

**UP AND UP MULTI ACTION ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b, pramoxine hcl ointment**

**Target Corporation**

*Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.*

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**Target Corporation Multi-Action Antibiotic Ointment Drug Facts**

**Active ingredients (in each gram)**

Bacitracin zinc 500 units

Neomycin sulfate equivalent to 3.5 mg of neomycin base

Polymyxin B 10,000 units as polymyxin B sulfate

Pramoxine HCl 10 mg

**Purpose**

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 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 right away (1-800-222-1222).

**Directions**

- adults and children 2 years of age and older:
- clean the affected area
- 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 20-25°C (68-77°F)

**Inactive ingredients**

cocoa butter, cottonseed oil, olive oil, petrolatum, sodium pyruvate, tocopheryl acetate

**Questions or comments?**

1-888-547-7400

**Package/Label Principal Display Panel**

Compare to active ingredients in Neosporin® + Pain Itch Scar

pain itch scar

multi-action antibiotic ointment

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 11673-562-58



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

pain itch scar  
**multi-action antibiotic ointment**

first aid antibiotic/pain relieving ointment  
bacitracin zinc/neomycin sulfate/polymyxin B sulfate/pramoxine HCl



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

pain itch scar  
**multi-action antibiotic ointment**

first aid antibiotic/pain relieving ointment  
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powerful pain and itch relief for cuts, scrapes and burns  
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\*This product is not affiliated with, manufactured by,  
or produced by the makers of Neosporin®.

4R058 UW C1

Code Area



2156

<b>Drug Facts</b>		<b>Drug Facts (continued)</b>	
<b>Active ingredients (in each gram)</b>	<b>Purpose</b>	<b>Stop use and ask a doctor if</b>	
Bacitracin zinc 500 units.....	First aid antibiotic	■ you need to use longer than 1 week	
Neomycin sulfate equivalent to 3.5 mg of neomycin base.....	First aid antibiotic	■ condition persists or gets worse	
Polymyxin B 10,000 units as		■ symptoms persist for more than 1 week, or clear up and occur again within a few days	
polymyxin B sulfate.....	First aid antibiotic	■ rash or other allergic reaction develops	
Pramoxine HCl 10 mg.....	External analgesic		
<b>Uses</b>	first aid to help prevent infection and for temporary relief of pain or discomfort in minor:	<b>Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away (1-800-222-1222).</b>	
■ cuts ■ scrapes ■ burns			
<b>Warnings</b>		<b>Directions</b>	
For external use only.		■ adults and children 2 years of age and older:	
<b>Do not use</b>		■ clean the affected area	
■ if you are allergic to any of the ingredients		■ 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	
■ in the eyes		■ may be covered with a sterile bandage	
■ over large areas of the body		■ children under 2 years of age: ask a doctor	
<b>Ask a doctor before use if you have</b>		<b>Other information</b>	■ store at 20-25°C (68-77°F)
■ deep or puncture wounds		<b>Inactive ingredients</b>	cocoa butter, cottonseed oil, olive oil, petrolatum, sodium pyruvate, tocopheryl acetate
■ animal bites		<b>Questions or comments?</b>	1-888-547-7400
■ serious burns			



**UP AND UP MULTI ACTION ANTIBIOTIC**  
 bacitracin zinc, neomycin sulfate, polymyxin b, pramoxine hcl ointment

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11673-562
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
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [USP]U in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g

Inactive Ingredients	
Ingredient Name	Strength
COCOA BUTTER (UNII: 512OYT1CRR)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
PETROLATUM (UNII: 4T6H12BN9U)	
SODIUM PYRUVATE (UNII: POD38AIF08)	
COTTONSEED OIL (UNII: H3E878020N)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	

### Product Characteristics

<b>Color</b>	YELLOW (Pale)	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11673-562-58	1 in 1 CARTON	05/11/2017	
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 final	part333B	05/11/2017	

**Labeler** - Target Corporation (006961700)

Revised: 5/2017

Target Corporation