

RAPIDOL TRIPLE ANTIBIOTIC- bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment
Pharmadel LLC

Rapidol Triple Antibiotic (KENIL)

Drug Facts

Active ingredients & Purpose

<i>Active ingredients</i>	<i>Purpose</i>
Bacitracin zinc 400 units	First aid antibiotic
Neomycin sulfate 3.5 mg.....	First aid antibiotic
Polymyxin B sulfate 5,000 units	First aid antibiotic

Uses

First aid to help prevent infection in minor cuts, scrapes, and burns.

Warnings

For external use only.

Do not use

- in the eyes
- over large areas of the body
- if you are allergic to any of the ingredients

Ask a doctor before use if you have

- deep or puncture wounds
- animal bites
- serious burns

Stop use and ask a doctor if

- condition persists or gets worse
- a rash or other allergic reaction develops
- you need to use longer than 1 week

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center immediately.

Directions

- clean the affected area
- apply a small amount of this product (an amount equal to the surface of the tip of a finger) on the area 1 to 3 times daily
- may be covered with a sterile bandage

Other information

- store between (68-77°F) 20-25°C
- don't use if foil seal under cap is punctured, torn, or missing

Inactive ingredients

cocoa butter, cottonseed oil, mineral oil, olive oil, paraffin, sodium pyruvate, tocopherol acetate (vit. E), white petrolatum

Principal Display Panel

NDC 55758-440-02



Triple Antibiótico

UNGUENTO DE PRIMEROS AUXILIOS

Bacitracina Zinc/Sulfato de Neomicina/Sulfato de Polimixina B

Peso Neto 2 oz (57g)



Triple Antibiotic / Antibiótico

EFFECTIVO
24- Horas
Protección
contra infecciones

EFFECTIVE
24-Hour
Infection
Protection



Triple Antibiotic

FIRST AID OINTMENT

Net Wt 2 oz (56g)

Bacitracin Zinc/ Neomycin Sulfate/ Polymyxin B Sulfate



RAPIDOL TRIPLE ANTIBIOTIC

bacitracin zinc, neomycin sulfate, polymyxin b sulfate ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55758-440
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [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	5000 [USP ^U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
PARAFFIN (UNII: I9O0E3H2ZE)	
COTTONSEED OIL (UNII: H3E878020N)	
COCOA BUTTER (UNII: 512OYT1CRR)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
OLIVE OIL (UNII: 6UYK2W1W1E)	
MINERAL OIL (UNII: T5L8T28FGP)	
SODIUM PYRUVATE (UNII: POD38AIF08)	
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Product Characteristics

Color	white (Translucent)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55758-440-02	1 in 1 CARTON	11/04/2024	
1		56 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	11/04/2024	

Labeler - Pharmadel LLC (030129680)

Revised: 10/2024

Pharmadel LLC