

PUBLIX ANTIBIOTIC PLUS PAIN RELIEF- neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream
Publix Super Markets Inc

Publix
antibioticcream plus Pain Relief

Drug Facts

<i>Active ingredients (each gram contains)</i>	<i>Purpose</i>
Neomycin sulfate 3.5 mg	First aid antibiotic
Polymyxin B sulfate 10,000 units	First aid antibiotic
Pramoxine hydrochloride 10 mg	Topical 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

- in the eyes
- over large areas of the body
- if you are allergic to any of the ingredients
- longer than 1 week

Ask a doctor before use

- on deep or puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if

- condition gets worse
- condition persists for more than 7 days
- condition clears up and occurs again within a few days
- 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

Adults and children 2 years of age and older:

- 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

Children under 2 years of age: consult a doctor

Other information

- To open: unscrew cap, pull tab to remove foil seal
- store at controlled room temperature
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

emulsifying wax, methylparaben, mineral oil, propylene glycol, purified water, white petrolatum

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SUPER MARKETS, INC.
3300 PUBLIX CORPORATE
PARKWAY
LAKELAND, FL 33811**

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

Publix

antibioticcream+

PAIN

RELIEF

NEOMYCIN SULFATE • POLYMYXIN B SULFATE • PRAMOXINE HCl

FIRST AID ANTIBIOTIC CREAM PLUS PAIN RELIEVER

NET WT 1 OZ (28.4g)

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41415-076	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
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 [iU] in 1 g	
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)		PRAMOXINE HYDROCHLORIDE	10 mg in 1 g	
Inactive Ingredients				
Ingredient Name			Strength	
METHYLPARABEN (UNII: A2I8C7HI9T)				
MINERAL OIL (UNII: T5L8T28FGP)				
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)				
WATER (UNII: 059QF0KO0R)				
PETROLATUM (UNII: 4T6H12BN9U)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41415-076-75	1 in 1 CARTON	03/21/2012	
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 Drug	M004	03/21/2012		

Labeler - Publix Super Markets Inc (006922009)

Establishment

Name	Address	ID/FEI	Business Operations
Sun Pharma Canada Inc.		243339023	manufacture(41415-076)

Revised: 7/2025

Publix Super Markets Inc