

MULTI-ACTION ANTIBIOTIC AND PAIN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment ointment
Target Corporation

Target Up &Up Multi-Action Antibiotic Plus Ointment

Drug Facts

Active Ingredients

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 Ingredients

Gossypium Herbaceum (Cotton) Seed Oil, Olea Europaea (Olive) Fruit Oil, Petrolatum, Sodium Pyruvate, Theobroma Cacao (Cocoa) Seed Butter, Tocopherol

Questions or comments ? 1-800-910-6874**Principal Display Panel**

Target Up &Up NDC 11673-

Multiaction Pain Itch Scar Antibiotic ointment

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

NET WT 0.5OZ (14g)



Multi-Action Pain Itch Scar

First Aid Antibiotic/
Pain Relieving Ointment

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

- Long-lasting infection protection
- Powerful pain and itch relief for cuts, scrapes and burns
- Minimizes the appearance of scars

NET WT 0.5 OZ (14 g)

NDC 11673-875-14

Active ingredients (in each gram) Bacitracin 500 units, Neomycin 3.5 mg, Polymyxin B 10,000 units, Pramoxine hydrochloride 10mg **Purpose** First aid antibiotic, First aid antibiotic, First aid antibiotic, 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. Do not use** • if you are allergic to any of the ingredients • in or near the eyes • over 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 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 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

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245 07 3559 R00 C-002262-01-035

Compare to active ingredients in Neosporin®* +Pain +Itch +Scar*

Multi-Action Pain Itch Scar



up&up™

First Aid Antibiotic/Pain Relieving Ointment

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

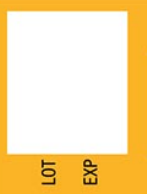
- Long-lasting infection protection
- Powerful pain and itch relief for cuts, scrapes and burns
- Minimizes the appearance of scars

NET WT 0.5 OZ (14 g)



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Minneapolis, MN 55403
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*This product is not manufactured or distributed by Johnson & Johnson Consumer Companies, Inc., owner of the registered trademark Neosporin®

<p>Drug Facts (continued)</p> <p>Directions Adults and children 2 years and older • clean affected area • apply a small amount (equal to the surface area of fingertip) 1 to 3 times daily or contact a Poison Control Center immediately.</p> <p>Keep out of reach of children. If swallowed, get medical help days • rash or other allergic reaction develops</p> <p>Stop use and ask a doctor if • you need to use longer than 1 week • condition persists or gets worse • symptoms persist for more than 1 week or clear up and occur again within a few days</p> <p>Other information Store at controlled room temperature 20°-25°C (68°-77°F).</p> <p>Inactive ingredients Gossypium Herbaceum (Cotton) Seed Oil, Olea Europaea (Olive) Fruit Oil, Petrolatum, Sodium Pyruvate, Theobroma Cacao (Cocoa) Seed Butter</p> <p>Questions or comments? 1-800-910-6874</p>	<p>Drug Facts (continued)</p> <p>Active ingredients (in each gram) Bacitracin 500 units Neomycin 3.5 mg Polymyxin B 10,000 units Pramoxine hydrochloride 10mg External analgesic</p> <p>Purpose First aid antibiotic First aid antibiotic First aid antibiotic</p> <p>Warnings For external use only</p> <p>Do not use • if you are allergic to any of the ingredients • in or near the eyes • over large areas of the body</p> <p>Ask a doctor before use if you have • deep or puncture wounds • animal bites • serious burns</p>
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MULTI-ACTION ANTIBIOTIC AND PAIN RELIEF

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-875
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
LEVANT COTTON OIL (UNII: N5CFT140R8)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM PYRUVATE (UNII: POD38AIF08)	
COCOA BUTTER (UNII: 512OYT1CRR)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-875-14	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

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