

LANOLIN 50%- lanolin cream
Aldermed Inc.

Lanolin 50%

Active Ingredient

Lanolin USP 50%

Purpose

Skin Protectant

Uses

- temporarily protects minor: cuts, scrapes, burns
- temporarily protects and helps relieve chafed, chapped or cracked skin and lips
- helps prevent and protect from the drying effects of wind and cold weather

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

If pregnant or breast-feeding

ask a health professional before use.

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

Directions

apply as needed

Other Information

- to open: unscrew cap, pull tab to remove foil seal
- store at 15° to 30°C (59° to 86°F)
- avoid excessive heat
- see tube crimp for lot number and expiration date

Inactive Ingredients

cetosteryl alcohol, edetate disodium, fragrance, lanolin alcohols, polysorbate 60, purified water, sodium borate, white petrolatum, white wax

Questions

Call 1-833-605-84-74 9am-5 pm EST M-F

Label



LANOLIN 50%

lanolin cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:87236-019
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LANOLIN (UNII: 7EV65EAW6H) (LANOLIN - UNII: 7EV65EAW6H)	LANOLIN	50 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
POLYSORBATE 60 (UNII: CAL22UVI4M)	
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
WHITE WAX (UNII: 7G1J5DA97F)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87236-019-04	113 g in 1 TUBE; Type 0: Not a Combination Product	12/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/01/2025	

Labeler - Aldermed Inc. (144887712)

Revised: 5/2026

Aldermed Inc.