

**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 68°-77°F (20°-25°C)

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)



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

Dist. by Target Corp., Mpls., MN 55403 TM & ©2024 Target Brands, Inc.

245 07 9271 R00 C-002262-01-035

Compare to active ingredients in Neosporin®* + Pain Relief*

Maximum Strength Pain Relieving



up&up™

First Aid Antibiotic 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)



Distributed by Target Corporation
Minneapolis, MN 55403
Made in U.S.A. of U.S. and imported ingredients and components
TM & ©2024 Target Brands, Inc.
This product is not manufactured or distributed by Johnson & Johnson Consumer Products Company - A Division of Johnson & Johnson
Companies, Inc., owner of the registered trademark Neosporin®

<p>Drug Facts (continued)</p> <p>Stop use and ask a doctor if</p> <ul style="list-style-type: none"> • 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 <p>Keep out of the reach of children. If swallowed, get medical help or contact a Poison Control Center right away.</p>	<p>Drug Facts (continued)</p> <p>Active Ingredients (in each gram)</p> <p>Bacitracin 500 units First aid antibiotic Neomycin 3.5 mg First aid antibiotic Polymyxin B 10,000 units First aid antibiotic Pramoxine hydrochloride 10 mg External analgesic</p> <p>Uses First aid to help prevent infection and for the temporary relief of pain in minor: • cuts • scrapes • burns</p>
<p>Drug Facts (continued)</p> <p>Directions</p> <ul style="list-style-type: none"> • 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 under: ask a doctor <p>Other Information</p> <p>Store at a controlled room temperature 20°-25°C (68°-77°F)</p> <p>Inactive ingredient White petrolatum</p> <p>Questions or comments? 1-800-910-6874</p>	<p>Drug Facts (continued)</p> <p>Warnings</p> <p>Do not use • if you are allergic to any of the ingredients For external use only</p> <p>Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns</p>



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:J2VZ.07J96K)	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