

PUBLIX BACITRACIN ZINC- bacitracin zinc ointment
Publix Super Markets Inc

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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

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PRINCIPAL DISPLAY PANEL - 28.4 g Tube Carton

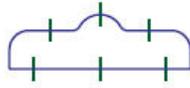
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bacitracinzinc
OINTMENT USP

FIRST AID ANTIBIOTIC OINTMENT

- **helps prevent infection in minor cuts, scrapes, and burns.**

NET WT 1 OZ (28.4 g)



bacitracin zinc

OINTMENT USP

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bacitracin zinc

OINTMENT USP



bacitracin zinc

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bacitracin
zinc
OINTMENT USP

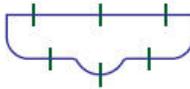
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OINTMENT USP

PUBLIX BACITRACIN ZINC

bacitracin zinc ointment

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:41415-051
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BACITRACIN ZINC (UNII: 89Y4M234ES) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [iU] in 1 g

Inactive Ingredients

Ingredient Name	Strength
MINERAL OIL (UNII: T5L8T28FGP)	
PETROLATUM (UNII: 4T6H12BN9U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:41415-051-75	1 in 1 CARTON	05/11/2006	
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 Drug	M004	05/11/2006	

Labeler - Publix Super Markets Inc (006922009)**Registrant** - Taro Pharmaceuticals U.S.A., Inc. (145186370)**Establishment**

Name	Address	ID/FEI	Business Operations
Taro Pharmaceuticals Inc.		206263295	manufacture(41415-051)

Revised: 2/2024

Publix Super Markets Inc