

**TRIPLE ANTIBIOTIC PLUS LIDOCAINE- bacitracin zinc, neomycin sulfate ,
polymyxin b sulfate, lidocaine ointment
TRIFECTA PHARMACEUTICALS USA LLC**

Triple Antibiotic Ointment + Lidocaine

Active ingredients (each gram contains)

Bacitracin zinc 500 units
Neomycin sulfate 3.5 mg
Polymyxin B sulfate 10,000 units

Purpose

First aid antibiotic

Active Ingredient

Lidocaine 40mg

Purpose

External Analgesic

Uses

Uses first aid to help prevent infection and for the temporary relief of pain or discomfort in minor:● cuts ● scrapes ● burns

Keep out of reach of children

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Warnings

For external use only.

Ask Doctor Before use if you have

Ask Doctor before use if you have

- Deep or puncture wounds
- Animal bites
- serious burns

Do Not Use

Do Not Use

- If you are allergic to any of the ingredients
- In the eyes
- Over large areas of the body
- In large quantities, particularly over raw surfaces or blistered areas

Stop Use and Ask a doctor if

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- 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

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

Questions

Questions? Call 1-888-296-9067

Inactive Ingredient

Petrolatum

Other information

● store at 20° to 25°C (68° to 77°F)

* Lidocaine known in 30 to 60 minutes.

Distributed By

Distributed by:

Trifecta Pharmaceuticals USA®

101 NE Third Avenue, Suite 1500

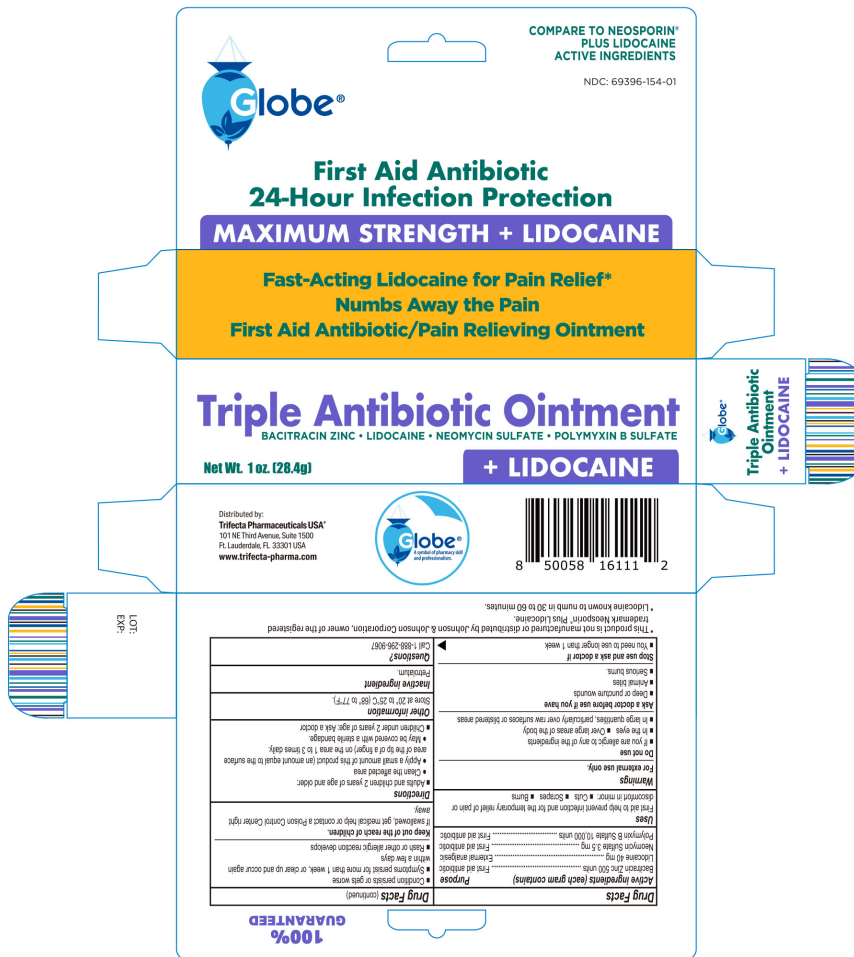
Ft. Lauderdale, FL 33301, USA

1-888-296-9067

* This product is not manufactured or distributed by Johnson & Johnson Corporation, owner of the registered trademark Neosporin® plus Lidocaine

Packaging

OUTSIDE BOX



INNER TUBE



Active ingredients (each gram contains)	Purpose
Diclofenac 250 mg	For relief of pain and inflammation
Lidocaine 40 mg	For relief of pain
Paracetamol 325 mg	For relief of pain and inflammation
Polysorb 10 200 mg	For relief of pain and inflammation

Use: Rub into the affected area 3-4 times a day for the temporary relief of pain or discomfort.

Warnings: Do not use if you are allergic to any of the ingredients, if you have a skin condition, if you are pregnant or breastfeeding, or if you are taking other painkillers.

Do not use if: You are allergic to any of the ingredients, if you have a skin condition, if you are pregnant or breastfeeding, or if you are taking other painkillers.

Ask a doctor before use if you have: A history of stomach ulcers, kidney or liver disease, or if you are taking other medicines.

Do not use for more than 1 week.

Other information: Contains 250 mg of paracetamol per gram.

Contains: Paracetamol, Lidocaine, Diclofenac.

Manufactured by: Takeda Pharmaceuticals USA, Inc. 101 NE Third Avenue, Suite 1500, Ft. Lauderdale, FL 33301 USA | www.takeda-pharma.com

Label

OUTSIDE BOX



TRIPLE ANTIBIOTIC PLUS LIDOCAINE

bacitracin zinc, neomycin sulfate, polymyxin b sulfate, lidocaine ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69396-154
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
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ 07J96K)	POLYMYXIN B	10000 [USP'U] in 1 g
LIDOCAINE (UNII: 98PI200987) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE	40 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:69396-154-01	1 in 1 BOX	03/31/2024	
1		28.4 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69396-154-22	2 in 1 BOX	06/26/2024	
2		28.4 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	03/31/2024	

Labeler - TRIFECTA PHARMACEUTICALS USA LLC (079424163)**Registrant** - Trifecta Pharmaceuticals USA (079424163)