LIVE BETTER BACITRACIN ZINC- bacitracin zinc ointment The Great Atlantic & Pacific Tea Company

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.

Live Better Bacitracin Zinc

Drug Facts

Active ingredient (each gram contains)

Bacitracin Zinc 500 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

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

Ask a doctor before use

• on deep or puncture wounds, animal bites, or serious burns

Stop use and ask a doctor if

- condition persists or gets worse
- a rash or other allergic reaction develops

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

Directions

- clean the affected area and dry thoroughly
- 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

Other information

- To open: unscrew cap, pull tab to remove foil seal
- store at room temperature
- see carton or tube crimp for lot number and expiration date

Inactive ingredients

mineral oil, white petrolatum

Questions?

Call **1-866-923-4914**

DISTRIBUTED BY ONPOINT, INC. 2 PARAGON DRIVE, MONTVALE, NJ 07645

PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

NDC 51143-075-02

Bacitracin Zinc Ointment USP

FIRST AID ANTIBIOTIC

NET WT 1 OZ (28.4g)



LIVE BETTER BACITRACIN ZINC

bacitracin zinc ointment

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51143-075	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
Bacitracin Zinc (UNII: 89 Y4M234ES) (Bacitracin - UNII:58 H6 RWO 52I)	Bacitracin	500 [iU] in 1 g		

Inactive Ingredients		
Ingredient Name	Strength	
MINERAL OIL (UNII: T5L8T28FGP)		
PETROLATUM (UNII: 4T6 H12BN9 U)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:51143-075-02	1 in 1 CARTON		
1	28.4 g in 1 TUBE		

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333B	05/11/2006		

Labeler - The Great Atlantic & Pacific Tea Company (001367366)

Registrant - Taro Pharmaceuticals U.S.A., Inc. (145186370)

Establishment				
Name	Address	ID/FEI	Business Operations	
Taro Pharmaceuticals Inc.		206263295	MANUFACTURE(51143-075)	

Revised: 3/2013 The Great Atlantic & Pacific Tea Company