# GENE 16 WHITENING FACTOR- allantoin tablet, soluble Shanghai Liang Liang International Inc

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# Shanghai Liang Liang International Inc

# **Active Ingredient**

Allantoin 0.5%

# **Purpose**

Allantoin Skin Lightener

Skin Protectant

## Warning

For External Use Only.

#### Use

Apply moderate amount of the product on the skin.

# Stop Using the Product if

Stop using the product when you have skin problems or the product disagrees with your skin.

Stop using the product immediately and consult a dermatologist if you have redness, swelling, itching or irritation on the skin while or after using the product.

 $\square$ If the product gets into the eyes, don't rub but rinse with water.

# Keep Out of Reach of Children

Keep out of reach of children.

# When Using the Product

Mix one tablet with water in palm to dissolve before applying to skin. Two tablets two times a day. One after face cleanse in the morning; and one before bed.

#### Other Information

Keep in a dry, cool place away from direct sunlight. Store at room temperature.

### Contact the Manufacturer or Distributor

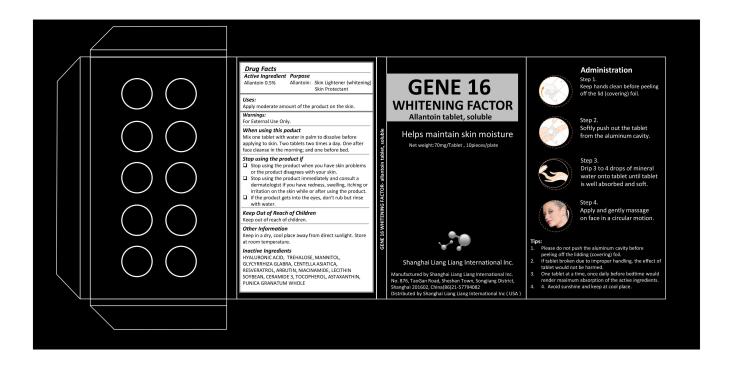
Manufactured by Shanghai Liang Liang International Inc. No. 876, TaoGan Road, Sheshan Town, Songjiang District, Shanghai 201602, China (86)21-57794082

Distributed by Shanghai Liang Liang International Inc (USA)

# **Inactive Ingredients**

HYALURONIC ACID, TREHALOSE, MANNITOL, GLYCYRRHIZA GLABRA, CENTELLA ASIATICA, RESVERATROL, ARBUTIN, NIACINAMIDE, LECITHIN SOYBEAN, CERAMIDE 3, TOCOPHEROL, ASTAXANTHIN, PUNICA GRANATUM WHOLE

# **Drug Facts**



# GENE 16 WHITENING FACTOR allantoin tablet, soluble Product Information Product Type HUMAN OTC DRUG Item Code (Source) Route of Administration TOPICAL

| Active Ingredient/Active Moiety                            |                          |                  |  |
|--|--------------------------|------------------|--|
| Ingredient Name  | <b>Basis of Strength</b> | Strength         |  |
| ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z) | ALLANTOIN                | 0.5 mg in 100 mg |  |

| Inactive Ingredients                     |          |  |
|--|----------|--|
| Ingredient Name                          | Strength |  |
| HYALURONIC ACID (UNII: S270N0TRQY)       |          |  |
| TREHALOSE (UNII: B8WCK70T7I)             |          |  |
| MANNITOL (UNII: 30WL53L36A)              |          |  |
| GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)    |          |  |
| CENTELLA ASIATICA (UNII: 7M867G6T1U)     |          |  |
| RESVERATROL (UNII: Q369O8926L)           |          |  |
| ARBUTIN (UNII: C5INA23HXF)               |          |  |
| NIACINAMIDE (UNII: 25X51I8RD4)           |          |  |
| LECITHIN, SOYBEAN (UNII: 1DI56QDM62)     |          |  |
| CERAMIDE 3 (UNII: 4370DF050B)            |          |  |
| TOCOPHEROL (UNII: R0ZB2556P8)            |          |  |
| ASTAXANTHIN (UNII: 8XPW32PR7I)           |          |  |
| PUNICA GRANATUM WHOLE (UNII: O2ZTS50U5E) |          |  |

| Product Characteristics |       |              |          |  |
|-------------------------|-------|--------------|----------|--|
| Color                   | white | Score        | no score |  |
| Shape                   | OVAL  | Size         | 8mm      |  |
| Flavor                  |       | Imprint Code |          |  |
| Contains                |       |              |          |  |

| Packaging |                      |  |                         |                       |
|-----------|----------------------|--|-------------------------|-----------------------|
| #         | Item Code            | Package Description                                | Marketing Start<br>Date | Marketing End<br>Date |
| 1         | NDC:69764-001-<br>10 | 1 in 1 PACKAGE                                     | 09/25/2014              |                       |
| 1         |                      | 7 mg in 1 POUCH; Type 0: Not a Combination Product |                         |                       |

| Marketing Information                                       |      |                         |                       |  |
|---|------|-------------------------|-----------------------|--|
| Marketing Application Number or Monograph Category Citation |      | Marketing Start<br>Date | Marketing End<br>Date |  |
| OTC Monograph Drug  | M016 | 09/25/2014              |                       |  |
|   |      |                         |                       |  |

# Labeler - Shanghai Liang Liang International Inc (547834726)

| Establishment                          |         |           |                            |  |
|--|---------|-----------|----------------------------|--|
| Name                                   | Address | ID/FEI    | <b>Business Operations</b> |  |
| Shanghai Liang Liang International Inc |         | 547834726 | manufacture(69764-001)     |  |

Revised: 12/2023 Shanghai Liang Liang International Inc