# 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.

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## **Drug Facts**

#### 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

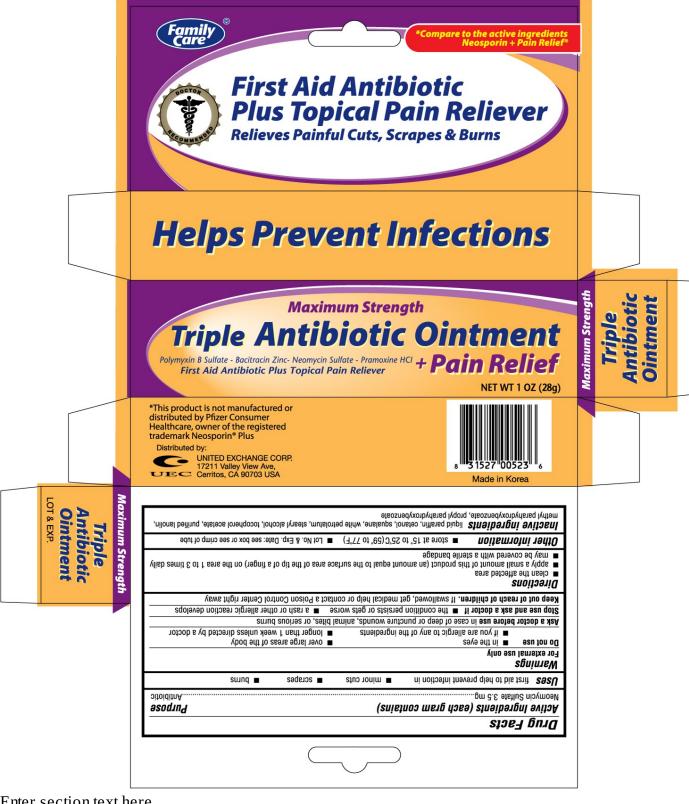
## Inactive ingredients

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

### Distributed by:

UNITED EXCHANGE CORP.

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:116QD7X297)	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: 2KR89I4H1Y)			
LANOLIN (UNII: 7EV65EAW6H)			
METHYLPARABEN (UNII: A218 C7HI9 T)			
PROPYLPARABEN (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 UNITED EXCHANGE CORP.