

**NEOSPORIN PLUS PAIN RELIEF CLUBTRAY- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride
Johnson & Johnson Consumer Inc.**

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.

Neosporin Plus Pain Relief Clubtray

Neosporin Plus Pain Relief

Drug Facts

Active ingredients (in each gram)

Bacitracin Zinc (500 units)

Neomycin Sulfate (3.5 mg)

Polymyxin B Sulfate (10,000 units)

Pramoxine HCl (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 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 Centre right away.

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° to 25°C (68° to 77°F)

Inactive ingredients

Petrolatum

Questions?

call **800-223-0182** or **215-273-8755** (collect)

Distributed by:

JOHNSON & JOHNSON CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - Kit Package Label

#1 DOCTOR

RECOMMENDED

BRAND

OINTMENT

NEOSPORIN®

+

PAIN RELIEF

Maximum Strength Pain Relief

24-Hour Infection

Protection

Every Cut.

Every Time.®

Use with

BAND-AID®

Brand

Bandages

FIRST AID ANTIBIOTIC

PAIN RELIEVING

OINTMENT

Bacitracin Zinc-Neomycin

Sulfate-Polymyxin B

Sulfate-Pramoxine HCl

For Home

& On-the-Go!

NO STING

NEW WT 2.0 oz (56.7g)

CONTAINS:

1 Tube -

NEOSPORIN® + Pain Relief

Ointment 1 oz (28.3 g)

2 Tubes -

NEOSPORIN® + Pain Relief

Ointment 0.5 oz (14.2 g) each



NEOSPORIN PLUS PAIN RELIEF CLUBTRAY

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

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0801
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Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0801-9	1 in 1 PACKAGE	06/01/2022	

Quantity of Parts

Part #	Package Quantity	Total Product Quantity
Part 1	1 TUBE	14.2 g
Part 2	1 TUBE	28.3 g

Part 1 of 2

NEOSPORIN PLUS PAIN RELIEF

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

Product Information

Item Code (Source)	NDC:69968-0057
Route of Administration	TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ.07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
PRAMOXINE HYDROCHLORIDE (UNII: 88AYB867L5) (PRAMOXINE - UNII:068X84E056)	PRAMOXINE HYDROCHLORIDE	10 mg in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0057-2	1 in 1 CARTON		
1		14.2 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	12/01/2009	

Part 2 of 2

NEOSPORIN PLUS PAIN RELIEF

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

Product Information

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Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0057-1	1 in 1 CARTON		
1		28.3 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	12/01/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	06/01/2022	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 2/2023

Johnson & Johnson Consumer Inc.