ZINC OXIDE- zinc oxide ointment ointment
Dynarex Corporation

1190, 1191, 1192 Zinc Oxide Ointment 20%

Active Ingredient

Zinc Oxide 20%

Purpose

Skin Protectant

Use(s)

• Helps treat chafed skin associated with diaper rash

• Dries the oozing and weeping of poison ivy, oak, and sumac

Warnings

For External Use Only

When using this product

• Avoid contact with 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 (1-800-222-1222) right away.

Directions

• For diaper rash: Change wet and soiled diapers promptly, cleanse the diaper area, and allow to dry. Apply ointment liberally as often as necessary with each diaper change, especially at bedtime or anytime when exposure to wet diapers may be prolonged.

• For poison ivy, oak, and sumac: Apply as needed.

Other Information

• Store at room temperature between 15º-30ºC (59º-86ºF)

- Avoid excessive heat
- Tamper evident. Do not use if seal is damaged.

Inactive Ingredients

Cetomacrogol 1000. Cetostearyl Alcohol Light Liquid Paraffin, White Soft Paraffin

Questions?

1-888-Dynarex Monday - Friday, 9AM - 5PM EST.

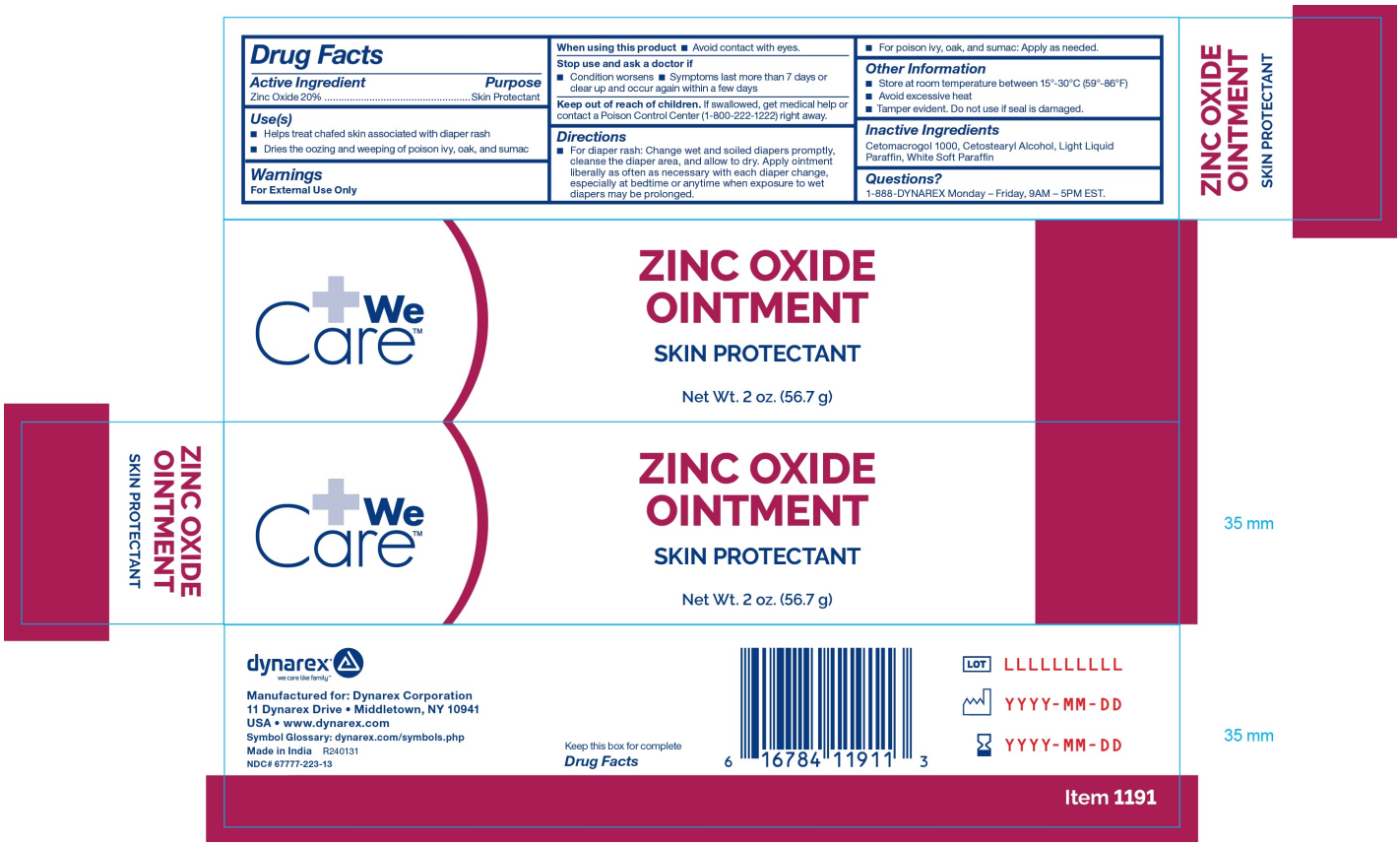
Label

Label 1192

Label

Label 1190

Label



Label 1191

ZINC OXIDE

zinc oxide ointment ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-223
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ZINC OXIDE (UNII: SOI2LOH54Z) (ZINC OXIDE - UNII:SOI2LOH54Z)	ZINC OXIDE	200 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-20 (UNII: I835H2IHHX)	

Product Characteristics			
Color		**Score**	
Shape	FREEFORM	**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:67777-223-02	72 in 1 CASE	02/29/2024	
1	NDC:67777-223-11	1 in 1 BOX		
1	NDC:67777-223-01	28.4 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:67777-223-04	72 in 1 CASE	02/29/2024	
2	NDC:67777-223-13	1 in 1 BOX		
2	NDC:67777-223-03	56.7 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:67777-223-06	12 in 1 CASE	02/29/2024	
3	NDC:67777-223-05	425 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	02/29/2024	

Labeler - Dynarex Corporation (008124539)

Revised: 4/2026 Dynarex Corporation