

**PREMIER VALUE MAX STRENGTH ANTIBIOTIC PLUS PAIN RELIEF- premier value neomycin sulfate, polymyxin b sulfate, pramoxine hcl cream
Pharmacy Value Alliance LLC**

Maximum Strength Antibiotic Cream plus Pain Relief

DRUG FACTS

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 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:

Cetostearyl alcohol, Ethylparaben, Glycerin, Glyceryl Sterate, Light Mineral Oil, Petrolatum, Polyoxyethylene lauryl ether, Purified Water, Sodium Lauryl Sulfate.

Questions or Comments

Call 1-866-923-4914

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

Distributed By:

Pharmacy Value Alliance, LLC.

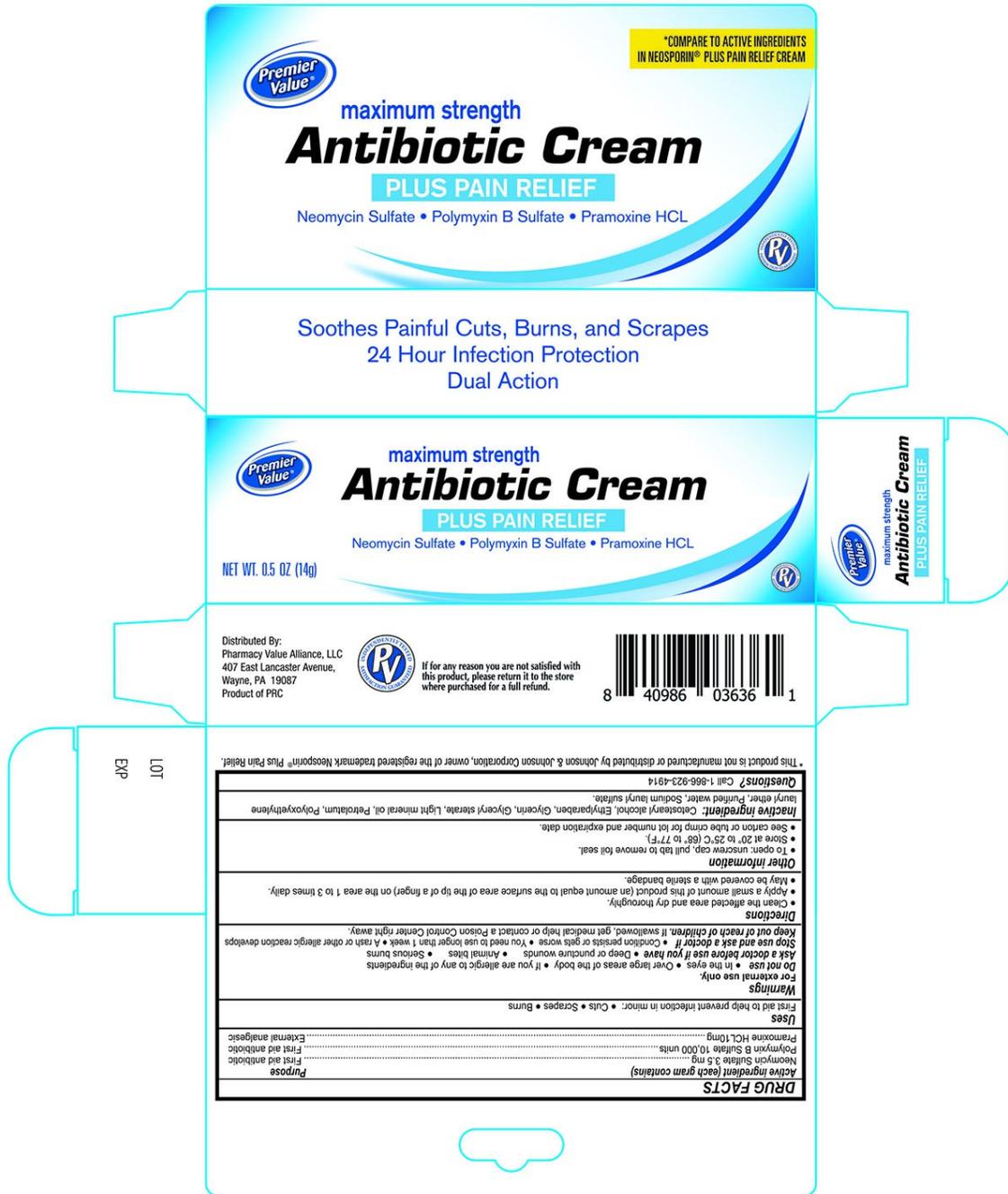
407 East Lancaster Avenue

Wayne, PA. 19087

Product of PRC

Packaging

OUTER BOX



INNER TUBE



Active ingredient (each gram contains) Neomycin Sulfate 3.5 mg Polymyxin B Sulfate 10,000 units Pramoxine HCL 10mg	Purpose First aid antibiotic First aid antibiotic External analgesic	Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.
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. ▶	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 ingredients: Cetostearyl alcohol, Ethylparaben, Glycerin, Glyceryl stearate, Light mineral oil, Petrolatum, Polyoxyethylene lauryl ether, Purified water, Sodium lauryl sulfate.
		Questions? Call 1-866-923-4914
Distributed By: Pharmacy Value Alliance, LLC 407 East Lancaster Avenue, Wayne, PA 19087 Product of PRC		

PREMIER VALUE MAX STRENGTH ANTIBIOTIC PLUS PAIN RELIEF

premier value neomycin sulfate, polymyxin b sulfate, pramoxine hcl cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:68016-296
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
SODIUM DODECYLBENZENESULFONATE (UNII: 554127163Y)	
PETROLATUM (UNII: 4T6H12BN9U)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
WATER (UNII: 059QF0KO0R)	
ETHYLPARABEN (UNII: 14255EXE39)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:68016-296-05	1 in 1 BOX	09/22/2019	
1		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 Drug	M004	09/22/2019	

Labeler - Pharmacy Value Alliance LLC (101668460)

Registrant - Trifecta Pharmaceuticals USA LLC (079424163)

Revised: 12/2024

Pharmacy Value Alliance LLC