TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment
Walgreen Company

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.

Triple Antibiotic
Plus Pain Relief

Drug Facts

Active ingredients (in each gram)	***Purpose***
Bacitracin zinc USP 500 units	First aid antibiotic
Neomycin sulfate USP 3.5 mg	First aid antibiotic
Polymyxin B sulfate USP 10,000 units	First aid antibiotic
Pramoxine HCl USP 10 mg	External analgesic

Uses

first aid to help prevent infection and for the temporary relief of pain or discomfort 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 1 week
- condition persists or gets worse
- symptoms persist for more than 1 week, or clear up and occur again within a few days
- 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

- 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

- **TAMPER EVIDENT: DO NOT USE IF THE SEAL ON THE TUBE IS PUNCTURED OR NOT VISIBLE.**
- store at 20° to 25°C (68° to 77°F)
- to open: unscrew cap, pull tab to remove foil seal
- see carton or tube crimp for lot number and expiration date

Inactive ingredient

white petrolatum

Questions?

Call **1-866-923-4914**

DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

Walgreens

Compare to Neosporin® + Pain Relief
active ingredients††

NDC 0363-8888-30

Triple Antibiotic
Ointment + Pain Relief

BACITRACIN ZINC / NEOMYCIN SULFATE /
POLYMYXIN B SULFATE / PRAMOXINE HCl

FIRST AID ANTIBIOTIC /
PAIN-RELIEVING OINTMENT

TRIPLE ANTIBIOTIC
MAXIMUM STRENGTH

- First aid antibiotic
- Pain-relieving ointment
- Helps prevent infection in minor cuts, scrapes & burns plus maximum strength pain relief

NET WT 1 OZ (28.4 g)

51
1220
5215025

Walgreens

Compare to Neosporin® + Pain Relief active ingredients††

NDC 0363-8888-30

Triple Antibiotic Ointment + Pain Relief

BACITRACIN ZINC / NEOMYCIN SULFATE / POLYMYXIN B SULFATE / PRAMOXINE HCl
FIRST AID ANTIBIOTIC / PAIN-RELIEVING OINTMENT

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PHARMACIST RECOMMENDED†
Walgreens
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Health expertise you rely on.™

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Drug Facts (continued)

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Walgreens Pharmacist Survey

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

PAPER BOX | PLASTIC TUBE

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DISTRIBUTED BY: WALGREEN CO.
200 WILMOT RD., DEERFIELD, IL 60015

Walgreens 100% SATISFACTION GUARANTEED

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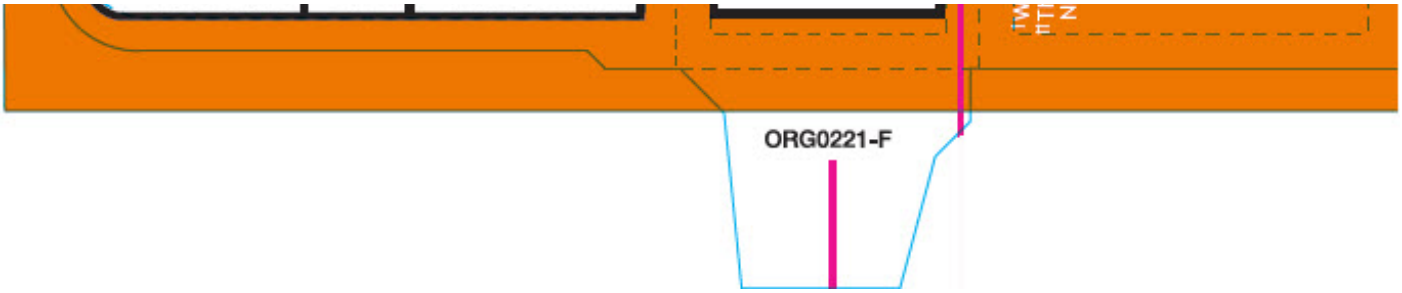

TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

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

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0363-8888
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Bacitracin zinc (UNII: 89Y4M234ES) (Bacitracin - UNII:58H6RWO52I)	Bacitracin	500 [USP'U] in 1 g
Neomycin sulfate (UNII: 057Y626693) (Neomycin - UNII:I16QD7X297)	Neomycin sulfate	3.5 mg in 1 g
Polymyxin B sulfate (UNII: 19371312D4) (Polymyxin B - UNII:J2VZ07J96K)	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:0363-8888-15	1 in 1 CARTON	08/21/2020	
1		14.2 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:0363-8888-30	1 in 1 CARTON	08/21/2020	
2		28.4 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	03/31/2012	

Labeler - Walgreen Company (008965063)

Revised: 4/2021 Walgreen Company