FAMILY CARE ANTIBIOTIC- 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.

Family Care Antibiotic Ointment 513

Active ingredient (in each gram) Purpose

Neomycin sulfate 3.5 mg.....First Aid-Antibiotic

Warnings

For external use only

Uses

first aid to help prevent infection in:

- minor cuts
- scrapes
- burns

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

- you need to use longer than 1 week
- rash or other allergic reaction develops
- the condition persists or gets worse

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately.

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

- Istore between 20° to 25°C (68° to 77°F)
- Lot No. & Exp. Date: see box or see crimp of tube

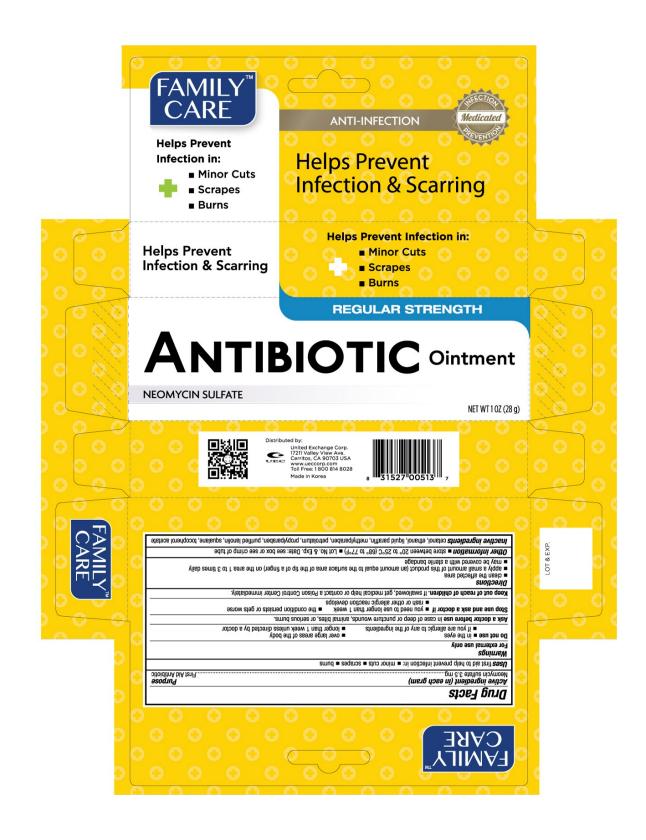
Inactive ingredients

cetanol, ethanol, liquid paraffin, methylparaben, petrolatum, propylparaben, purified lanolin, squalane, tocopherol acetate

Distributed by:

United Exchange Corp.

17211 Valley View Ave. Cerritos, CA 90703 USA www.ueccorp.com 1 800 814 8028 Made in Korea



FAMILY CARE ANTIBIOTIC

neomycin sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-517
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g	

Inactive Ingredients		
Ingredient Name	Strength	
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
PETROLATUM (UNII: 4T6H12BN9U)		
SQUALANE (UNII: GW89575KF9)		
METHYLPARABEN (UNII: A2I8 C7HI9 T)		
LANOLIN (UNII: 7EV65EAW6H)		
CETYL ALCOHOL (UNII: 936JST6JCN)		
.ALPHATO CO PHERO L ACETATE (UNII: 9E8X80D2L0)		

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-517-28	1 in 1 BOX	12/29/2017	
1		28 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333B	12/29/2017		

Labeler - United Exchange Corp. (840130579)

Revised: 12/2017 United Exchange Corp.