LINCOLN- bacitracin zinc neomycin sulfate polymyxin b sulfate ointment
Lincoln Pharmaceutical Ltd.

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.

Drug Facts

Bacitracin Zinc 400 units- Neomycin Sulfate 5 mg (Equivalent to 3.5 mg Neomycin) Polymyxin B sulfate 5000 units

Drug Facts

First Aid antibiotic

Drug Facts

Uses

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

Drug Facts

For External use only

Drug Facts

- in the eyes
- if you are allergenic to any of the ingredients
- over the large areas of the body
- longer than 1 week unless directed by a doctor

Drug Facts

- deep or puncture wounds
- animal bites
- serious burns

Drug Facts

- the condition persists or get worse
- a rash or other allergic reaction develops

Drug Facts

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

Drug Facts

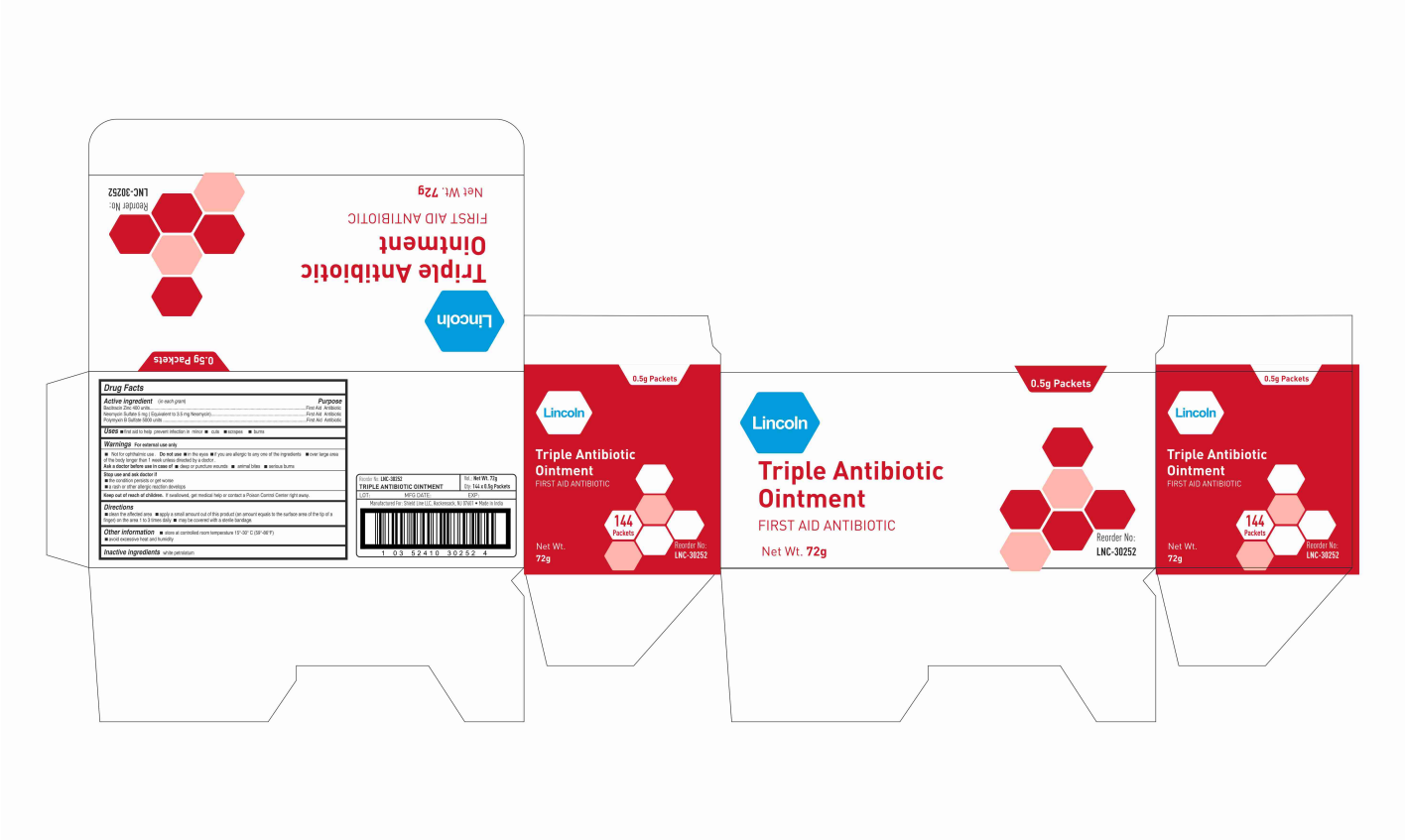
- clean the affected area

- 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 sterile bandage

- store at room temperature 15- 30° C (59-86F)
- avoid excessive heat and humidity

Drug Facts

white petrolatum



LINCOLN

bacitracin zinc neomycin sulfate polymyxin b sulfate ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69636-3025
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POLYMYXIN B SULFATE (UNII: 19371312D4) (POLYMYXIN B - UNII:J2VZ07J96K)	POLYMYXIN B	5000 [USP'U] in 1 g
NEOMYCIN SULFATE (UNII: 057Y626693) (NEOMYCIN - UNII:I16QD7X297)	NEOMYCIN	3.5 mg in 1 g
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	400 [USP'U] in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69636-3025-2	0.5 g in 1 PACKET; Type 0: Not a Combination Product	05/17/2013	
2	NDC:69636-3025-3	0.9 g in 1 PACKET; Type 0: Not a Combination Product	05/17/2013	
3	NDC:69636-3025-5	28.3 g in 1 TUBE; Type 0: Not a Combination Product	05/17/2013	
4	NDC:69636-3025-4	14 g in 1 TUBE; Type 0: Not a Combination Product	05/17/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	05/17/2013	

Labeler - Lincoln Pharmaceutical Ltd. (915839373)

Revised: 4/2016 Lincoln Pharmaceutical Ltd.