

FAMILY WELLNESS PLUS PAIN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, pramoxine hcl ointment

Trifecta Pharmaceutical USA LLC

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 Ointment + Pain Relief

DRUG FACTS

Active Ingredient

Bacitracin 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 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:

Petrolatum

Questions?

Call 1-888-296-9067

Weekdays 9AM - 4PM EST

DISTRIBUTED BY: MIDWOOD BRANDS, LLC.

500 Volvo Parkway, Chesapeake, VA. 23320

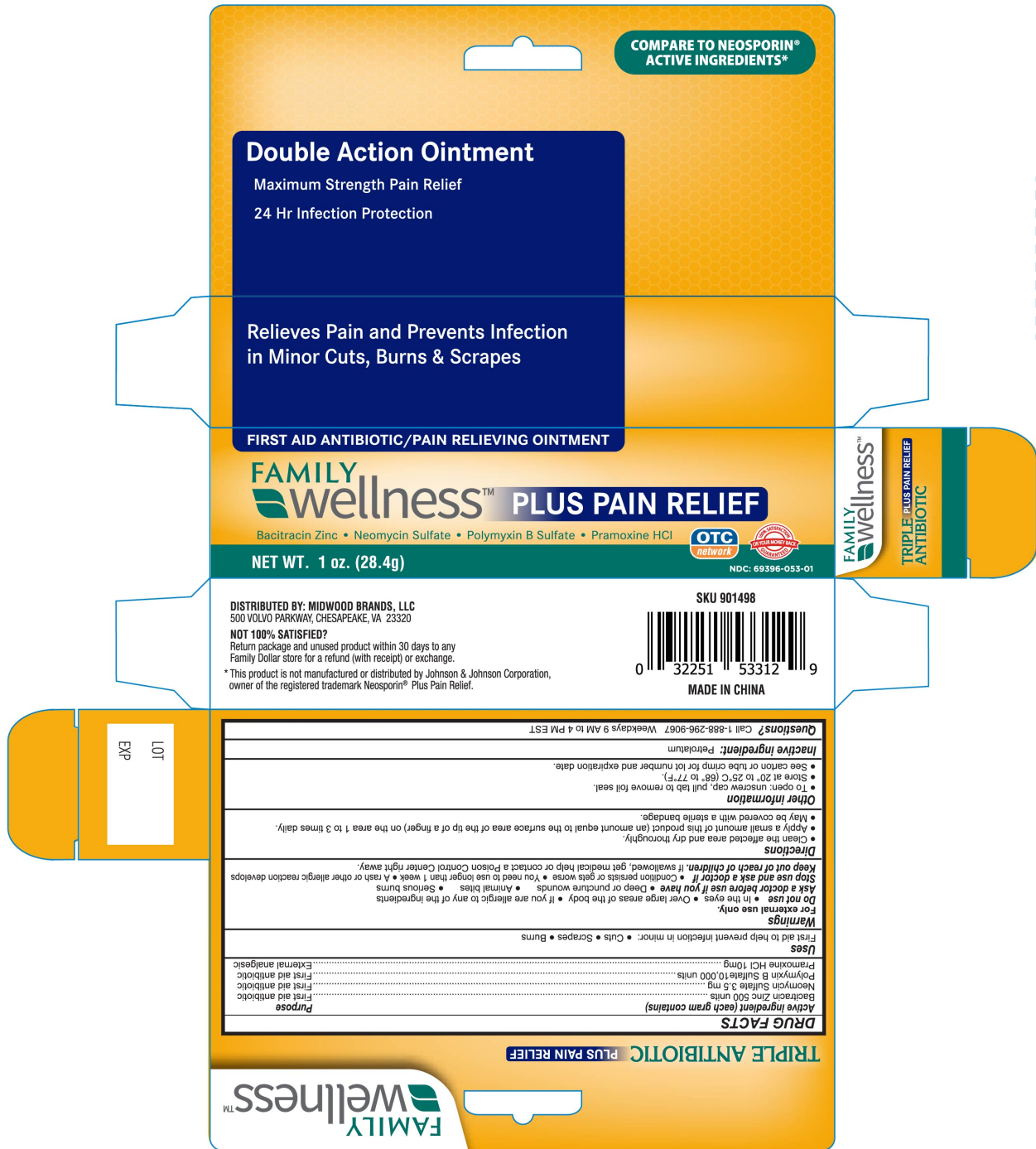
NOT SATISFIED?

Return package and unused product within 30 days to any Family Dollar store for a refund (with receipt) or exchange.

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

Packaging

OUTSIDE BOX



INNER TUBE



Pramoxine HCl 10mg External analgesic	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.
Uses: First aid to help prevent infection in minor: • Cuts • Scrapes • Burns	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.
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. ▶	Inactive ingredients: Petrolatum
Questions? Call 1-888-295-9267 Weekdays 9 AM to 4 PM EST	
DISTRIBUTED BY: MIDWOOD BRANDS, LLC 500 VOLVO PARKWAY, CHESAPEAKE, VA 23320 MADE IN CHINA	
NOT 100% SATISFIED? Return package and unused product within 30 days to any Family Dollar store for a refund (with receipt) or exchange.	

FAMILY WELLNESS PLUS PAIN RELIEF

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, pramoxine hcl ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-053
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
PRAMO XINE HYDRO CHLORIDE (UNII: 88AYB867L5) (PRAMO XINE - UNII:068X84E056)	PRAMO XINE HYDROCHLORIDE	10 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	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:69396-053-01	1 in 1 BOX	09/22/2019	
1		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	08/30/2019	

Labeler - Trifecta Pharmaceutical USA LLC (079424163)