NEOMYCIN- neomycin ointment ointment
Dynarex Corporation

1164UB-10
1164-25

Active Ingredient (in each gram)

Neomycin Sulfate 5mg (equivalent to 3.5 mg Neomycin)

Purpose

First Aid Antibiotic

Use(s)

First aid to help prevent infection in minor cuts, scrapes, and burns

Warnings

For External Use Only

Do not use

- In the eyes or over large areas of the body
- If you are allergic to any of the ingredients
- Longer than 1 week unless directed by a doctor

Ask a doctor before use if

You have deep or puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if

- The condition persists or gets worse
- A rash or other allergic reaction develops

Keep out of reach of children

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

Directions

- Clean the affected area
- Apply a small amount of this product (an amount equal to the surface area of the tip

of a finger) on the area 1 to 3 times daily

• May be covered with a sterile bandage

Other Information

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

• Tamper evident. Do not use if seal is torn, cut, or opened.

Inactive Ingredient(s)

White Petrolatum

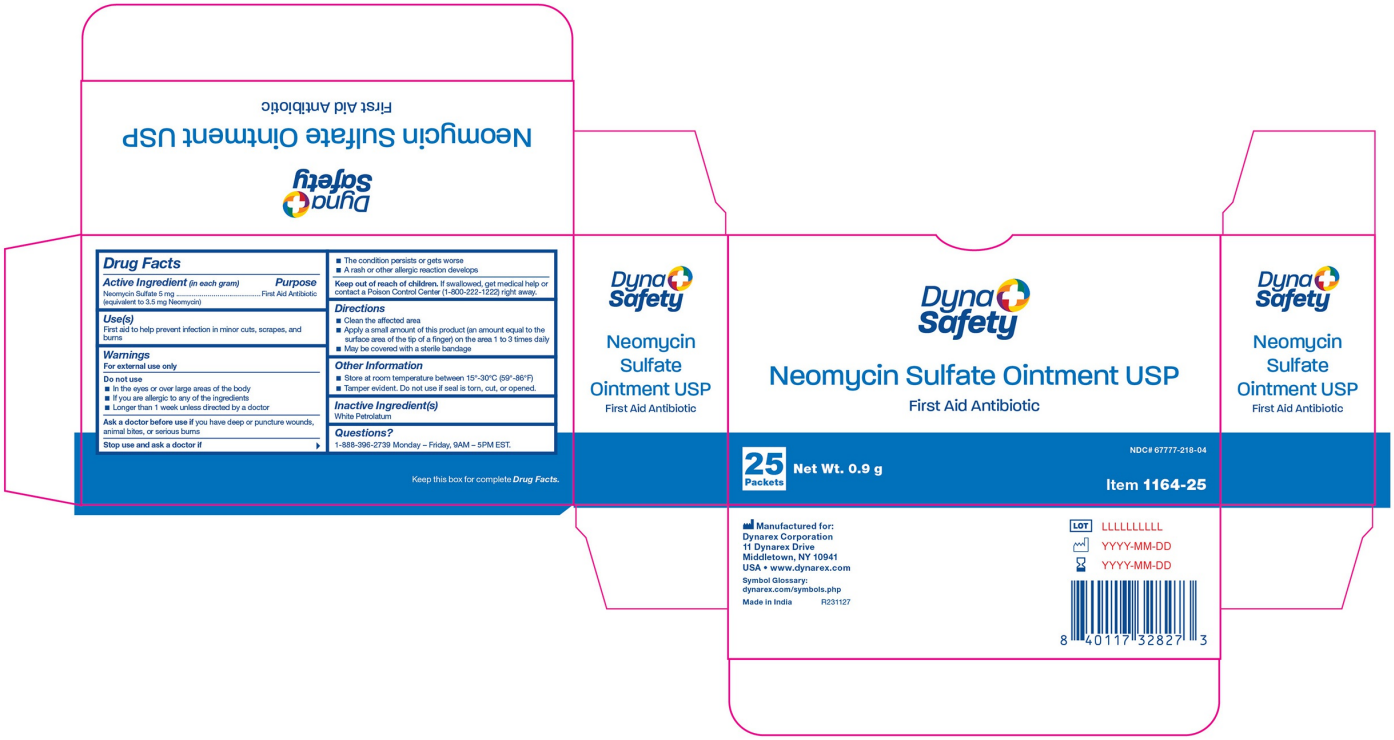
Questions?

1-888-396-2739 Monday - Friday, 9AM - 5PM EST.

Label

1164UB-10

Label



1164-25

NEOMYCIN

neomycin ointment ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-218
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-218-01	1000 in 1 CASE	01/30/2024	
1	NDC:67777-218-02	10 in 1 BOX		
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product		
2	NDC:67777-218-03	1800 in 1 CASE	01/30/2024	
2	NDC:67777-218-04	25 in 1 BOX		
2		0.9 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	01/30/2024	

Labeler - Dynarex Corporation (008124539)

Revised: 1/2024 Dynarex Corporation