NEOSPORIN PLUS BURN RELIEF- bacitracin zinc, neomycin sulfate, polymyxin b sulfate, and pramoxine hydrochloride ointment
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 ® + BURN RELIEF

Drug Facts

Active ingredients (in each gram)	***Purpose***
Bacitracin Zinc (500 units)	First aid antibiotic
Neomycin Sulfate (3.5 mg)	First aid antibiotic
Polymyxin B Sulfate (10,000 units)	First aid antibiotic
Pramoxine HCl (10 mg)	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 outside the US **215-273-8755** (collect)

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 14.2 g Tube Carton

#1 **DOCTOR**
RECOMMENDED
BRAND
NEOSPORIN ®
+
BURN RELIEF
Maximum Strength
Pain Relief to
Soothe Minor Burns

24-Hour Infection
Protection

Triple Antibiotic
Formula

NO STING

OINTMENT
FIRST AID ANTIBIOTIC/ PAIN RELIEVING OINTMENT

Bacitracin Zinc-Neomycin Sulfate-Polymyxin B Sulfate-
Pramoxine HCl
NET WT. 0.5 OZ (14.2 g)

NEOSPORIN®

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Maximum Strength Pain Relief to Soothe Minor Burns

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NET WT. 0.5 OZ (14.2 g)

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The makers of NEOSPORIN® do not make store brand products.

how2recycle.info

PAPER | PLASTIC TUBE BOX

Distributed by:
JOHNSON & JOHNSON CONSUMER INC.
Skillman, NJ 08558
 www.neosporin.com

Every Cut. Every Time.®
Use with BAND-AID® Brand Bandages

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Drug Facts (continued)

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Drug Facts (continued)

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EXP
LOT
CODE AREA

NEOSPORIN®
+ BURN RELIEF
OINTMENT

NEOSPORIN PLUS BURN RELIEF

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

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0518
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [iU] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	10000 [iU] 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:69968-0518-1	1 in 1 CARTON	02/01/2019	
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	02/01/2019	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 1/2023 Johnson & Johnson Consumer Inc.