FAMILY CARE TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin neomycin sulfate polymyxin b sulfate pramoxine hydrochloride 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.

Active ingredients (in each gram)	Purpose

Bacitracin 500 units..... First aid antibiotic

Neomycin sulfate 3.5 mg..... First aid antibiotic

Polymyxin B sulfate 10,000 units...... First aid antibiotic

Pramoxine hydrochloride 10mg...... First aid antibiotic

Uses

first aid to help prevent infection and for temporarily relief of pain or discomfort in minor:

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- if you are allergic to any of the ingredients
- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- you need to use longer than 1 week
- symptoms persist for more than 1 week, or clear up and occur again within a few days
- the condition persists or gets worse
- rash or allerfic reaction develops

Keep out of reach of children

If swallowed, get medical help or contact Poison Control Center immediately

Directions

- adults and children 2 years of age and older:
- 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 stearile bandage
- children under 2 years of age: ask a doctor

Other information

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

Inactive ingredients

ethanol, liquid paraffin, methylparaben, polyoxyl 40 stearate, propylparaben, white petrolatum

Distributed by:

United Exchange Corp.

17211 Valley View Ave.

Cerritos CA, 90703 USA





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bacitracin neomycin sulfate polymyxin b sulfate pramoxine hydrochloride ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65923-523
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
BACITRACIN (UNII: 58 H6 RWO 52I) (BACITRACIN - UNII: 58 H6 RWO 52I)	BACITRACIN	500 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
PRAMO XINE HYDRO CHLO RIDE (UNII: 88 AYB867L5) (PRAMO XINE - UNII: 068 X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
PARAFFIN (UNII: 1900E3H2ZE)		
METHYLPARABEN (UNII: A2I8C7HI9T)		
POLYOXYL 40 STEARATE (UNII: 13A4J4NH9I)		
PROPYLPARABEN (UNII: Z8IX2SC1OH)		
PETROLATUM (UNII: 4T6H12BN9U)		

P	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65923-523-14	1 in 1 CARTON	09/15/2016	
1		14 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	11/24/2014	

Labeler - United Exchange Corp. (840130579)

Registrant - United Exchange Corp. (840130579)

Revised: 9/2016 United Exchange Corp.