# CURAD MAXIMUM STRENGTH TRIPLE ANTIBIOTIC- bacitracin, neomycin, polymyxin b, pramoxine hydrochloride ointment Medline Industries, LP

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## 031 Curad Maximum Strength Triple Antibiotic

### Active ingredients (in each gram)

Bacitracin 500 units

Neomycin 3.5 mg

Polymyxin B 10,000 units

Pramoxin hydrochloride 10 mg

### **Purposes**

First aid antibiotic

First aid antibiotic

First aid antibiotic

Pain reliever

#### Uses

- First aid to help prevent infection in and temporarily relieves pain due to
- minor cuts
- scrapes
- burns

## Warnings

• For external use only

#### Do not use

- if 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 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 68°-77°F (20°-25°C).

## Inactive ingredient

petrolatum

## **Manufacturing Information**

Manufactured for:

Medline Industries, LP

Three Lakes Drive Northfield, IL 60093 USA

Made in USA of domestic and imported materials www.medline.com

1-800-MEDLINE (633-5463)

REF: CUR001232

V1 RD22SFZ

## Package Label

NDC: 53329-031-01



## MAXIMUM STRENGTH

## TRIPLE ANTIBIOTIC + PAIN RELIEF

BACITRACIN ZINC/NEOMYCIN SULFATE/POLYMYXIN-B SULFATE/PRAMOXINE HCI FIRST AID ANTIBIOTIC/PAIN RELIEVING OINTMENT



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# NET WT. 1 OZ (28.4 g)

**REF** CUR001232

#### **Drug Facts**

# Active ingredients (in each gram) Bacitracin 500 units Neomycin 3.5 mg First aid antibiotic Pramoxine hydrochloride 10 mg Pain reliever Pain reliever

**Uses** ■ first aid to help prevent infection in and temporarily relieves pain due to ■ minor cuts ■ scrapes ■ burns

Warnings

■ For external use only

Do not use ■ if 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

#### Drug Facts (continued)

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 reactions 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

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Inactive ingredient petrolatum

2404982

#### **CURAD MAXIMUM STRENGTH TRIPLE ANTIBIOTIC**

bacitracin, neomycin, polymyxin b, pramoxine hydrochloride ointment

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53329-031
Floudti ivbe	HOMAN OTC DRUG	item code (Source)	NDC.33323-031

Route of Administration TOPICAL

#### **Active Ingredient/Active Moiety Basis of Strength Ingredient Name** Strength POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B -10000 [USP'U] POLYMYXIN B UNII: J2VZ 07J96K) in 1 g NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN -**NEOMYCIN SULFATE** 3.5 mg in 1 g UNII:116QD7X297) PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE -**PRAMOXINE** 10 mg in 1 g UNII:068X84E056) **HYDROCHLORIDE** 500 [USP'U] BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I) **BACITRACIN ZINC** in 1 g

Inactive Ingredients	
Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

#### **Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53329-031- 01	1 in 1 BOX	12/01/2019	
1		28.4 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	12/01/2019			

## Labeler - Medline Industries, LP (025460908)

Revised: 11/2024 Medline Industries, LP