PROCUREFIRST AID ANTIBIO FIRST AID ANTIBIOTIC- bacitracin ointment TWIN MED, LLC

Drug Facts

Active Ingredients(in each gram)

Bacitracin Zinc (equal to 500 Bacitracin units)

Purpose

Antibiotic

Uses

First aid to prevent infection in Minor cuts, Scrapes & Burns

Directions

- Clean the affected area
- Apply small amount of this product(an amount equal to the surface area of the tip of the finger) on the area 1 to 3 times daily
- May be covered with a sterile bandage

Warnings

For External Use Only.

Do not use

- In the eyes
- If you are allergic to any of the ingredients, due to possibility of anaphylactic shock
- 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 allergic reaction develops

Keep out of reach of children

• If swallowed, get medical help or contact a Poison Control Centre right away

Other information

- Store at controlled room temperature 15 ° 30 °C(59 °--86 ° F)
- Avoid excessive heat and humidity
- Tamper Evident
- Do not use if packet is torn, cut, or opened.

Inactive Ingredients

Hard Paraffin, Lanolin, Liquid Paraffin, White Soft Paraffin

Principal Display Panel

PARAFFIN (UNII: 1900E3H2ZE)



PROCUREFIRST AID ANTIBIO FIRST AID ANTIBIOTIC bacitracin ointment **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:55681-218 **TOPICAL Route of Administration Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I) **BACITRACIN** 500 U in 1 g **Inactive Ingredients Ingredient Name** Strength

LANOLIN (UNII: 7EV65EAW6H)	
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	

l	P	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
	1	NDC:55681-218- 01	0.9 g in 1 PACKET; Type 0: Not a Combination Product	09/22/2021			
	2	NDC:55681-218- 02	28.3 g in 1 TUBE; Type 0: Not a Combination Product	08/18/2022			

Marketing In	nformation			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC Monograph Drug	M004	02/05/2013		

Labeler - TWIN MED, LLC (009579330)

Revised: 1/2024 TWIN MED, LLC