LEADER TRIPLE ANTIBIOTIC PLUS- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, pramoxine hydrochloride ointment CARDINAL HEALTH

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.

Leader Triple Antibiotic Ointment Plus 1 oz. NBE Neosporin + Pain Relief

Uses

help prevent infection and temporarily relieves pain due to

- minor cuts
- scrapes
- burns

Warnings

For external use only

Allergy alert: do not use if allergic to any of the ingredients

Do not use

- in or near the eyes
- over large areas of the body
- longer than 1 week

Ask a doctor before use in case of deep or puncture wounds, animal bites, or serious burns

When using this product do not use longer than 1 week

Stop use and ask a doctor if

- condition persists or gets worse
- symptoms last for more than 7 days or clear up and comes back within a few days
- a rash or other allergic reaction occurs

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

Directions

Adults and children 2 years and older:

- clean affected area
- apply a small amount (equal to the surface area of the tip of a finger) on 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 crimp of tube

Inactive ingredients

white petrolatum, water

DISTRIBUTED BY CARDINAL HEALTH

DUBLIN, OHIO 43017

CIN 2372167

www.myleader.com

1-800-200-6313

Made in Korea



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Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:37205-266	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	BACITRACIN	500 [USP'U] in 1 g		
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	3.5 mg in 1 g		
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] 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	
PETROLATUM (UNII: 4T6H12BN9U)		
WATER (UNII: 059QF0KO0R)		

]	Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:37205-266-10	1 in 1 CARTON	0 3/0 1/20 12		
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	0 3/0 1/20 12		

Labeler - CARDINAL HEALTH (097537435)

Revised: 12/2017 CARDINAL HEALTH