FIRST AID ANTIBIOTIC BACITRAYCIN PLUS MAXIMUM STRENGTH- bacitracin and pramoxine hclointment First Aid Research 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.

First Aid Antibiotic Bacitraycin Plus Maximum Strength Ointment

Active ingredient (each gram contains)

Bacitracin 500 units

Pramoxine HCl 10 mg

Purpose

First Aid Antibiotic

Pain Reliever

Uses

first aid to help prevent infection in

- minor cuts
- scrapes
- burns

Warnings

For external use only.

Do not use

- in the eyes
- over large areas of the body
- if you are allergic to any of the ingredients
- 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. & Exp. Date: see box or see crimp of tube

Inactive ingredients

aloe vera leaf, methylparaben, mineral oil, petrolatum, propylparaben, stearyl alcohol

Distributed by:

First Aid Research Corp.

Jupiter, FL 33478

Made in India



FIRST AID ANTIBIOTIC BACITRAYCIN PLUS MAXIMUM STRENGTH

bacitracin and pramoxine hcl ointment

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:75983-004	
Route of Administration	TOPICAL				
Active Ingredient/Active Moi	A 4-7				
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Ingree	lient Name		Basis of Stren	gth	Strength
BACITRACIN (UNII: 58H6RWO52I) (BA	BACITRACIN		500 [USP'U] in 1 g		
PRAMO XINE HYDRO CHLO RIDE (UNII: 88AYB867L5) (PRAMO XINE - UNII:068X84E056)			PRAMOXINE HYDROCHLORIDE		10 mg in 1 g
Inactive Ingredients					
Ingredient Name					Strength
ALOE VERA LEAF (UNII: ZY81Z83H02	K)				
METHYLPARABEN (UNII: A2I8C7HI9T	")				
MINERAL OIL (UNII: T5L8T28FGP)					

PETROLATUM (UNII: 4T6H12BN9U)

STEARYL ALCOHOL	(UNII: 2KR8914H1Y)				
Dl					
Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
NDC:75983-004-28	1 in 1 CARTON	02/20/2018			
1	28 g in 1 TUBE; Type 0: Not a Combination Product				
Marketing Info	rmation				
Marketing Info		Marketing Start Date	Marketing End Date		

Labeler - First Aid Research Corp. (089405927)

Registrant - Anicare Pharmaceuticals Pvt. Ltd (916837425)

Establishment

Name	Address	ID/FEI	Business Operations
Anicare Pharmaceuticals Pvt. Ltd		916837425	manufacture(75983-004)

Revised: 12/2020

First Aid Research Corp.