TRIPLE ANTIBIOTIC AND PAIN RELIEF- triple antibiotic ointment
Target Corporation

Target Up &Up Maximum Strength Triple Antibiotic+Pain Relief

Drug Facts

Active ingredients (in each gram)

Bacitracin 500

Neomycin 3.5 mg

Polymyxin B 10,000 units

Pramoxine HCL 10 mg

Purpose

First aid antibiotic

First aid antibiotic

First aid antibiotic

Pain Relief

Uses

helps prevent infection in and temporarily relieves pain due to minor:

- cuts
- scrapes
- burns

Warnings
For external use only

Do not use

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

- you need to use longer than 1 week
- 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 information

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

Inactive ingredient

Petrolatum

Questions or comments? 1-800-910-6874

Principal Display panel

Target Up & Up NDC 11673-746-28

Maximum Strength Pain Relieving

First aid Antibiotic Ointment

Polymixin B Sulfate/Bacitracin Zinc/Neomycin Sulfate/ Pramoxine HCl

NET WT 1.0 OZ (28g)

up&up™

Maximum Strength Pain Relieving

First Aid Antibiotic/Pain Relieving Ointment

Polymyxin B Sulfate/Bacitracin Zinc/ Neomycin Sulfate/Pramoxine HCl

- Soothes cuts, scrapes and burns
- Helps prevent infection in minor cuts, scrapes and burns

NET WT 1 OZ (28 g)

NDC 11673-746-28

Active ingredients (in each gram) Bacitracin 500 units, Neomycin 3.5 mg, Polymyxin B 10,000 units, Pramoxine hydrochloride 10 mg ***Purpose*** First aid antibiotic, First aid antibiotic, First aid antibiotic, External analgesic ***Uses*** first aid to help prevent infection and for the temporary relief of pain in minor: ▪ cuts ▪ scrapes ▪ burns ***Warnings*** **For external use only Do not use** ▪ if you are allergic to any of the ingredients ▪ 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 **Stop use and ask a doctor if** ▪ you need to use longer than 1 week ▪ 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 the 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 surface area of tip of finger) on the area 1 to 3 times daily. ▪ may be covered with a sterile bandage. Children 2 years and older: ask a doctor ***Other information*** Store at a controlled room temperature 20°-25°C (68°-77°F)

Questions or comments? **1-800-910-6874**

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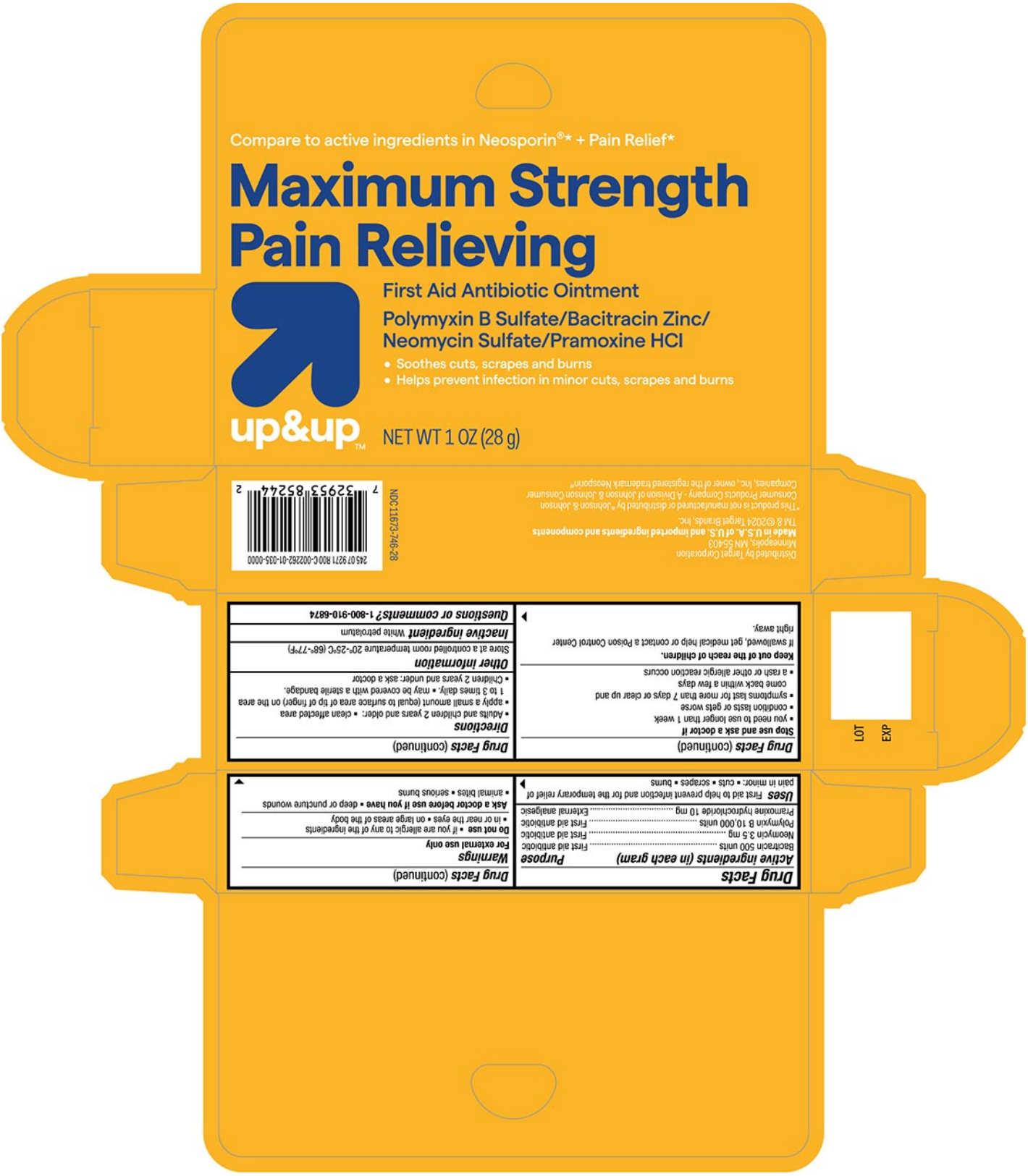

TRIPLE ANTIBIOTIC AND PAIN RELIEF

triple antibiotic ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11673-746
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g
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
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-746-28	1 in 1 CARTON	01/31/2024	
1		28 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