LANOLIN - lanolin ointment
Dynarex Corporation

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

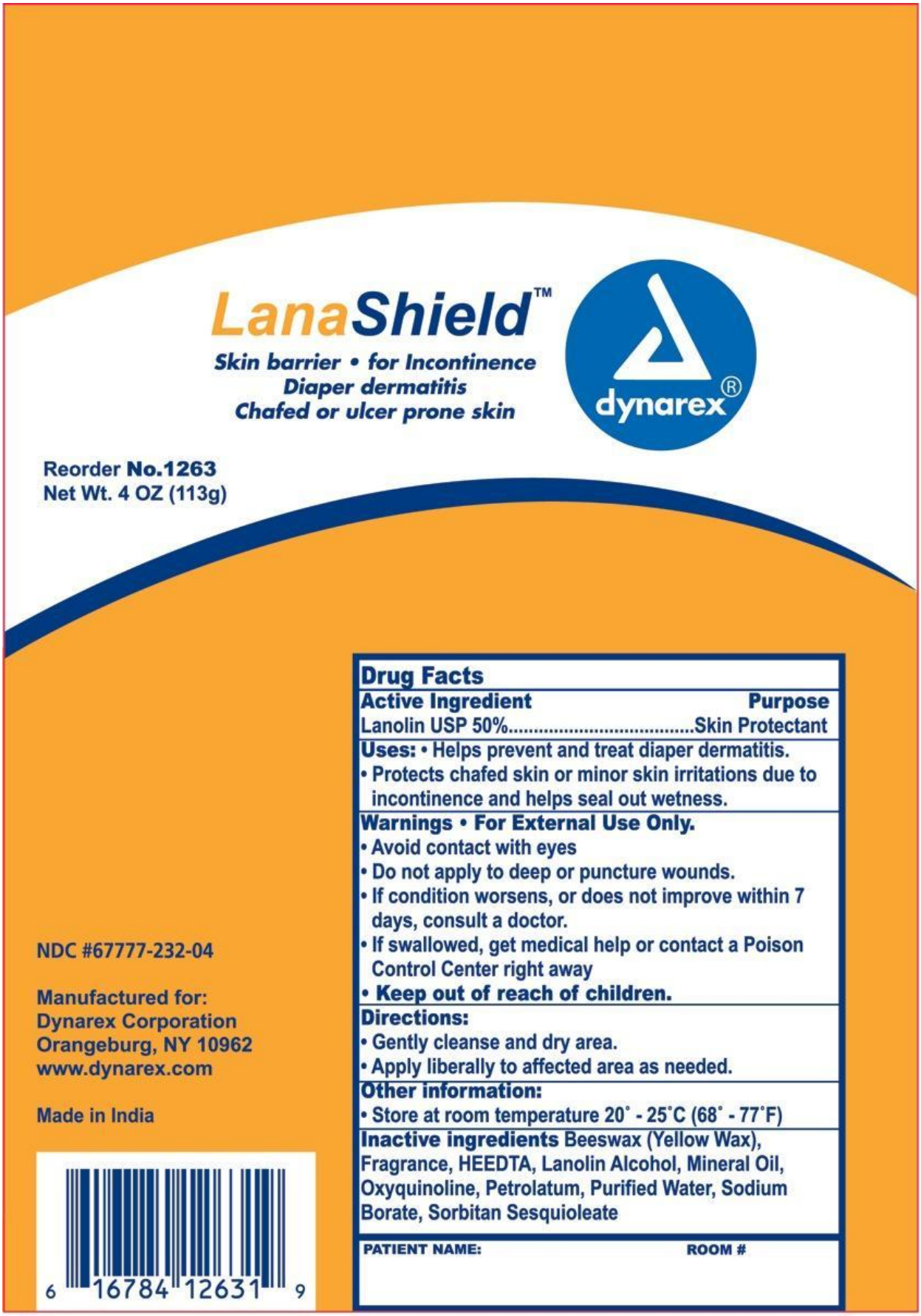
Dosage & Administration
- Gently cleanse and dry area
- Apply liberally to affected area as needed

Other information:
- Store at room temperature 20 deg C to 25 deg C 68 deg F to 77 deg F

Inactive Ingredients

Beeswax (Yellow Wax), Fragrance, HEEDTA, Lanolin alcohol, Mineral oil, Oxyguinoline, Petrolatum, Purified water, Sodium borate, Sorbitan sesquioleate

Principal Display Panel



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LANOLIN

lanolin ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-232
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**
SORBITAN SESQUIOLEATE (UNII: 0W8RRI5W5A)	
PETROLATUM (UNII: 4T6H12BN9U)	
YELLOW WAX (UNII: 2ZA36H0S2V)	
LANOLIN ALCOHOLS (UNII: 884C3FA9HE)	
MINERAL OIL (UNII: T5L8T28FGP)	
OXYQUINOLINE (UNII: 5UTX5635HP)	
SODIUM BORATE (UNII: 91MBZ8H3QO)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:67777-232-01	2 in 1 CASE		
1		144 in 1 BOX		
1		5 g in 1 PACKET		
2	NDC:67777-232-02	4 in 1 CASE		
2		36 in 1 BOX		
2		15 g in 1 PACKET		
3	NDC:67777-232-03	36 in 1 CASE		
3		71 g in 1 JAR		
4	NDC:67777-232-04	24 in 1 CASE		
4		113 g in 1 TUBE		
5	NDC:67777-232-05	24 in 1 CASE		
5		127.5 g in 1 JAR		
6	NDC:67777-232-06	12 in 1 CASE		
6		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	05/12/2010	

Labeler - Dynarex Corporation (008124539)

Registrant - Dynarex Corporation (008124539)