### TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate. ointment Front Pharmaceutical PLC

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**Triple Antibiotic Ointment** 

### Active ingredients (each gram contains)

Bacitracin zinc 400 units Neomycin sulfate 3.5 mg Polymyxin B sulfate 5,000 units

### **Purpose**

First aid antibiotic

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

**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.

## Ask a doctor before use if you have

deep or puncture wounds

animal bites

serious burns.

**Stop use and ask a doctor if** condition persists or gets worse

- you need to use longer than 1 week
- a rash or other allergic reaction develops

### **Directions**

- clean the affected area and dry thoroughly.
- 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.

Inactive ingredient Mineral Oil, White Petrolatum

### Other information

store at 20° to 25°C (68° to 77°F)

**Questions?** Call 1-888-296-9067

Distributed by:

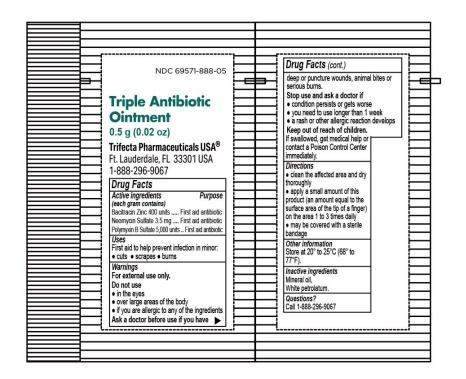
#### Trifecta Pharmaceuticals USA®

Ft. Lauderdale, FL 33301, USA

1-888-296-9067

### Package Label

# INNER PACKET



bacitracin zinc, neomycin sulfate, polymyxin b sulfate. ointment

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Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69571-888
FIGURE INDE	HOMAN OTC DROG	itelli code (Source)	NDC.03371-00

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	400 [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: J2VZ 07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g

## **Inactive Ingredients**

Ingredient Name	Strength	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)		

MINERAL OIL (UNII: T5L8T28FGP)

## **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69571-888- 05	8000 in 1 BAG	04/08/2025	
1		0.5 g in 1 PACKET; Type 0: Not a Combination Product		

## **Marketing Information**

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Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M004	04/08/2025	

# Labeler - Front Pharmaceutical PLC (530897792)

# Registrant - Trifecta Pharmaceuticals USA (079424163)

Establishment				
Name	Address	ID/FEI	<b>Business Operations</b>	
FRONT PHARMACEUTICAL PLC		530897792	manufacture(69571-888)	

Revised: 4/2025 Front Pharmaceutical PLC