LINCOLN- bacitracin zinc ointment
Lincoln Pharmaceuticals 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

Active Ingredient (in each gram)

Bacitracin Zinc (equal to 500 bacitracin units)

Drug Facts

First Aid Antibiotic

Drug Facts

Uses

- first aid to help prevent infection in minor
- cuts scrapes
- burns

Drug Facts

For External use only

Do not use

- in the eyes
- if you are allergenic to any of the ingredients
- over large area 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

Keep out of reach of children

- 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 equals 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

Drug Facts

Other information:

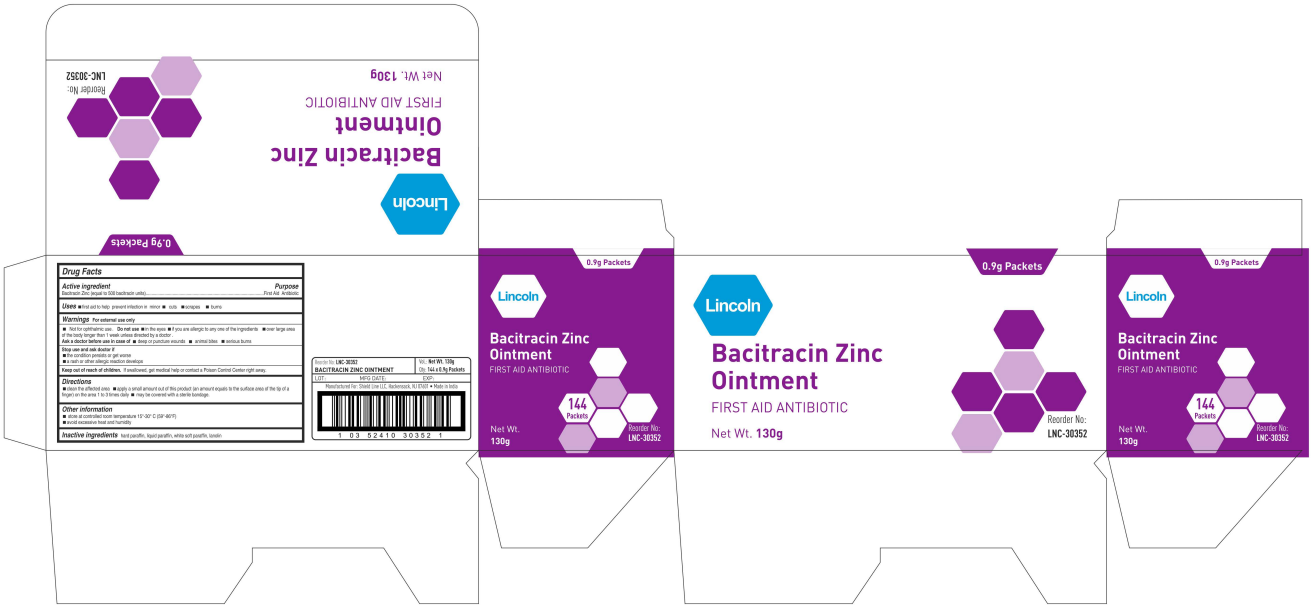
- store at controlled temperature 15°-30° C
- Avoid excessive heat and humidity

Drug Facts

Inactive ingredients

hard paraffin, liquid paraffin, white soft paraffin, lanolin

Principal Display Panel



LINCOLN

bacitracin zinc ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69636-3035
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

Inactive Ingredients	
Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
LANOLIN (UNII: 7EV65EAW6H)	
PETROLATUM (UNII: 4T6H12BN9U)	
PARAFFIN (UNII: I9O0E3H2ZE)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69636-3035-2	0.9 g in 1 PACKET; Type 0: Not a Combination Product	05/30/2013	
2	NDC:69636-3035-3	14 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2013	
3	NDC:69636-3035-4	28 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2013	
4	NDC:69636-3035-5	113 g in 1 TUBE; Type 0: Not a Combination Product	05/30/2013	
5	NDC:69636-3035-6	425 g in 1 JAR; Type 0: Not a Combination Product	05/30/2013	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333B	05/30/2013	

Labeler - Lincoln Pharmaceuticals Ltd. (915839373)

Establishment

Name	Address	ID/FEI	Business Operations
Lincoln Pharmaceuticals Ltd		915839373	manufacture(69636-3035)

Revised: 4/2016 Lincoln Pharmaceuticals Ltd.