ANTIBIOTIC AND PAIN RELIEF- neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream
Target Corporation

Target Up &Up Antibiotic+Pain Relief Cream

Drug Facts

Active ingredients

Neomycin Sulfate (3.5mg)
Polymixin B Sulfate (10,000 units)
Pramoxine Hydrochloride (10 mg)

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

Allergy alert

- do not use if you are allergic to any of the ingredients

Do not use

- in or near the eyes
- or on large areas of the body

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
- symptoms last for more than 7 days or clear up and come back within a few days

- a rash or other allergic reaction develops

Keep out of the reach of children. If swallowed get medical help or contact a Poison Control Center immediately.

Directions

- clean the affected area
- apply a small amount (equal to surface area of tip of finger) on the area 1 to 3 times daily.
- may be covered with a sterile bandage

Other inofrmation

Store at a controlled room temperature 68º-77ºF (20º-25ºC)

Inactive ingredients

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

Questions or comment? 1-800-910-6874

Principal Display Panel

Target up&Up NDC 11673-770-14

Maximum Strength Pain Releiving & Infection Protection

Neomycin Sulfate/ Polymyxin B Sulfate / Pramoxine HCL

NET WT 0.5oz (14g)

Bend Line

up&up™

Maximum Strength Pain Relieving & Infection Protection

First Aid Antibiotic Cream

Neomycin Sulfate/Polymyxin B Sulfate/ Pramoxine HCl

- Helps soothe minor cuts, scrapes and burns
- Maximum strength pain relief and infection protection

NET WT 0.5 OZ (14 g)

NDC 11673-770-14

Center Front Panel

Bend Line

Active ingredients (in each gram) Neomycin 3.5 mg, Polymyxin B 10,000 units, and Pramoxine HCl 10 mg ***Purposes*** First aid antibiotic and External analgesic ***Uses*** first aid to help prevent infection and for temporary relief of pain or discomfort in minor: • cuts • scrapes • burns ***Warnings*** **For external use only. Allergy alert:** • do not use if allergic to any of the ingredients **Do not use** • in or near the eyes • on large areas of the body. **Ask a doctor before use if you have** • deep or puncture wounds • animal bites • serious burns. **When using this product** • do not use longer than 1 week. **Stop use and ask a doctor if** • condition lasts or gets worse • symptoms last for more than 7 days or clear up and come back within a few days • a rash or other allergic reaction occurs. **Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center immediately. ***Directions*** Adults and children 2 years and older: • clean affected area • apply a small amount (equal to the surface area of tip of finger) on area 1 to 3 times daily. • may be covered with a sterile bandage. Children under 2 years: ask a doctor. ***Other information*** Store at controlled room temperature 20°-25°C (68°-77°F). ***Questions or comments?*** **1-800-910-6874**

Center Back Panel

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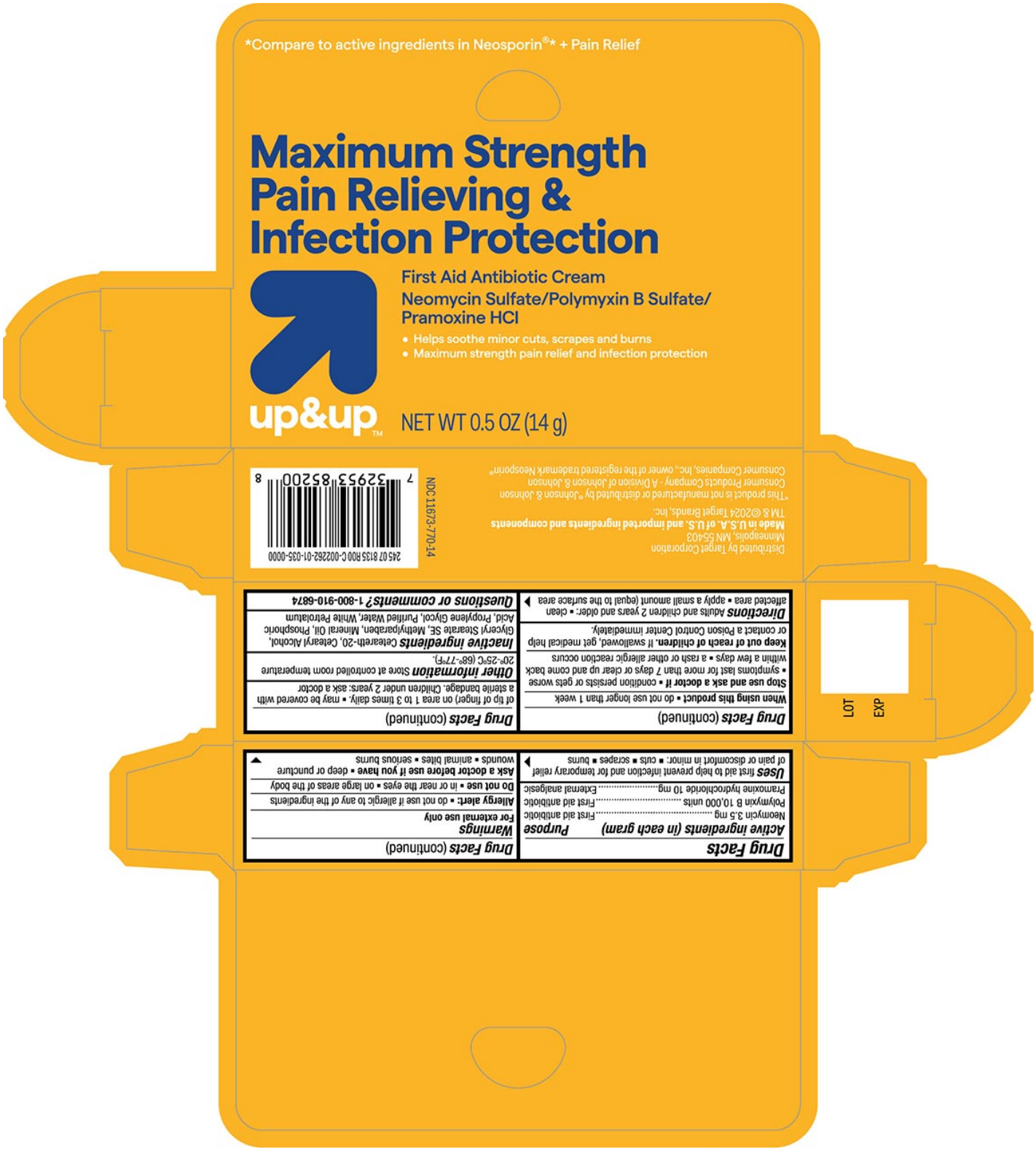

ANTIBIOTIC AND PAIN RELIEF

neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-770
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)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-770-14	1 in 1 CARTON	01/31/2024	
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	01/31/2024	

Labeler - Target Corporation (006961700)

Revised: 1/2024 Target Corporation