PENIMIXCINA- bacitracin, neomycin sulfate, polymixin b sulfate, pramoxine hydrochloride ointment OPMX LLC

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.

Penimixcina - First Aid Ointment

ACTIVE INGREDIENTS (IN EACH GRAM)

BACITRACIN ZINC 500 UNITS

NEOMYCIN SULFATE 5 MG (EQUIVALENT TO 3.5 MG NEOMYCIN)

POLYMYXIN B SULFATE 5000 UNITS

PRAMOXINE HYCHLORIDE 10 MG

PURPOSE

FIRST AID ANTIBIOTIC
PAIN RELIEVER

USES

- FIRST AID TO HELP PREVENT INFECTION IN
- MINOR CUTS
- SCRAPES
- BURNS

WARNINGS

For external use only.

DO NOT USE

- IN THE EYES
- IF YOU ARE ALLERGIC TO ANY OF THE INGREDIENTS
- OVER LARGE AREAS OF THE BODY
- LONGER THAN 1 WEEK UNLESS DIRECTED BY A DOCTOR

ASK A DOCTOR BEFORE USE IN CASE OF

- DEEP OR PUNCTURE WOUNDS
- ANIMAL BITES
- OR SERIOUS BURNS

Keep out of reach of children

• If swallowed get medical help or contact Poison Control Center right away.

DIRECTIONS

- 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

INACTIVE INGREDIENTS

WHITE PETROLATUM

OTHER INFORMATION

- STORE AT 15° 30°C (59°F 86°F)
- AVOID EXCESSIVE HEAT AND HUMIDITY

Penimixcina First Aid Ointment

NDC 69729-615-05



bacitracin, neomycin sulfate, polymixin b sulfate, pramoxine hydrochloride ointment

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69729-615	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RW052I)	BACITRACIN	500 [iU] in 1 g		
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:116QD7X297)	NEOMYCIN	5 mg in 1 g		
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII: J2VZ 07J96K)	POLYMYXIN B	5000 [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)		

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:69729-615- 05	1 in 1 BOX	09/05/2018			
1	NDC:69729-615- 01	14 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	09/05/2018		

Labeler - OPMX LLC (029918743)

Revised: 3/2022 OPMX LLC