

ANTIBIOTIC PLUS PAIN RELIEF- neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream
TARGET Corporation

Antibiotic 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	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

- 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
- 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 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: ask a doctor

Other information

- store at 20° to 25°C (68° to 77°F)
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

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

Questions?

Call **1-800-910-6874**

Dist. by Target Corp.
Mpls., MN 55403

PRINCIPAL DISPLAY PANEL - 14.2 g Tube Carton

up & up

maximum strength

antibiotic cream + pain relief

neomycin sulfate, polymyxin b sulfate, pramoxine HCl

first aid antibiotic / pain relieving cream

helps prevent infection in minor cuts, scrapes and burns plus
maximum strength pain relief

NET WT 0.5 OZ (14.2 g)

NDC 11673-393-01



Compare to active ingredients in Neosporin® + Pain Relief*

maximum strength
antibiotic cream + pain relief
neomycin sulfate, polymyxin b sulfate, pramoxine HCl
first aid antibiotic / pain relieving cream

5213029
1020
17



maximum strength
antibiotic cream + pain relief
neomycin sulfate, polymyxin b sulfate, pramoxine HCl
first aid antibiotic / pain relieving cream
helps prevent infection in minor cuts, scrapes and burns plus
maximum strength pain relief

NET WT 0.5 OZ (14.2 g)



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245 07 0071 00000-0X
11673 39301 2

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Drug Facts (continued)
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• see carton or tube crimp for lot number and expiration date
Inactive ingredients emulsifying wax, methylparaben, mineral oil, propylene glycol, purified water, white petrolatum
Questions? Call 1-800-910-8774

NO COPY / NO COLOR
THIS FLAP FOR LOT #
AND EXP DATE PRINT

*This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin® + Pain Relief.
Dkt. by Target Corp., Mpls., MN 55403
Made in Canada
TM & ©2021
Target Brands, Inc.

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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-393
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
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	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:11673-393-01	1 in 1 CARTON	10/16/2020	
1		14.2 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	M017	10/16/2020	

Labeler - TARGET Corporation (006961700)

Establishment

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	manufacture(11673-393)