PENTREXCILINA TRIPLE ANTIBIOTIC PLUS PAIN RELIEF- bacitracin, neomycin, polymyxin b, pramoxine ointment
OPMX LLC

Pentrexcilina Triple Antibiotic

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

Pentrexilina Triple Antibiotic

NDC 69729-616-61

PENTREXCILINA TRIPLE ANTIBIOTIC PLUS PAIN RELIEF

bacitracin, neomycin, polymyxin b, pramoxine ointment

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69729-616
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	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69729-616-61	1 in 1 BOX	09/05/2018	
1	NDC:69729-616-01	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 Drug	M004	09/05/2018	

Labeler - OPMX LLC (029918743)

Revised: 5/2024 OPMX LLC