

FAMILY WELLNESS- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment
FRONT PHARMACEUTICAL PLC

Active ingredients (in each gram)

Bacitracin 400 units

Neomycin 3.5 mg

Polymyxin B 5,000 units

Purpose

First aid antibiotic

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first aid to help prevent infection in minor: **Uses**

-uses -scrapes -burns

Warnings

. **For external use only**

Do not use

Ask a doctor before use if you have

Stop use and ask a doctor if

- if you are allergic to any of the ingredients in this product
- in contact with the eyes or mucous membranes
- over large areas of the body

- deep or puncture wounds
- animal bites
- serious burns

- symptoms persist for more than 7 days, the condition persists or gets worse
- symptoms clear up and then occur again within a few days
- a rash or other allergic reaction occurs

. If swallowed, get medical help or **Keep out of reach of children**

contact a Poison Control Center right away.

Directions

Adults and children 2 years of age and older:

Children under 2 years of age: ask a doctor before use.

- clean the affected area
- apply a small amount of this product to affected area 1 to 3 times daily
- May be covered with a sterile bandage.

store at room temperature 68 - 77 F **Other information**

(20-25 C).

1-800-639-3803 Weekdays 9 AM to 4 PM EST **Questions?**

mineral oil, petrolatum **Inactive ingredients**



FAMILY WELLNESS

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69571-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	0.41 g in 100 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	0.0035 g in 100 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: 232323232323)	POLYMYXIN B	0.52 g in 100 g

UNII:J2VZ07J96K) POLYMERIN B 0.52 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69571-003-02	1 in 1 BOX	02/22/2017	
1	NDC:69571-003-01	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 Drug	M004	02/22/2017	

Labeler - FRONT PHARMACEUTICAL PLC (530897792)

Establishment

Name	Address	ID/FEI	Business Operations
FRONT PHARMACEUTICAL PLC		530897792	manufacture(69571-003)

Revised: 10/2023

FRONT PHARMACEUTICAL PLC