

SHEFFIELD TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin zinc, polymyxin b sulfate, neomycin sulfate, pramoxine hydrochloride ointment
Sheffield Pharmaceuticals LLC

Drug Facts

Active ingredients (In each gram)

Bacitracin 500

Neomycin 3.5 mg

Polymyxin B 10,000 units

Pramoxine hydrochloride 10mg

Purpose

First aid antibiotic

First aid antibiotic

First aid antibiotic

Pain reliever

Uses

helps prevent infection in and temporarily relieves pain due to:

- cuts
- scrapes
- burns

Warnings

For external use only

Allergy alert:

- do not use if you are allergic to any of the ingredients

Do not use

- in or near the eyes
- on large areas of the body

Consult a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

When using this product

- do not use longer than 1 week

Stop use and consult a doctor if

- 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 ingredient

Petrolatum

Questions? 1-800-222-1087

Principal Display Panel - Carton Label

Dr. Sheffield's NDC 11527-161-55

Triple Antibiotic Plus Pain Relief

Bacitracin Zinc • Neomycin Sulfate • Polymyxin-B Sulfate • Pramoxine HCL

First Aid Antibiotic Plus Pain Relieving Ointment

NET WT. 1 OZ (28 g)



Compare to the active ingredients
in Neosporin®+ Pain Relief*

MULTI ACTION OINTMENT

- INFECTION PROTECTION FOR MINOR CUTS, SCRAPES & BURNS
- MAXIMUM STRENGTH PAIN RELIEF



NDC 11527-161-55

Maximum Strength Triple Antibiotic + Pain Relief

• Bacitracin Zinc • Neomycin Sulfate • Polymyxin-B Sulfate • Pramoxine HCl

FIRST AID ANTIBIOTIC / PAIN RELIEVING OINTMENT

NET WT. 1 OZ (28g)



Distributed by: Sheffield Pharmaceuticals
170 Broad Street, New London, CT 06320
www.sheffieldpharma.com
Questions? 1-800-222-1087

Made in USA
OF U.S. & IMPORTED
INGREDIENTS AND COMPONENTS



5442613

**learn more at www.sheffieldpharma.com

*This product is not manufactured or distributed by ©Johnson & Johnson Consumer Products Company - A Division of Johnson & Johnson Consumer Companies, Inc., owner of the registered trademark Neosporin®

Drug Facts	
Active ingredients (in each gram)	Bacitracin 500 units Neomycin 5 mg Polymyxin B 10,000 units Pramoxine hydrochloride 10 mg
Purpose	First aid antibiotic First aid analgesic Pain reliever
Uses	Helps prevent infection in and temporarily relieves pain due to • minor cuts • scrapes • burns
Warnings	For external use only Allergy alert: • do not use if allergic to any of the ingredients Do not use • 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 When using this product • do not use longer than 1 week Stop use and ask a doctor if • condition 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 occurs Keep out of the reach of children. If swallowed get medical help or contact a Poison Control Center immediately.
Directions	Adults and children 2 years and older • 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. Children 2 years and older, ask a doctor.
Other information	• Store at a controlled room temperature 68°-77° (20°-25°) Inactive ingredient: White Petroleum



Compare to the active ingredients
in Neosporin®+ Pain Relief*

MULTI ACTION OINTMENT

- INFECTION PROTECTION FOR MINOR CUTS, SCRAPES & BURNS
- MAXIMUM STRENGTH PAIN RELIEF



NDC 11527-161-55

Maximum Strength Triple Antibiotic + Pain Relief

• Bacitracin Zinc • Neomycin Sulfate • Polymyxin-B Sulfate • Pramoxine HCl

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Drug Facts	
Active Ingredients (in each gram)	Bacitracin 500 units Neomycin 3.5 mg Polymyxin B 10,000 units Pramoxine Hydrochloride 10 mg
Uses	Helps prevent infection in and temporarily relieves pain due to • minor cuts • scrapes • burns
Warnings	For external use only Allergy alert: • do not use if allergic to any of the ingredients Do not use • 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
Directions	When using this product • do not use longer than 1 week Stop use and ask a doctor if • a rash or other allergic reaction occurs Keep out of the reach of children. If swallowed get medical help or contact a Poison Control Center immediately.
Other information	Adults and children 2 years and older • 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. Children 2 years and older, ask a doctor
Store at a controlled room temperature 68°F-77°F (20°-25°C)	
Inactive ingredient	White Petroleum

Principal Display Panel - Tube Label

Dr. Sheffield's NDC 11527-161-55

Triple Antibiotic Plus Pain Relief

**Bacitracin Zinc • Neomycin Sulfate • Polymyxin-B Sulfate • Pramoxine HCL
First Aid Antibiotic/Pain Relieving Ointment**

NET WT. 1 OZ (28 g)

First Aid Treatment for Minor Cuts, Scrapes and Burns

Compare to the active ingredients
in Neosporin® + Pain Relief

Maximum Strength Triple Antibiotic + Pain Relief

• Bacitracin Zinc • Neomycin Sulfate • Polymyxin-B Sulfate • Pramoxine HCl
FIRST AID ANTIBIOTIC / PAIN RELIEVING OINTMENT

NET WT. 1 OZ (28g)



Active Ingredients (In each gram)

Bacitracin 500 units, Neomycin 3.5 mg, Polymyxin B 10,000 units First aid antibiotic
Pramoxine hydrochloride 10 mg Pain reliever

Uses helps prevent infection in and temporarily relieves pain due to • minor cuts • scrapes • burns **Warnings** For external use only.

Allergy alert: • do not use if allergic to any of the ingredients Do not use • 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 When using this product • do not use longer than 1 week **Stop use and ask a doctor if** • 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 occurs

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). **Questions?** 1-800-222-1087

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Distributed by: Sheffield Pharmaceuticals 170 Broad Street, New London, CT 06320 www.sheffieldpharma.com

Made in USA OF U.S. & Imported Ingredients and Components

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SHEFFIELD TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

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

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:11527-161
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:J2VZ 07J96K)	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
Petrolatum (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11527-161-55	1 in 1 CARTON	01/23/2011	
1		28 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	01/23/2011	

Labeler - Sheffield Pharmaceuticals LLC (151177797)**Registrant** - Sheffield Pharmaceuticals LLC (151177797)**Establishment**

Name	Address	ID/FEI	Business Operations
Sheffield Pharmaceuticals LLC		151177797	manufacture(11527-161)

Revised: 11/2023

Sheffield Pharmaceuticals LLC