LANA1263- lanolin ointment Blossom Pharmaceuticals

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Lanashield Ointment

ACTIVE INGREDIENT

Active ingredient Purpose

Lanolin USP 50% Skin Protectant

Purpose:

- Helps prevent and treat diaper dermatitis
- Protects chafed skin or minor skin irritations due to incontinence and helps seal out wetness

Warnings

For External Use Only

Keep Out Of Reach Of Children

KEEP OUT OF REACH OF CHILDREN

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

Indications & Usage

- Avoid contact with eyes
- Do not apply to deep or puncture wounds
- If condition worsens, or does not improve within 7 days, consult a doctor

Dosage & Administration

- Gently cleanse and dry area
- Apply liberally to affected area as needed

Other information:

• Store at room temperature 20 deg C to 25 deg C 68 deg F to 77 deg F

Inactive Ingredients

Beeswax (White), EDTA, Emulsifying wax, Lanolin alcohol, Lavender perfume, Mineral oil, Petrolatum, Purified water, Sodium borate

Principal Display Panel

Lana1263 Ointment

Lana1263.jpg



LANA1263

lanolin ointment

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:61767-232

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			
WHITE WAX (UNII: 7G1J5DA97F)				
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)				
MINERAL O IL (UNII: T5L8T28FGP)				
SODIUM BORATE (UNII: 91MBZ8H3QO)				
WATER (UNII: 059QF0KO0R)				
EDETIC ACID (UNII: 9G34HU7RV0)				
PETROLATUM (UNII: 4T6H12BN9U)				

Packaging					
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:61767-232-01	5 g in 1 PACKET			
2	NDC:61767-232-02	15 g in 1 PACKET			
3	NDC:61767-232-03	71 g in 1 JAR			
4	NDC:61767-232-04	113 g in 1 TUBE			
5	NDC:61767-232-05	127.5 g in 1 JAR			
6	NDC:61767-232-06	396.8 g in 1 JAR			

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph final	part347	10/23/2014			

Labeler - Blossom Pharmaceuticals (677381470)

Revised: 10/2014 Blossom Pharmaceuticals