

**TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin-b sulfate,
pramoxine hydrochloride ointment
Rite Aid Corporation**

Triple Antibiotic Ointment (Bacitracin Zinc, Neomycin Sulfate, Polymyxin B Sulfate)

Drug Facts

Active ingredients (in each gram)

Bacitracin Zinc USP, 500 units

Neomycin 3.5 mg

Polymyxin B sulfate USP, 10,000 units

Pramoxine hydrochloride USP, 10 mg

Purpose

First aid antibiotic

Pain reliever

Uses

First aid to help prevent infection and for the temporary relief of pain in minor

- cuts
- scrapes
- burns

Warnings

For external use only

Do not use

- if you are allergic to any of the ingredients
- in the eyes
- over large areas of the body

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- you need to use for longer than one week
- condition persists or gets worse
- rash or other allergic reaction develops
- symptoms persist for more than one week, or clear up and occur again with a few days

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

- adults and children 2 years of age and older
 - 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
- children under 2 years of age: ask a doctor

Other information

- Store at 15°-30°C (59°-86°F)
- Before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredient

white petrolatum

Questions or comments?

Call **1-877-753-3935** Monday-Friday 9AM-5PM EST

Principal display panel

Compare to the active ingredients in Neosporin® + Pain Relief Ointment*

MAXIMUM STRENGTH

TRIPLE ANTIBIOTIC OINTMENT + PAIN RELIEF

BACITRACIN ZINC

NEOMYCIN SULFATE

POLYMYXIN B SULFATE

PRAMOXINE HCl

FIRST AID ANTIBIOTIC /

PAIN-RELIEVING OINTMENT

FOR EXTERNAL USE ONLY

HELP PREVENT INFECTION AND

TEMPORARILY RELIEVES PAIN OF MINOR CUTS, SCRAPES, AND BURNS

NET WT OZ (g)

*This product is not manufactured or distributed by Johnson & Johnson Consumer, Inc distributor of Neosporin® + Pain Relief Ointment.

DISTRIBUTED BY:

RITE AID, 30 HUNTER LANE,

CAMP HILL, PA 17011

www.riteaid.com

Package label



Compare to the active ingredients in Neosporin® + Pain Relief Ointment*

NDC 11822-2700-2

MAXIMUM STRENGTH

TRIPLE ANTIBIOTIC OINTMENT + PAIN RELIEF

BACITRACIN ZINC
NEOMYCIN SULFATE
POLYMYXIN B SULFATE
PRAMOXINE HCl

FIRST AID ANTIBIOTIC / PAIN RELIEVING OINTMENT FOR EXTERNAL USE ONLY HELPS PREVENT INFECTION AND TEMPORARILY RELIEVES PAIN OF MINOR CUTS, SCRAPES, AND BURNS

2-1 OZ (28 g) TUBES
NET WT 2.0 OZ (56 g)

DISTRIBUTED BY:
RITE AID, 30 HUNTER LANE,
CAMP HILL, PA 17011
www.riteaid.com

SATISFACTION GUARANTEE

If you're not satisfied, we'll happily refund your money.

PLD-A692B FC007994



Code: GO/DRUGS/362 ABM3B/01

Drug Facts

Active ingredients (in each gram)	Purposes
Bacitracin zinc, USP 500 units	}First aid antibiotic
Neomycin 3.5 mg	
Polymyxin B sulfate, USP 10,000 units	
Pramoxine hydrochloride, USP 10 mgPain reliever

Uses First aid to help prevent infection and for the temporary relief of pain in minor ■ cuts ■ scrapes ■ burns

Warnings
For external use only.

Do not use
■ if you are allergic to any of the ingredients ■ in the eyes
■ over large areas of the body

Ask a doctor before use if you have ■ deep or puncture wounds
■ animal bites ■ serious burns

Stop use and ask a doctor if
■ you need to use longer than one week
■ condition persists or gets worse
■ rash or other allergic reaction develops
■ symptoms persist for more than one week, or clear up and occur again within a few days

Drug Facts (continued)

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions
■ adults and children 2 years of age and older
■ 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
■ children under 2 years of age: ask a doctor

Other information
■ store at 15°-30°C (59°-86°F)
■ before using any medication, read all label directions. Keep carton, it contains important information.

Inactive ingredient white petrolatum

Questions or comments?
Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

*This product is not manufactured or distributed by Johnson & Johnson Consumer, Inc., distributor of Neosporin® + Pain Relief Ointment.

MAXIMUM STRENGTH

TRIPLE ANTIBIOTIC OINTMENT + PAIN RELIEF

BACITRACIN ZINC • NEOMYCIN SULFATE
POLYMYXIN B SULFATE • PRAMOXINE HCl

FIRST AID ANTIBIOTIC / PAIN RELIEVING OINTMENT

RITE AID Maximum Strength Triple Antibiotic Ointment and Pain relief

TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin-b sulfate, pramoxine hydrochloride ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11822-2700
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11822-2700-5	1 in 1 CARTON	02/25/2022	
1		14 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:11822-2700-1	1 in 1 CARTON	02/25/2022	
2		28 g in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:11822-2700-2	2 in 1 CARTON	02/25/2022	
3		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	02/25/2022	

Labeler - Rite Aid Corporation (014578892)

Revised: 5/2024

Rite Aid Corporation