

WALGREENS KIDS MULTI-ANTIBIOTIC WITH PAIN RELIEF CREAM- polymyxin b sulfate, neomycin sulfate, pramoxine hydrochloride cream
Walgreens Company

Drug Facts

Active ingredients (in each gram)	Purpose
Neomycin Sulfate (3.5mg)	First Aid Antibiotic
Polymixin B Sulfate (10,000 units)	First Aid Antibiotic
Pramoxine Hydrochloride (10 mg)	External Analgesic

Purpose

First Aid Antibiotic
First Aid Antibiotic
External Analgesic

Uses

first aid to prevent infection and for the temporary relief of pain or discomfort in minor:
cuts
scapes
burns

Warnings

For external use only

Keep this and all drugs out of the reach of children

If swallowed, get medical help or contact 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

store at 20° to 25°C (68° to 77°F)

Inactive ingredients

Ceteareth-20, Cetearyl Alcohol, Glyceryl Stearate SE, Methylparaben, Mineral Oil, Phosphoric Acid, Propylene Glycol, Purified Water, White Petrolatum

Questions?

Call 1-800-222-1087

Principal Display Panel -Tube

Walgreens Children Multi Antibiotic +Pain Relief Cream NDC 0363-2122-14

Neomycin Sulfate /Polymyxin B Sulfate /Pramoxine HCL

First Aid Antibiotic / Pain Relieving Cream

NET WT 0.5 OZ (14.2g)



Principal Display Panel -Carton

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Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:0363-2122

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
POLYOXYL 20 CETOSTEARYL ETHER (UNII: YRC528SWUY)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PHOSPHORIC ACID (UNII: E4GA8884NN)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-2122-14	1 in 1 CARTON	12/01/2022	
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	12/01/2022	

Labeler - Walgreens Company (008965063)

Revised: 11/2024

Walgreens Company