

**TRIPLE ANTIBIOTIC AND PAIN RELIEF- bacitracin,neomycin,polymyxin,
pramoxine ointment
Meijer**

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.

Meijer Triple Antibiotic + Pain Relief Ointment

DRUG FACTS

Active Ingredient

Bacitracin zinc 500 Units

Purpose

First Aid Antibiotic

Active Ingredient

Neomycin Sulfate 3.5mg

Purpose

First Aid Antibiotic

Active Ingredient

Polymyxin B Sulfate 10,000 Units

Purpose

First Aid Antibiotic

Active Ingredient

Pramoxine HCL 10mg

Purpose

External Analgesic

Uses

First Aid to help prevent infection in minor:

- Cuts
- Scrapes
- Burns

Warnings

For external use only. Do not use:

- In eyes
- Over large areas of the body
- If you are allergic to any of the ingredients

Ask a Doctor before Use

Ask 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

Keep out of Reach of Children

If Swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- Clean the affected area 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

Other Information

- To Open: Unscrew cap, pull tab to remove foil seal
- Store at 20° to 25°C (68° to 77°F)
- See carton or tube crimp for lot number and expiration date

Inactive Ingredient:

Petrolatum

TRIPLE ANTIBIOTIC AND PAIN RELIEF

bacitracin,neomycin,polymyxin, pramoxine ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79481-0016
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN ZINC	500 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ 07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79481-0016-2	1 in 1 BOX	06/25/2021	
1		2 in 1 PACKAGE, COMBINATION		
1		28 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:79481-0016-1	1 in 1 BOX	06/25/2021	
2		28 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:79481-0016-5	1 in 1 BOX	06/25/2021	
3		14 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 final	part333B	06/25/2021	

Labeler - Meijer (006959555)

Registrant - Trifecta Pharmaceuticals USA (079424163)

Revised: 1/2023

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