TRIPLE ANTIBIOTIC- triple antibiotic ointment
SOLA Pharmaceuticals

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.

Triple Antibiotic Ointment

Active Ingerdients (in each gram)

Bacitracin Zinc USP 400 Units

Neomycin Sulfate USP 5mg (equivalent to 3.5mg of Neomycin)

Polymixin B Sulfate USP 5,000 Units

Purpose

First aid antibiotic

Uses

First aid to help prevent infection in minor:
- cuts
- scrapes
- burns

Warnings

For external use only

Do not use:
- If you are allergic to any of the ingredients
- In the eyes
- Over large areas of the body

Ask a doctor before use if you have
- Deep or puncture wounds
- Animal bites
- Serious burns

Stop use and ask a doctor if:
- You need to use longer than 1 week
- Condition persists or gets worse
- Rash or other allergic reaction developes

Keep out of reach of children.

If swallowed, get medical help or contact a 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
- Children under 2 years of age - Ask the doctor

Other Information

- Store at 20-25°C (68-77°F)
- Store away from heat

Inactive ingredients

White Petrolatum

Questions or comments?

Call 1-866-747-7365

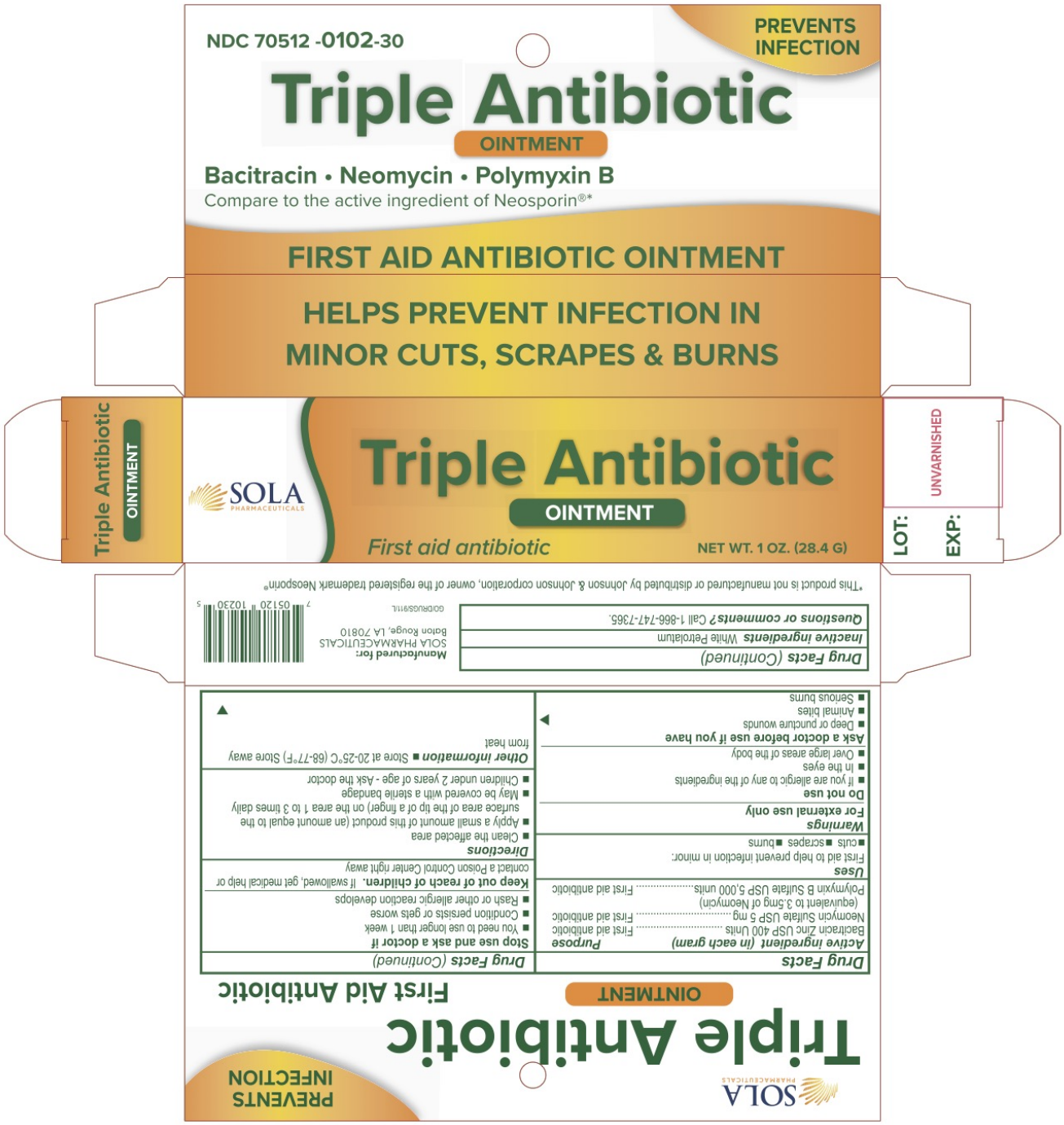
Manufactured for : SOLA Pharmaceuticals

Baton Rouge, LA 70810

Triple Antibiotic

NDC 70512-102-30

Qty: 28.4g



TRIPLE ANTIBIOTIC

triple antibiotic ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70512-102
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
		Basis of

Ingredient Name	Basis of Strength	Strength
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN SULFATE	5 mg in 1 g
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [USP'U] in 1 g

Inactive Ingredients

Ingredient Name	Strength
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70512-102-30	1 in 1 CARTON	02/12/2021	
1		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 final	part333B	02/12/2021	

Labeler - SOLA Pharmaceuticals (080121345)

Revised: 7/2022 SOLA Pharmaceuticals