

**SKINCEUTICALS ADVANCED SCAR CONTROL SKIN PROTECTANT- allantoin gel**  
**L'Oreal USA Products Inc**

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**Drug Facts**

**Active ingredient**

Allantoin 0.5%

**Purpose**

Skin protectant

**Uses**

temporarily protects minor:

- cuts
- scrapes
- burns

**Warnings**

For external use only

**Do not use on**

- deep or puncture wounds
- animal bites
- serious burns

**When using this product**

do not get into eyes

**Stop use and ask a doctor if**

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- apply as needed

### **Inactive ingredients**

dimethicone, dimethicone crosspolymer, helianthus annuus (sunflower) seed oil unsaponifiables, silica silylate, acrylates/dimethicone copolymer, jojoba esters, helianthus annuus (sunflower) seed wax, acacia decurrens flower wax, polyglycerin-3

### **Questions or comments?**

Call 1-800-811-1660

Monday - Friday (9 a.m. - 5 p.m. CST)

CORRECT

95% SILICONES

SKINCEUTICALS

ADVANCED  
SCAR CONTROL

ALLANTOIN  
SKIN PROTECTANT

IMPROVES APPEARANCE,  
COLOR, AND TEXTURE OF  
NEW AND RECENT SCARS

50 ml / 1.7 fl oz

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855262 5 (CODE F.I.L.: D251660/1)



SKINCEUTICALS®

SkinCeuticals LLC, Dallas, TX 75241

Made in USA of US and/or Imported

Ingredients [www.skinceuticals.com](http://www.skinceuticals.com)

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allantoin gel

**Product Information**

|   |   |  |                             |                           |
|---|---|--|-----------------------------|---------------------------|
| <b>Product Type</b>   | HUMAN OTC DRUG                                  | <b>Item Code (Source)</b>                            | NDC:49967-783               |                           |
| <b>Route of Administration</b>  | TOPICAL   |  |                             |                           |
| <b>Active Ingredient/Active Moiety</b>  |   |  |                             |                           |
| <b>Ingredient Name</b>  |   | <b>Basis of Strength</b>                             | <b>Strength</b>             |                           |
| ALLANTOIN (UNII: 344S277G0Z) (ALLANTOIN - UNII:344S277G0Z)                              |   | ALLANTOIN  | 5 mg in 1 mL                |                           |
| <b>Inactive Ingredients</b>   |   |  |                             |                           |
| <b>Ingredient Name</b>  |   |  |                             | <b>Strength</b>           |
| DIMETHICONE (UNII: 92RU3N3Y1O)  |   |  |                             |                           |
| DIMETHICONE CROSSPOLYMER (450000 MPA.S AT 12% IN CYCLOPENTASILOXANE) (UNII: UF7620L1W6) |   |  |                             |                           |
| SUNFLOWER OIL (UNII: 3W1JG795YI)  |   |  |                             |                           |
| HYDROGENATED JOJOBA OIL, RANDOMIZED (UNII: Q47ST02F58)                                  |   |  |                             |                           |
| HELIANTHUS ANNUUS SEED WAX (UNII: 42DG15CHXV)   |   |  |                             |                           |
| ACACIA DECURRENS FLOWER WAX (UNII: AU6XZ E9IY9)   |   |  |                             |                           |
| POLYGLYCERIN-3 (UNII: 4A0NCJ6RD6)   |   |  |                             |                           |
| <b>Packaging</b>  |   |  |                             |                           |
| <b>#</b>  | <b>Item Code</b>                                | <b>Package Description</b>                           | <b>Marketing Start Date</b> | <b>Marketing End Date</b> |
| 1   | NDC:49967-783-01                                | 1 in 1 CARTON  | 06/01/2021                  |                           |
| 1   |   | 50 mL in 1 TUBE; Type 0: Not a Combination Product   |                             |                           |
| 2   | NDC:49967-783-02                                | 4 mL in 1 TUBE; Type 0: Not a Combination Product    | 06/01/2021                  |                           |
| 3   | NDC:49967-783-03                                | 1 in 1 CARTON  | 06/01/2021                  |                           |
| 3   |   | 50 mL in 1 BOTTLE; Type 0: Not a Combination Product |                             |                           |
| <b>Marketing Information</b>  |   |  |                             |                           |
| <b>Marketing Category</b>   | <b>Application Number or Monograph Citation</b> | <b>Marketing Start Date</b>                          | <b>Marketing End Date</b>   |                           |
| OTC Monograph Drug  | M016  | 06/01/2021   |                             |                           |

**Labeler** - L'Oreal USA Products Inc (002136794)

### Establishment

| Name                                 | Address | ID/FEI    | Business Operations    |
|--------------------------------------|---------|-----------|------------------------|
| Beauty Manufacturing Solutions Corp. |         | 783200723 | manufacture(49967-783) |