

**FAMILY CARE TRIPLE ANTIBIOTIC PAIN RELIEF - neomycin sulfate ointment
UNITED EXCHANGE CORP.**

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.

Drug Facts

Active Ingredients (each gram contains)	Purpose
Neomycin Sulfate 3.5 mg.....	Antibiotic

Uses

first aid to help prevent infection in

- minor cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- in the eyes
- if you are allergic to any of the ingredients
- over large areas of the body
- longer than 1 week unless directed by a doctor

Ask a doctor before use in case of 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 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 15° to 25°C (59° to 77°F)
- Lot No. and Exp. Date: see box or see crimp of tube

Inactive ingredients

liquid paraffin, cetanol, squalane, white petrolatum, stearyl alcohol, tocopherol acetate, purified lanolin, methyl parahydroxybenzoate, propyl parahydroxybenzoate

Distributed by:

UNITED EXCHANGE CORP.

17211 Valley View Ave.
 Cerritos, CA 90703 USA
 Made in Korea



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FAMILY CARE TRIPLE ANTIBIOTIC PAIN RELIEF
 neomycin sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-007
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
SQUALANE (UNII: GW89575KF9)	
PETROLATUM (UNII: 4T6H12BN9U)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
LANOLIN (UNII: 7EV65EAW6H)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
PROPYL PARABEN (UNII: Z8IX2SC1OH)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-007-23	1 in 1 CARTON		
1		28 g in 1 TUBE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	02/07/2012	

Labeler - UNITED EXCHANGE CORP. (840130579)

Revised: 2/2012

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