extract. The
10 patent is hereby incorporated by reference in its entirety.

In the aforesaid US patent, the inventors found that the water-soluble extract can inhibit the growth of tumor/cancer cells (specifically liver tumor cells, lung cancer cells, and breast cancer cells). In this invention, the inventors unexpectedly found that the water-soluble extract effectively cure
15 and/or prevent inflammation and cutaneous photodamage.

Therefore, according to the first aspect, this invention provides a use of a water-soluble extract from a plant of *Solanum* genus in the preparation of a pharmaceutical composition for preventing and treating cutaneous photodamage, wherein the water-soluble extract comprises at least 60% of
20 solamargine and solasonine.

In the second aspect, this invention provides a cosmetic method of preventing and treating cutaneous photodamage, the method comprising administering a composition comprising a water-soluble extract from a plant of *Solanum* genus, wherein the water-soluble extract comprises at least 60%
25 of solamargine and solasonine.

In the third aspect, this invention provides a use of a water-soluble extract from a plant of *Solanum* genus in the preparation of a pharmaceutical

composition for preventing and treating inflammation caused by exposure to sunlight or UV light, wherein the water-soluble extract comprises at least 60% of solamargine and solasonine.

Other features and advantages of the present invention will become apparent in the following detailed description of the preferred embodiments of the invention, with reference to the accompanying drawings in which:

Figure 1 is a western blot, showing COX-2 expression in RAW264.7 cell
5 after LPS-induced inflammatory response. The cells in normal control group were not treated with LPS and a water-soluble extract from a plant of *Solanum* genus, whereas cells in pathological control group were treated with 100ng/mL of LPS. Experimental groups 1-6 were treated with various concentrations of the water-soluble extract (0.01, 0.05, 0.1, 0.5, 1 and 5 μ
10 g/mL) along with 100ng/mL of LPS;

Figure 2 shows transepidermal water loss (TEWL) of dorsal skin of HRS/J hairless mice in each of normal control group, pathological control group, and experimental groups 1 and 2 after 6 weeks of UV-B radiation. HRS/J hairless mice in normal control group did not receive any treatment
15 including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. In experimental group 1, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. solution containing the water-soluble extract. In experimental
20 group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract;

Figure 3 is a western blot showing expressions of NF-kB, COX-2 and iNOS from dorsal skin tissue of HRS/J hairless mice after the 60 week experimental period. HRS/J hairless mice in normal control group did not
25 receive any treatment including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. In experimental

group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract;

Figure 4 shows an immunohistochemistry of NF-kB expression on dorsal skin tissue from HRS/J hairless mice at the end of 60 weeks after the first time of UV-B radiation. HRS/J hairless mice in normal control group did not receive any treatment including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. In experimental group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract;

Figure 5 shows a toluidine blue stain of dorsal skin tissue from HRS/J hairless mice at the end of the 60week experimental period. HRS/J hairless mice in normal control group did not receive any treatment including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. In experimental group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. Arrows indicate the locations of infiltrated mast cells;

Figure 6 is a bar diagram showing average mast cell number from dorsal skin tissue of HRS/J hairless mice in normal control group, pathological control group, and experimental group 2 after the 60 week experimental period. HRS/J hairless mice in normal control group did not receive any treatment including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. In experimental group 2,

HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. “*” indicates $p < 0.05$ between experimental group 2 and normal control group. “****” indicates $p < 0.01$ between experimental group 2 and pathological control group. “###” indicates $p < 0.001$ between pathological control group and normal control group;

Figure 7 is a plot showing average tumor number in HRS/J mice formed at various time points during the 60 week experimental period. HRS/J hairless mice in pathological control group had photodamage induced by UV-B radiation and were not treated with the water-soluble extract. HRS/J hairless mice in experimental group 1 were exposed to UV-B radiation and were treated with *Solanum incanum* L. solution containing the water-soluble extract. In experimental group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. “*” indicates $p < 0.05$ between experimental group 2 and pathological control group. “****” indicates $p < 0.01$ between experimental group 2 and pathological control group;

Figure 8 is a plot showing average tumor size in HRS/J hairless mice at various time points during the 60 week experimental period. HRS/J hairless mice in pathological control group had photodamage induced by UV-B radiation and were not treated with the water-soluble extract. HRS/J hairless mice in experimental group 1 were exposed to UV-B radiation and were treated with *Solanum incanum* L. solution containing the water-soluble extract. HRS/J hairless mice in experimental group 2 were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. “#” indicates $p < 0.05$ between experimental group 1 and pathological control group. “##” indicates $p < 0.01$ between experimental

group 1 and pathological control group. "###" indicates $p < 0.001$ between experimental group 1 and pathological control group. "***" indicates $p < 0.01$ between experimental group 2 and pathological control group. "****" indicates $p < 0.001$ between experimental group 2 and pathological control group;

5 Figure 9 is a bar diagram showing recovery time in pinch test on HRS/J hairless mice after the 60 week experimental period. HRS/J hairless mice in normal control group did not receive any treatment including UV-B radiation and the water-soluble extract. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the
10 water-soluble extract. In experimental group 2, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. "****" indicates $p < 0.001$ between experimental group 2 and pathological control group. "###" indicates $p < 0.001$ between pathological control group and normal control group;

15 Figure 10 shows images of elastic fibers of dorsal skin tissue stained with resorcin-fuchsin solution, Weigert's iron hematoxylin and van Gieson solution. HRS/J hairless mice in pathological control group had photodamage induced by UV-B radiation and were not treated with the water-soluble extract. In experimental group 2, HRS/J hairless mice were exposed to UV-B
20 radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. Arrows indicate the locations of elastic fiber denaturation;

 Figure 11 shows average tumor number in HRS/J hairless mice over the experimental period. HRS/J hairless mice were exposed to UV-B radiation
25 and were treated with AIROL vanishing cream in tretinoin group. In *Solanum incanum* L. gel group, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble

extract;

Figure 12 is a graph showing transepidermal water loss (TEWL) over the experimental period. HRS/J hairless mice in pathological control group were exposed to UV-B radiation and were not treated with the water-soluble extract. HRS/J hairless mice in tretinoin group were exposed to UV-B radiation and were treated with AIROL vanishing cream. In *Solanum incanum* L. gel group, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. “*” indicates $p < 0.05$ between tretinoin group and *Solanum incanum* L. gel group; and

Figure 13 is a graph showing water content of dorsal skin from HRS/J hairless mice over the experimental period. HRS/J hairless mice in pathological control group had photodamage induced by UV-B radiation and were not treated with the water-soluble extract. In tretinoin group, HRS/J hairless mice were exposed to UV-B radiation and were treated with AIROL vanishing cream. In *Solanum incanum* L. gel group, HRS/J hairless mice were exposed to UV-B radiation and were treated with *Solanum incanum* L. gel containing the water-soluble extract. “*” indicates $p < 0.05$ between tretinoin group and *Solanum incanum* L. gel group.

For the purpose of this specification, it will be clearly understood that the word “ comprising ” means “ including but no limited to ”, and that the words “ comprises, ” “ contain ” and variants thereof have a corresponding meaning.

Unless otherwise defined, all technical and scientific terms used herein have the meaning commonly understood by a person skilled in the art to

which this invention belongs. One skilled in the art will recognize many methods and materials similar or equivalent to those described herein, which could be used in the practice of this invention. Indeed, this invention is in no way limited to the methods and materials described. For clarity, the following
5 definitions are used herein.

The present invention provides a pharmaceutical composition for preventing and treating inflammation, which includes a water-soluble extract from a plant of *Solanum* genus. The water-soluble extract comprises at least 60% of solamargine and solasonine. Preferably, the water-soluble extract
10 comprises at least 60%-90% of solamargine and solasonine.

A process for preparing the water-soluble extract has been disclosed in US 7,078,063 B2, and includes the following steps:

- (a) subjecting a plant material of a plant of *Solanum* genus to an extraction treatment using an acidic aqueous solution with a pH value of 3~5, such
15 that an aqueous solution is obtained;
- (b) adjusting the pH value of the aqueous solution obtained in step (a) to pH 8~10 with a base, such that a precipitate is formed;
- (c) washing the precipitate formed in step (b) with water, followed by drying, such that a dried product is obtained;
- 20 (d) admixing the dried product obtained in step (c) with chloroform, followed by addition of a suitable amount of a 100% alcohol, such that a chloroform-alcohol mixture is formed;
- (e) mixing the chloroform-alcohol formed in step (d) with a water/alcohol solution having a predetermined water/alcohol ratio, such that a mixture
25 containing a chloroform-based layer and a non-chloroform-based layer is obtained;
- (f) removing the chloroform-based layer from the mixture obtained in step (e),

followed by addition of a suitable amount of water; and

(g) obtaining a supernatant from the resultant mixture of step (f), followed by drying the supernatant, wherein the resultant dried product is able to be directly dissolved in water to form a yellowish clear and transparent aqueous solution.

5 Preferably, the water-soluble extract of a plant of *Solanum* genus has been obtained at least from the fruit, root, stem and leaf of the plant of *Solanum* genus. The plant of *Solanum* genus has been chopped in a preliminary treatment. In a preferred embodiment of this invention, the plant material used in step (a) is the fruit of the plant of *Solanum* genus.

The inventors found that certain factors might affect the content and proportion of solasonine and solamargine in the water-soluble extract obtained using the aforesaid process. These factors include the species of the plant of *Solanum* genus and the part(s) of the plant used in the extracting process, as well as the types of alcohol and base used. Therefore, a skilled artisan can prepare a desired water-soluble extract by selecting suitable species of the plant, in conjunction with appropriate operating conditions.

20 Preferably, the water-soluble extract is obtained from a plant of *Solanum* genus selected from the group consisting of *Solanum incanum* L., *Solanum indicum*, *Solanum nigrum*, *Solanum capsicastrum*, *Solanum xanthocarpum*, *Solanum melongena*, *Solanum coagulans*, *Solanum tunigrum*, *Solanum sodomeum*, *Solanum turburosum*, *Solanum aculeastrum*, *Solanum lycocarpum*, *Solanum khasianum*, *Solanum suaveolens*, *Solanum uporo*, *Solanum abutiloides*, *Solanum coccineum*, *Solanum unguiculatum*, *Solanum robustum*, *Solanum anguivi*, *Solanum platanifolium* and *Solanum mammosum*. In a preferred embodiment of this invention, the water-soluble extract is obtained from *Solanum incanum* L..

In preliminary experiments, *in vitro* data have shown that the water-soluble extract from *Solanum* genus can inhibit inflammatory response induced by LPS. It has also been proven that, after hairless mice with UV-B induced inflammation are treated with the water-soluble
5 extract from *Solanum* genus, the expression of NF- κ B, COX-2 and iNOS on skin tissue thereof was markedly decreased, together with an alleviation of mast cell infiltration and inflammatory response.

Therefore, this invention provides a method for preventing and treating inflammation, comprising applying to a subject in need of such
10 treatment the water-soluble extract or the pharmaceutical composition.

The administration route of the aforesaid pharmaceutical composition provided by this invention comprises oral, topical and parenteral routes.

The pharmaceutical composition provided by this invention can be
15 formulated into an oral dosage form using technology well known to a skilled artisan. Examples of the oral dosage form include: aseptic power, tablet, troche, lozenge, pellet, capsule, dispersible powder or granule, solution, suspension, emulsion, syrup, elixir, and slurry.

The pharmaceutical composition of this invention can further include
20 a pharmaceutically acceptable carrier that is widely employed in drug-manufacturing technology.

The pharmaceutically acceptable carrier comprises one or more reagents, including: solvent, buffer, emulsifier, suspending agent, decomposer, disintegrating agent, dispersing agent, binding agent,
25 excipient, stabilizing agent, chelating agent, preservative, wetting agent, lubricant, diluent, absorption delaying agent, liposome, sweetening agent, flavoring agent, and coloring agent. The choice and amount of the pharmaceutically acceptable carrier are within the expertise of those skilled in the art.

30 The pharmaceutically acceptable carrier comprises one or more

reagents including, for example, water, normal saline, phosphate buffered saline (PBS), glucose containing solution, aqueous solution containing alcohol (for example, ethanol, propanediol, glycol, mannitol), oil (for example, peanut oil, olive oil, sesame oil, castor oil, cottonseed oil, soybean oil), glycerol, organic solvent and liposome. In an embodiment of this invention, the solvent is water.

The pharmaceutical composition according to this invention can be formulated into a suitable dosage form for topical administration using technology well known to those skilled in the art, which includes external preparations, effervescent tablets, suppositories, and the like.

In a preferred embodiment of this invention, the pharmaceutical composition of this invention is formulated into an external preparation in a gel form by admixing the water-soluble extract with a base that is well known and commonly used in the art.

In this invention, the suitable base may include one or more of the following additives: water, alcohols, glycol, hydrocarbons (such as petroleum jelly and white petrolatum), waxes (such as paraffin and yellow wax), preserving agents, antioxidants, surfactants, absorption enhancers, stabilizing agents, gelling agents (such as carbopol®974P, microcrystalline cellulose and carboxymethylcellulose), active agents, humectants, odor absorbers, fragrances, pH adjusting agents, chelating agents, emulsifiers, occlusive agents, emollients, thickeners, solubilizing agents, penetration enhancers, anti-irritants, colorants and propellants. The choice and amount of these additives are within the expertise of those skilled in the art.

The dosage and frequency of administration of this pharmaceutical composition may vary depending on the following factors: the severity of the disease to be treated, the route of administration, and the weight, age, physical condition and response of the subject to be treated. For instance, the daily dosage of the pharmaceutical composition for topical

administration according to this invention may be 10-20 mg/cm² of the lesion area and one to six times per day. The dosage of this pharmaceutical composition for oral administration may be 1-30 mg/Kg, 1-4 times a day.

5 The water-soluble extract from the plant of *Solanum* genus has been proven to alleviate skin pathological change induced by UV-B-induced cutaneous photodamage on hairless mice. In addition, it can effectively improve skin elasticity and solar elastosis, while maintaining moisture-retaining capacity at the same time.

10 Therefore, the present invention provides a pharmaceutical composition for treating and/or preventing cutaneous photodamage, which comprises the water-soluble extract from a plant of *Solanum* genus as described above. The present invention also provides a method for preventing and treating cutaneous photodamage, comprising
15 applying to a subject in need of such treatment the aforesaid water-soluble extract or the pharmaceutical composition.

 In this invention, the term "cutaneous photodamage" indicates skin damage caused by exposure of skin to sunlight or UV light (especially, UV-B). The cutaneous photodamage comprises: thinning of the skin,
20 skin atrophy, decrease in collagen fiber and elastic fiber, elastosis, loss of skin elasticity, dryness, wrinkle formation, inflammatory cell infiltration, premature skin aging, vascular change (such as diffuse erythema, ecchymosis, or telangiectasia), pigmentary change (such as lentigines,

freckles, hypopigmentation or hyperpigmentation), comedone, cyst and skin pathological change.

According to this invention, the administration route, dosage and pharmaceutically acceptable carrier are similar to those used in the pharmaceutical composition for preventing and treating inflammation. In a preferred embodiment of the current invention, the pharmaceutical composition is formulated into topical form for administration onto the skin.

Due to prevention/protection of cutaneous photodamage of the water-soluble extract, one can predict that the water-soluble extract can be used as a cosmetic component to prepare a cosmetic composition.

Therefore, this invention provides a cosmetic composition with photoprotective effect, which comprises the water-soluble extract as described above.

In this invention, the term "photoprotective" indicates to block or alleviate adverse clinical, histological and immunological effects caused by the sun or ultraviolet. These effects include acute reactions (such as erythema and inflammation) and chronic effects (such as elastosis and wrinkle formation).

According to this invention, the cosmetic compositions further comprise cosmetically acceptable adjuvant that is widely employed in cosmetic-manufacturing technology.

The acceptable adjuvant comprises one or more reagents including solvent, gelling agent, activating agents, preservatives, antioxidants, screening agent, chelating agent, surfactants, coloring agent, thickening agent, filler, fragrance and odor absorbent. The choice and amount of these additives are within the expertise of those skilled in the art.

The cosmetic composition provided by this invention can be prepared using technology well known to a skilled artisan into the form of skincare,

haircare or makeup products. The form includes aqueous solution, emulsion, gel, ointment, cream, mask, patch, powder, aerosol, spray, lotion, sunblock and other body cleansing products.

5 The cosmetic composition of this invention can be used with the following agents: whitening agents, humectants, bactericides, ultraviolet absorbers, anti-acne agent, antipruritic, antihyperkeratolytic agents, antipsoriatic agents, antiager, antiwrinkle agents, antiseborrheic agents, self-tanning agents and wound-healing agents. The choice and amount of these agents are within the expertise of those skilled in the art.

10 ***Experimental Materials:***

1. Preparations of water-soluble extract from *Solanum incanum* L. and *Solanum incanum* L. solution containing the water-soluble extract:

15 The water-soluble extract from *Solanum incanum* L. was prepared based on the procedure disclosed in Example 1 of US 7,078,063 B2. Specifically, 500 g of ripe fruit of *Solanum incanum* L. was ground subsequent to addition of 1000 ml pure water. To the resultant aqueous mixture, 99.5% of acetic acid was added dropwise to adjust the pH value to 4.0, followed by shaking at room temperature for 12 hrs. A
20 supernatant was obtained by centrifuging the aqueous mixture, and 33% NH₄OH basic solution was added thereto dropwise to adjust the pH value of the supernatant to 9.0, and a precipitate was formed. The precipitate was obtained by conducting centrifugation at 4,500 rpm (Beckman Coulter, Avanti J-25, JA-14 Rotor), and the residual basic
25 solution present therein was removed by washing the precipitate with water, followed by centrifugation at 4,500 rpm. The precipitate thus obtained was suspended in distilled water and subjected to lyophilization (Virtis, Freezemobile 12ES) to get 5 g of dried powder.

2 g of the dried powder was dissolved in 50 ml chloroform in reagent

grade, followed by addition of 40 ml of 100% methanol and was shaken to form a uniform suspension. A supernatant was obtained by centrifugation at 4,500 rpm or filtration. 70 ml of methanol:water solution (1:1) was added to the supernatant and mixed well. The mixture obtained was centrifuged at
5 12,000 rpm for 10 min. The resultant supernatant was taken out, and 120 ml distilled water was added thereto and shaken well. Meanwhile, the supernatant became murky. The supernatant was further centrifuged at 12,000 rpm for 10 min so as to remove the precipitate. The resultant supernatant was subjected to decompress concentrating under reduced
10 pressure at 55° C to remove methanol, followed by lyophilization to obtain dried powder of the water-soluble extract.

The water-soluble extract was dissolved in sterilized water to obtain a *Solanum incanum* L. solution for further use in experiments described below.

**2. Preparation of *Solanum incanum* L. gel containing the water-soluble
15 extract:**

4 g of carbopol®974P, which was used as a gelling agent and is commercially available from Lubrizol Advanced Materials, Inc., KY 40258, USA, was dissolved in 50g pure water, followed by sequentially adding 30g propylene glycol and 7 g of the dried powder of the water-soluble extract
20 obtained from the section of "1. Preparations of water-soluble extract from *Solanum incanum* L. and *Solanum incanum* L. solution containing the water-soluble extract" under "Experimental Materials" and mixing well. The mixture was heated in a heating vessel at a temperature of 50°C-60°C for 20 minutes, followed by cooling at room temperature. The cooled mixture was
25 added with triethanolamine to adjust pH to 7.0±0.5. Subsequently, water was added until the total weight of the mixture reached 100g, thereby obtaining a gel containing 7% (w/w) of the water-soluble extract (hereinafter

referred to as *Solanum incanum* L. gel).

3. Animal model

HRS/J hairless mice (6-8 weeks old, body weight 20-22g) were purchased from Jackson Laboratory (Bar Harbor, USA). The mice were kept
5 in a room with 12 hr/12 hr light/dark cycle, temperatures of 21-22°C, and 30-70% humidity. Diet and water were sufficient and accessible to the mice at all times. All animal experiments were conducted according to Guide for the Care and Use of Laboratory Animals of National Institute of Health (NIH).

General Methods:

10 1. Analysis of protein products:

Sodium dodecyl sulfate-polyacrylamide gel electrophoresis (SDS-PAGE) and western blot were used for protein analysis in this invention. Apparatus and reagents for SDS-PAGE and western blot are as follows:

- 15 a. Vertical electrophoresis system (Hoefer SE600, GE Healthcare) were used for SDS-PAGE analysis.
- b. Polyvinylidene difluoride membrane (PVDF, Millipore) and semi-dry blotters (Hoefer TE70X, GE Healthcare) were used for protein transfer.
- c. Primary and secondary antibodies used in western blot method are listed in Table 1.
- 20 d. Proteins were visualized by chemiluminescence staining using Immobilon™ Western Chemiluminescent HRP Substrate (Millipore, Cat No. WBKLS0500), followed by detection on autoradiography films (Kodak Biomax, Kodak Cat. No. 1788207).

Table 1. Primary and secondary antibodies used in western blot analysis

Target protein	Primary antibody	Secondary antibody
COX-2	Mouse anti COX-2 monoclonal antibody (BD Bioscience, Cat. No. 610203)	Sheep anti mouse IgG-horseradish peroxidase (HRP) antibody (Amersham, Cat. No. NA931)
NF- κ B	Rabbit anti NF- κ B monoclonal antibody (Abcam, Cat. No. Ab7970)	Donkey anti rabbit IgG-HRP antibody (Amersham, Cat. No. NA934)
iNOS	Rabbit anti iNOS polyclonal antibody (Santa Cruz, Cat. No. Sc-651)	Donkey anti rabbit IgG-HRP antibody
β -actin	Mouse anti β -actin monoclonal antibody (Sigma, Cat. No. A5441)	Sheep anti mouse IgG-HRP antibody
GAPDH	Mouse anti GAPDH monoclonal antibody (Chemicon, Cat. No. Mab374)	Sheep anti mouse IgG-HRP antibody

2. Induction of inflammation and photodamage:

Dorsal skin of each of the HRS/J mice was exposed to UV-B radiation three times a week for 14 weeks to induce inflammatory response and photodamage. The UV-B radiation was carried out using BLX-312 ultraviolet crosslinker (BIO-LINK, Vilber Lourmat, France), which is equipped with 6x8W T-8M UV-B lamp (312 nm). The single dosage of UV-B radiation for each exposure, the UV-B dosage for one week (dosage/wk), and the cumulative dosage are listed in Table 2.

Table 2.

Experimental time point (wk)	Single UV-B dosage (mJ/cm ²)	Dosage/wk (mJ/cm ²)	Accumulated dosage (mJ/cm ²)
1	36	108	108
2	54	162	270
3	72	216	486
4	90	270	765
5	108	324	1080
6	126	378	1458
7	144	432	1890
8	162	486	2376
9	180	540	2916
10	216	594	3510
11	216	648	4158
12	216	648	4806
13	216	648	5454
14	216	648	6102

3. Tissue slice:

Tissues (harvested at room temperature) were fixed in 4% paraformaldehyde in PBS for at least 12 hrs, followed by ethanol dehydration. The dehydrated tissues were embedded in paraffin and sliced to obtain longitudinal sections.

4. Statistical analysis:

Results are shown as mean $\pm$ standard error mean (SEM). All statistical

data analyses were performed by Student's t-test (2-tailed). $P < 0.05$ is considered statistically significant.

Example 1: Evaluation of *in vitro* anti-inflammatory effect of the water-soluble extract from *Solanum incanum* L.

5 Murine macrophage cell line RAW264.7 that was subjected to lipopolysaccharides (LPS)-induced inflammation was used to determine the anti-inflammatory effect of the water-soluble extract from *Solanum incanum* L.

RAW264.7 cells (purchased from American Type Culture Collection, ATCC, TIB-71) were seeded at a density of 1×10^6 cells/well (6-well plate) in 10 Dulbecco's Modified Eagle's Medium (DMEM) (HyClone SH 30022.01, Logan, Utah, USA) with 10% fetal bovine serum (FBS), 4mM L-glutamine and 4.5 g/L glucose (without sodium pyruvate), and incubated in a 37°C, 5% CO₂ incubator. The cells were divided into three groups: normal control group, 15 pathological control group and experimental groups (experimental groups 1, 2, 3, 4, 5 and 6). After 72hrs of incubation, the pathological control group was changed to be incubated in fresh DMEM medium containing 100ng/mL of LPS (*Escherichia coli* serotype 0111:B4, Sigma), and the experimental groups were incubated in fresh DMEM media containing 100ng/mL of LPS 20 and different concentrations of the water-soluble extracts obtained from the section of "1. Preparations of water-soluble extract from *Solanum incanum* L. and *Solanum incanum* L. solution containing the water-soluble extract" under "Experimental Materials". The concentrations of the water-soluble extracts for experimental groups 1, 2, 3, 4, 5 and 6 were respectively 0.01, 0.05, 0.1, 0.5, 25 1 and 5 µg/mL. The normal control group was still incubated in DMEM with 10% FBS, 4mM L-glutamine and 4.5 g/L glucose (without sodium pyruvate).

After 24hrs of incubation (in a 37°C, 5% CO₂ incubator), the medium in

each well was removed followed by addition of 100 μ L T-PER reagent (purchased from PIERCE) and mixing to obtain a cell mixture. The cell mixture was placed in a centrifugation tube and shaken for 5 min followed by cooling on ice for 30 min. Subsequently, the cell mixture was centrifuged at
5 14,000 rpm for 10 min at 4°C and a supernatant was collected as a protein sample for further analyses.

The resultant protein sample was subjected to COX-2 western blot analysis using the method described in the section of “1. Analysis of protein products” under “General Methods”. β -actin was used as an internal control.

10 Figure 1 is a western blot showing COX-2 expression in RAW264.7 cells. As shown in Figure 1, COX-2 expression was markedly increased in the pathological control group when compared to the normal control group, which indicates a successful induction of inflammation by LPS in RAW264.7 cells. A decreased expression of COX-2 was observed in experimental group 1-6
15 when compared to the pathological control group, and the decreased expression is more obvious when a higher concentration of water-soluble extract was added. These data demonstrate that the water-soluble extract from *Solanum incanum* L. can inhibit LPS-induced inflammation *in vitro*, and the effect of the water-soluble extract on anti-inflammatory response is
20 dose-dependent.

Example 2: Evaluation of the therapeutic effects of the *Solanum incanum* L. solution and the *Solanum incanum* L. gel on hairless mice with UV-B-induced inflammation

HRS/J hairless mice was randomly grouped into normal control group,
25 pathological control group and two experimental groups (i.e. experimental group 1 and 2) (n=4/group). HRS/J hairless mice in normal control group did not receive any UV-B radiation. The mice in pathological control group and

two experimental groups were induced inflammation using the procedure as described in the section of “2. Induction of inflammation and photodamage” under “General Methods”.

Starting from the initiation of UV-B radiation, HRS/J mice in experimental group 1 was administered orally with *Solanum incanum* L. solution, in which the solution was prepared as described in the section of “1. Preparations of water-soluble extract from *Solanum incanum* L. and *Solanum incanum* L. solution containing the water-soluble extract” under “Experimental Materials”. These mice were administered with *Solanum incanum* L. solution once a day (1500 mg/kg), 7 times a week, during a 14-week (wk) period, and were continuously observed for 60 wks (i.e., 60 wk experimental period). On the UV-B exposed dorsal skin of each of the HRS/J hairless mice in experimental group 2, the *Solanum incanum* L. gel obtained from the section of “2. Preparation of *Solanum incanum* L. gel containing the water-soluble extract” under “Experimental Materials” was applied. The gel was applied once a day (dosage: 10-20 mg of gel per cm²), 5 times a week, during a 14-wk period and was observed for 60 wks. HRS/J hairless mice in normal control and pathological control group were not treated with the *Solanum incanum* L. solution nor the *Solanum incanum* L. gel.

After 6 wks of UV-B radiation, HRS-J hairless mice were analyzed for Items A and B listed below. In addition, dorsal skin tissues were collected using surgical scissors from the mice in normal control group, pathological control group and experimental group 2 after the 60 wk experimental period. These tissues were analyzed for Items C to E listed below.

A. Diagnosis of clinical signs:

HRS/J hairless mice were observed for any signs of inflammation, e.g., itching, redness, after 6 wks of UV-B radiation.

B. Analysis of transepidermal water loss (TEWL)

After 6wks of UV-B radiation, TEWL was evaluated on dorsal skin of each HRS/J hairless mouse from each group using an evaporimeter (Tewameter TM 201R, Courage & Khazaka, Cologne, Germany).

5 C. Western blot analysis of NF- κ B, COX-2 and iNOS

0.5 g of skin tissue was added to T-PER tissue protein extraction reagent (Thermo Scientific, Cat. No. 78510), followed by tissue disruption by a homogenizer (purchased from Biospec Products). The well homogenized product was centrifuged at 15,000rpm and 4°C for 10hrs. Supernatant was
10 collected for protein analysis.

The isolated protein was further used to detect NF- κ B, COX-2 and iNOS by western blot analysis according to the procedure set forth in the section of "1. Analysis of protein products" under "General Methods". GAPDH was used as an internal control.

15 D. Immunohistochemistry stain of NF- κ B

The skin tissues were sectioned into 6 μ m thickness using the procedure set forth in the section of "3. Tissue slice" under "General Methods". The tissue sections were subjected to immunohistochemistry stain using primary antibody rabbit anti NF- κ B monoclonal antibody and donkey anti rabbit IgG
20 antibody conjugated with biotin (DarkoCytomation). The stained tissues were examined under an optical microscope (Olympus BX51, Japan) under 100 to 400X magnification, and was photographed using a digital camera (Olympus DP50, Japan).

E. Histochemistry stain of mast cells

25 The skin tissues were sectioned into 6 μ m thickness using the procedure set forth in the section of "3. Tissue slice" under "General Methods". The tissue sections were subjected to histochemistry stain with toluidine blue

(Sigma, Cat No. T3260). The stained tissue sections were examined under an optical microscope with 100 to 400X magnification, and were photographed with a digital camera (Olympus DP50, Japan). Quantification of mast cell number was conducted by randomly selecting 5 tissue sections, and counting randomly 10 observing fields in each of the tissue sections. Data are shown as mean $\pm$ SEM ($p < 0.05$ is statistically significant).

A. Diagnosis of clinical signs

HRS/J hairless mice in pathological control group showed symptoms of inflammation as compared to the normal control group, suggesting UV-B radiation successfully induced inflammatory response in HRS/J hairless mice. Experimental groups 1 and 2 showed a significant reduction in inflammatory response when compared to the pathological control group. These data suggest that *Solanum incanum* L. solution and *Solanum incanum* L. gel have anti-inflammatory effects.

15 B. Analysis of transepidermal water loss (TEWL)

Figure 2 is a graph showing the results of TEWL of the dorsal skins of the HRS/J hairless mice in normal control group, pathological group and experimental groups 1 and 2. The HRS/J hairless mice from pathological group showed an increased level of TEWL as compared to the normal control group. This indicates that UV-B radiation induced TEWL in HRS/J hairless mice and damaged the barrier function of stratum corneum. HRS/J hairless mice from experimental groups 1 and 2 showed a marked decrease in TEWL when compared to pathological control group. In all, these data suggest that *Solanum incanum* L. solution and *Solanum incanum* L. gel can improve stratum corneum damage and inflammation caused by UV-B radiation.

C. Western blot analysis of NF- κ B, COX-2 and iNOS

Figure 3 is a western blot analysis of NF- κ B, COX-2 and iNOS of dorsal

skin tissue from HRS/J hairless mice in normal control group, pathological group and experimental group 2. The skin tissues from the pathological control group showed an increased expression of NF- κ B, COX-2 and iNOS as compared to normal control group, suggesting that UV-B radiation induces
5 inflammatory response in HRS/J hairless mice. A significant reduction of NF- κ B, COX-2 and iNOS levels were observed in experimental group 2 when compared to pathological control group. In all, these data suggest that *Solanum incanum* L. gel have anti-inflammatory effects.

D. Immunohistochemistry stain of NF- κ B

10 Figure 4 shows an immunohistochemistry stain of NF- κ B expression on dorsal skin tissue obtained from HRS/J hairless mice. As shown in Figure 4, pathological control group showed an increased expression level in NF- κ B as compared to the normal control group, suggesting UV-B radiation induces inflammatory response in the HRS/J hairless mice. When *Solanum incanum*
15 L. gel was applied to HRS/J hairless mice, as in experimental group 2, the expression of NF- κ B showed a significant decrease when compared to the pathological control group. In all, these data suggest that *Solanum incanum* L. gel of the present invention has anti-inflammatory effect.

E. Histochemistry stain of mast cells

20 Figure 5 shows a histochemistry stain of mast cells obtained from the dorsal skin tissue of HRS/J hairless mice. As shown in Fig. 5, toluidine blue staining revealed an extensive mast cell infiltration in dorsal skin tissue from the pathological control group. This phenomenon was not visibly clear in the normal control group. Apparently, mast cell infiltration was significantly
25 reduced in experimental group 2 as compared to the pathological control group.

A comparison of mast cell number in each of normal control group,

pathological group and experimental group 2 is shown in Figure 6. Experimental group 2 showed a significant decrease of mast cell number when compared to pathological control group. These data suggest that *Solanum incanum L.* gel can improve the inflammatory effect induced by UV-B radiation in HRS/J hairless mice.

Example 3: Evaluation of the therapeutic effects of the *Solanum incanum L.* solution and the *Solanum incanum L.* gel on hairless mice with UV-B-induced photodamage.

HRS/J hairless mice was randomly grouped into normal control group, pathological control group and two experimental groups (i.e., experimental groups 1 and 2) (n=4/group). HRS/J hairless mice in normal control group did not receive any UV-B radiation treatment. Inflammation was induced in mice from pathological control group and two experimental groups as described in the section of "2. Induction of inflammation and photodamage" under "General Methods".

Starting from the initiation of UV-B radiation, HRS/J mice in experimental group 1 was administered orally with *Solanum incanum L.* solution, in which the solution was prepared as described in the section of "1. Preparations of water-soluble extract from *Solanum incanum L.* and *Solanum incanum L.* solution containing the water-soluble extract" under "Experimental Materials". These mice were administered with *Solanum incanum L.* solution once a day (1500 mg/kg), 7 times a week, during a 14-week period, and was continuously observed for 60 wks (i.e., 60 wk experimental period). In experimental group 2, the *Solanum incanum L.* gel prepared according to the section of "2. Preparation of *Solanum incanum L.* gel containing the water-soluble extract" under "Experimental Materials" was applied to dorsal skin of the mice subjected to UV-B radiation. The gel was applied once a day

(dosage: 10-20 mg of gel per cm²), 5 times a week, during a 14-wk period, and was observed for 60 wks. HRS/J hairless mice in normal control and pathological control group were not treated with the *Solanum incanum* L. solution or the *Solanum incanum* L. gel.

5 At the end of 26, 35, 38, 41, 52, 56, and 60 wks during the 60 wk experimental period, the dorsal skin from HRS/J hairless mice in pathological control and experimental groups 1 and 2 were subjected to analyses in Item A (see below). In addition, after the 60 wk experimental period, the dorsal skin obtained from HRS/J hairless mice in normal control, pathological control
10 and experimental group 2 were subjected to analyses in Item B as described below. The dorsal skin tissues of the mice from pathological control and experimental group 2 were obtained using surgical scissors and were subjected to analyses in Item C as described below.

A. Analysis of skin tumor formation

15 Tumor number on the dorsal skin of HRS/J hairless mice from pathological control and experimental groups 1 and 2 were observed by naked eyes. Tumor size was recorded using digital caliper (Mitutoyo, NTD15P-6"CX). Data for tumor number and volume are shown as mean $\pm$ SEM ($p < 0.05$ is statistically significant).

20 B. Evaluation of skin elasticity

 Skin elasticity was evaluated by referencing pinch testing that is mentioned in K. Tsukahara et al (2005), *Biolo. Pharm. Bull.*, 28 (12):2302-2307. In brief, HRS/J hairless mice from normal control, pathological control and experimental group 2 were placed on a platform.
25 The dorsal skin at the middle line of mouse body was pinched upward as much as possible (without lifting the mouse) and then released. The recovery time for the pinched skin to rebound to the original statue was measured.

C. Histochemical staining of elastic fiber

After the 60 wk experimental period, the skin tissues obtained from HRS/J hairless mice in pathological control and experimental group 2 were sectioned into 6µm thickness using the procedure set forth in the section of "3. Tissue slice" under "General Methods". Tissue sections were stained with resorcin-fuchsin solution, Weigert's iron hematoxylin and van Gieson's solution (all purchased from Muto Pure Chemicals Co., Ltd). Stained tissues were observed under an optical microscope with a 100-400X magnification and recorded with a digital camera (Olympus DP50, Japan).

10 A. Analysis of skin tumor formation

Tumor number over the course of the 60 wk experimental period is shown in Figure 7. Tumor size measurement over the course of the 60 wk experimental period is shown in Figure 8.

As shown in Figure 7 and 8, at 38 wk, the dorsal skin of the HSR/J hairless mice from the pathological control group showed an acceleration of tumor development. Conversely, a slower increase in tumor number and size was observed in experimental groups 1 and 2 during the 60 wk experimental period. These data suggest that *Solanum incanum* L. solution and *Solanum incanum* L. gel can improve the photodamage caused by UV-B radiation and alleviate skin pathologic change caused by photodamage.

B. Evaluation of skin elasticity

Pinch testing was done after the 60 wk experimental period on HRS/J hairless mice from normal control, pathological control and experimental group 2. As shown in Figure 9, pathological control group showed a decreased elasticity compared to the normal control group since a longer recovery time is required in the pathological control group; suggesting UV-B exposure induces photodamage in HRS/J hairless mice. When *Solanum*

incanum L. gel was applied to dorsal skin of hairless mice in experimental group 2, skin elasticity was greatly increased when compared to the pathological control group. Data from these experiments suggest that *Solanum incanum L.* gel can alleviate photodamage caused by UV-B and improve skin elasticity.

C. Histochemical staining of elastic fiber

As shown in Figure 10, the occurrence of elastic fiber denaturation was markedly increased in pathological control group. On the other hand, when *Solanum incanum L.* gel was applied to the dorsal skin of hairless mice in experimental group 2, there was less occurrence of elastic fiber denaturation and the fibers remained long and intact. Data from these experiments suggest that *Solanum incanum L.* gel of the present invention can alleviate photodamage caused by UV-B exposure and improve solar elastosis.

In all, data from the above experiments support that the water-soluble extract from *Solanum incanum L.* can prevent photodamage and improve pathological changes on the skin caused by photodamage.

Example 4: Comparison of the effect of the *Solanum incanum L.* gel and tretinoin on preventing UV-B-induced photodamage.

HRS/J hairless mice was randomly grouped into pathological control group, tretinoin group and *Solanum incanum L.* gel group (n=5/group). Induction of photodamage on HRS/J hairless mice from each group was performed according to the section of "2. Induction of inflammation and photodamage" under "General Methods".

Starting from the initiation of UV-B radiation, HRS/J mice in tretinoin group was applied with AIROL vanishing cream (contains 0.05% tretinoin, a conventional active component for treating photodamage, topical dosage: 10 to 20 mg of AIROL vanishing cream per cm²). The cream was applied once a

day, 5 times a week, for 14 weeks, and was continuously observed for 35 weeks (i.e., 35 week experimental period). The dorsal skin of HRS/J hairless mice in *Solanum incanum* L. gel group was applied with the *Solanum incanum* L. gel obtained from the section of "2. Preparation of *Solanum incanum* L. gel containing the water-soluble extract" under "Experimental Materials". The gel was applied once a day (dosage: 10-20 mg of gel /cm²), 5 times a week, during a 14-week period and was observed for 35 weeks. HRS/J hairless mice in pathological control group were not treated with the water-soluble extract.

10 Tumor number on the dorsal skin of the mice from tretinoin group and *Solanum incanum* L. gel group was quantified at 0 week (i.e., before exposure to UV-B radiation), and at the end of 2, 7, 12, 17, 22, 27, and 32 weeks during the 35 week experimental period based on the method mentioned in Item A (Analysis of skin tumor formation) of Example 3. In
15 addition, HRS/J hairless mice from pathological control, tretinoin group, and *Solanum incanum* L. gel group were subjected to analyses as mentioned in Item B (Analysis of transepidermal water loss) of Example 2 and the following Item A at 0 week and the end of 5, 10, 15, 20, 25, and 30 weeks during the 35 week experimental period.

20 A. Examination of water content in skin

Determination of water content in skin was examined by using an evaporimeter (Tewameter TM 201R, Courage & Khazaka, Cologne, Germany).

(1) Analysis of skin tumor formation

25 As shown in Fig. 11, at week 17, tretinoin group showed a dramatic increase in average tumor number. Conversely, HRS/J hairless mice from *Solanum incanum* L. gel group showed a slow increase in average tumor

number. These data suggest that *Solanum incanum* L. gel of the present invention can more effectively improve photodamage caused by UV-B when compared to tretinoin group, and alleviates pathological changes of the skin.

(2) Evaluation of transdermal water loss (TEWL)

5 As shown in Figure 12, during the 35 week experimental period, TEWL rate was lower in *Solanum incanum* L. gel group when compared to pathological and tretinoin groups. Specifically, at the end of 20, 25 and 30 weeks during the 35 week experimental period, there was a statistically significant difference in TEWL between the pathological control group and
10 *Solanum incanum* L. gel group ($p < 0.05$). These data suggest that *Solanum incanum* L. gel of the present invention can effectively alleviate photodamage induced by UV-B radiation and improve damage on stratum corneum caused by photodamage.

(3) Evaluation of water content in skin

15 Water content in the dorsal skin of HRS/J hairless mice from each group (pathological control, tretinoin and *Solanum incanum* L. gel groups) was determined at various time points throughout the 35 week experimental period. As shown in Fig. 13, the mice treated with *Solanum incanum* L. gel had a higher skin water content when compared to pathological control group
20 and tretinoin group. Specifically, at the end of 25 weeks, the mice treated with *Solanum incanum* L. gel showed a statistically significant water content as compared to the pathological group or tretinoin group. These data suggest that *Solanum incanum* L. gel of this invention is more effective in alleviating photodamage caused by UV-B and improve skin water loss than tretinoin
25 treatment.

In all, the water-soluble extract of this invention has been proven to exhibit prevention of photodamage, improvement of pathological changes

caused by photodamage, and maintenance of moisture-retaining capacity more effectively than what is often clinically used to treat and prevent photodamage (such as tretinoin).

CLAIMS:

1. Use of a water-soluble extract from a plant of *Solanum* genus in the manufacture of a medicament for preventing and treating cutaneous photodamage, wherein said water-soluble extract comprises at least
5 60% of solamargine and solasonine.
2. The use of claim 1, wherein said water-soluble extract comprises at least 60%-90% of solamargine and solasonine.
3. The use of claim 1 or claim 2, wherein said cutaneous photodamage comprises one of the following symptoms: thinning of the
10 skin, skin atrophy, decrease in collagen fiber and elastic fiber, elastosis, loss of skin elasticity, dryness, wrinkle formation, inflammatory cells infiltration, premature skin aging, vascular change, pigmentary change, comedone, cyst and skin pathologic change.
4. The use according to any one of claims 1 to 3, wherein the
15 water-soluble extract is administered by an oral route.
5. The use according to any one of claims 1 to 3, wherein the water-soluble extract is administered by a topical route.
6. Use of a water-soluble extract from a plant of *Solanum* genus in the manufacture of a medicament for preventing and treating
20 inflammation, wherein said water-soluble extract comprises at least 60% of solamargine and solasonine.
7. The use of claim 6, wherein said water-soluble extract comprises at least 60%-90% of solamargine and solasonine.
8. The use according to claim 6 or claim 7, wherein the
25 water-soluble extract is administered by an oral route.
9. The use according to claim 6 or claim 7, wherein the water-soluble extract is administered by a topical route.