

**LANTISEPTIC BY DERMARITE DRY SKIN THERAPY- lanolin cream**  
**Dermarite Industries LLC**

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**Lantiseptic by Dermarite Dry Skin Therapy**

***Active Ingredient***

Lanolin USP 30%

***Purpose***

Skin Protectant

***Uses***

Temporarily protects minor cuts, scrapes, and burns.

Helps prevent and temporarily protects chafed, chapped or cracked skin.

***Warnings***

- **For external use only.**
- **Avoid contact with eyes.** In case of contact, flush thoroughly with water
  - **Do not use on**
  - Deep or puncture wounds
  - animal bites
  - serious burns

**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.** In case of accidental ingestion contact a physician or Poison Control Center right away.

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***Directions***

Apply liberally to affected area as needed or as directed by a physician.

***Other Information***

Store at room temperature (59-86°F)

You may report a serious adverse event to DermaRite Industries, PO Box 7209, North Bergen, NJ 07047.

***Inactive Ingredients***

Water, Mineral Oil, Petrolatum, Cera Alba, Sodium Borate, DMDM Hydantoin, Sorbitan Sesquioleate, Lanolin Alcohol, Disodium EDTA.

**Questions or Comments?**

Call 1-800-337-6296 Mon-Fri 9AM-5PM EST

**Package Labeling:**

NDC 61924-504-04



Dry Skin Therapy

SKIN PROTECTANT

LANOLIN ENRICHED™

30% Lanolin  
Softens and moisturizes  
damage prone skin

REORDER #LS0410

Net Wt. 113 g (4 oz.)

**Drug Facts**

Active ingredient	Purpose
Lanolin USP 30%	Skin protectant

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Patient Name

Room #



DermaRite®

DermaRite Industries LLC • 7777 West Side Avenue  
North Bergen, NJ 07047 • www.dermarite.com

MADE IN THE USA  
100238

**LANTISEPTIC BY DERMARITE DRY SKIN THERAPY**

lanolin cream

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:61924-504
<b>Route of Administration</b>	TOPICAL		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
LANOLIN (UNII: 7EV65EAW6H) (LANOLIN - UNII:7EV65EAW6H)	LANOLIN	300 mg in 1 g

**Inactive Ingredients**

Ingredient Name	Strength
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	
WATER (UNII: 059QF0KO0R)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:61924-504-14	14.2 g in 1 PACKET; Type 0: Not a Combination Product	12/25/2018	
2	NDC:61924-504-04	113 g in 1 TUBE; Type 0: Not a Combination Product	12/25/2018	
3	NDC:61924-504-05	5 g in 1 PACKET; Type 0: Not a Combination Product	12/25/2018	

**Marketing Information**

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/25/2018	

**Labeler** - Dermarite Industries LLC (883925562)**Registrant** - Dermarite Industries LLC (883925562)