

**TRIPLEANTIBIOTICOINTMENTANDPAINRELIEF- bacitracin,neomycin,polymxin,
pramoxine ointment
Trifecta Pharmaceuticals USA**

Triple Antibiotic Ointment + Pain Relief

DRUG FACTS

Active Ingredient

Bacitracin Zinc 500 Units

Purpose

First Aid Antibiotic

Active Ingredient

Neomycin 3.5mg

Purpose

First Aid Antibiotic

Active Ingredient

Polymyxin B 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 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

- 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:

mineral oil, white Petrolatum

Questions?

Call 1-888-296-9067

Distributed By:

Trifecta Pharmaceuticals USA ®


101 NE Third Avenue, Suite 1500

Fort Lauderdale, FL. 33301 USA

www.trifecta-pharma.com

Packaging

OUTER BOX



**COMPARE TO
NEOSPORIN® PLUS PAIN
RELIEF® ACTIVE INGREDIENTS**

NDC: 69396-030-01

**Soothes Painful Cuts, Scrapes and Burns
While Preventing Infection**

TRIPLE ANTIBIOTIC INFECTION PROTECTION

**First Aid Antibiotic
Pain Relieving Ointment**

BACITRACIN ZINC • NEOMYCIN SULFATE • POLYMYXIN B SULFATE • PRAMOXINE HCL


Trip

Antibiotic Ointment

**Compare to Neosporin® Plus Pain Relief® active ingredients*


Net Wt. 1 oz. (28.4g)

+Pain Relief





X001GMSO0B

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Product of PRC
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0 402269 2268 7





* This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin® Plus Pain Relief®.

DRUG FACTS

Active ingredient (each gram contains)
Bacitracin 500 units
Neomycin B 10,000 units
Polymyxin B 10,000 units
Pramoxine HCl 10mg

Uses
First aid to help prevent infection in minor: • Cuts • Scrapes • Burns

Warnings
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.

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
Light mineral oil, Petrolatum

Questions? Call 1-888-296-9067

LOT: _____

EXP: _____

100%
GUARANTEED

Triple Antibiotic Ointment
BACITRACIN ZINC • NEOMYCIN SULFATE
POLYMYXIN B SULFATE • PRAMOXINE HCL
FIRST AID ANTIBIOTIC + PAIN RELIEF

INNER TUBE

NDC: 69396-030-01



Triple Antibiotic Ointment + Pain Relief

BACITRACIN ZINC • NEOMYCIN SULFATE • POLYMYXIN B SULFATE
PRAMOXINE HCL

Net Wt. 1 oz. (28.4g)

FIRST AID ANTIBIOTIC OINTMENT

Active ingredient (each gram contains)	Purpose
Bacitracin 500 units	First aid antibiotic
Neomycin 3.5 mg	First aid antibiotic
Polymyxin B 10,000 units	First aid antibiotic
Pramoxine HCL 10mg	External analgesic

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. Keep out of reach of children. 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: • 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: Light mineral oil, Petrolatum

Questions? Call 1-888-296-9067

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OUTSIDE BOX



INNER TUBE



TRIPLEANTIBIOTICOINTMENTANDPAINRELIEF

bacitracin,neomycin,polymxin, pramoxine ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 100 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 100 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 100 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ 07J96K)	POLYMYXIN B	10000 [USP'U] in 100 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:69396-030-01	1 in 1 BOX	07/20/2017	
1		28.4 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69396-030-44	4 in 1 BOX	10/20/2024	
2		30 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	07/20/2017	

Labeler - Trifecta Pharmaceuticals USA (079424163)