# BACITRACIN ZINC- first aid antibiotic ointment Dynarex Corporation

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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1171 Bacitracin Zinc Ointment NDC 67777-008-10

1172 Bacitracin Zinc Ointment NDC 67777-008-20

1173 Bacitracin Zinc Ointment NDC 67777-008-30

1175 Bacitracin Zinc Ointment NDC 67777-008-50

1176 Bacitracin Zinc Ointment NDC 67777-008-60

# **Active Ingredient**

Bacitracin Zinc (500 units in each gram)

#### **Purpose**

First Aid Antibiotic

#### Uses

First aid to help prevent infection in minor cuts, scrapes, and burns

# **Warnings**

# For external use only

#### Do not use

- In the eyes or apply 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 if you have

deep or puncture wounds, animal bites, or serious burns

# Stop use and ask a doctor if

the condition persists or gets worse, or if a rash or other allergic reaction develops

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 room temperature
- Avoid excessive heat and humidity
- Tamper Evident. Do not use if inner seal is torn, cut or opened.

# **Inactive Ingredients**

Hard Paraffin, Liquid Paraffin, White Soft Paraffin

#### 1171 Label



### 1172 Label





1175 Label



# 1176 Label



# **BACITRACIN ZINC**

first aid antibiotic ointment

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67777-008	
Route of Administration	TOPICAL			

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89 Y4M234ES) (BACITRACIN - UNII:58 H6 RWO 52I)	BACITRACIN	500 [USP'U] in 1 g

Inactive Ingredients			
Ingredient Name	Strength		
MINERAL OIL (UNII: T5L8T28FGP)			
PARAFFIN (UNII: 19 O 0 E 3 H 2 Z E )			
PETROLATUM (UNII: 4T6 H12BN9 U)			

P	Packaging					
#	Item Code	Package Description	<b>Marketing Start Date</b>	Marketing End Date		
1	NDC:67777-008-10	1728 in 1 CASE	02/16/2018			
1		144 in 1 BOX				
1		0.9 g in 1 PACKET; Type 0: Not a Combination Product				
2	NDC:67777-008-20	72 in 1 CASE	02/16/2018			
2		1 in 1 BOX				
2		14.2 g in 1 TUBE; Type 0: Not a Combination Product				
3	NDC:67777-008-30	72 in 1 CASE	02/16/2018			
3		1 in 1 BOX				
3		28.4 g in 1 TUBE; Type 0: Not a Combination Product				
4	NDC:67777-008-50	72 in 1 CASE	02/16/2018			
4		1 in 1 BOX				
4		113 g in 1 TUBE; Type 0: Not a Combination Product				
5	NDC:67777-008-60	12 in 1 CASE	02/16/2018			
5		425 g in 1 JAR; 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	02/16/2018		

# Labeler - Dynarex Corporation (008124539)

Revised: 2/2019 Dynarex Corporation