BACITRACIN- bacitracin ointment
Dynarex Corporation

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.

1161 Bacitracin NDC 67777-116-10
1162 Bacitracin NDC 67777-116-20
1163 Bacitracin NDC 67777-116-30

Active Ingredient

Bacitracin (500 units in each gram)

Purpose

First Aid Antibiotic

Use

First aid to help prevent infection in minor cuts, scrapes, and burns

Warnings

For External Use Only

Do not use

- In the eyes or apply 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 if you have

deep or puncture wounds, animal bites, or serious burns

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.

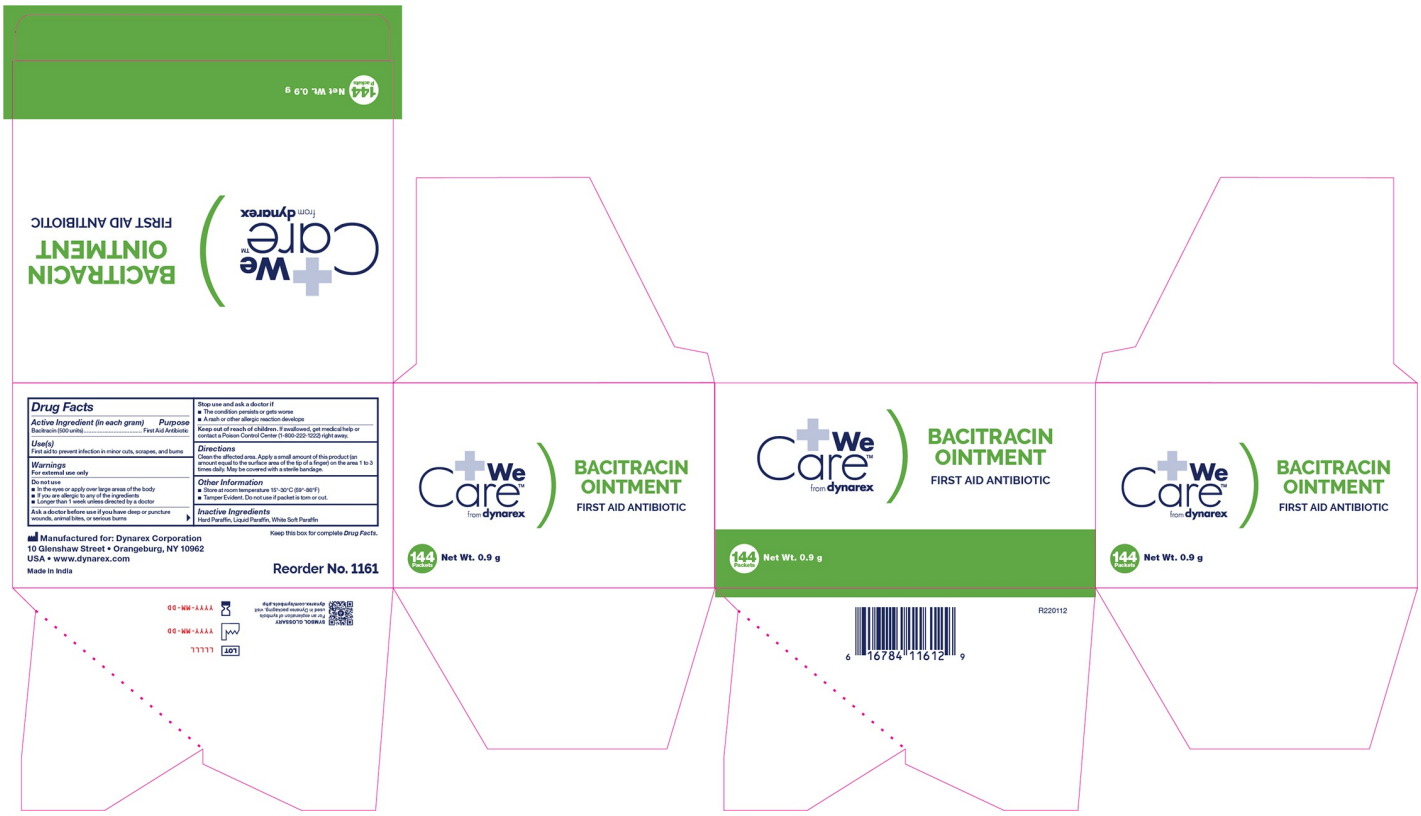
Other information

Store at room temperature

Inactive ingredients

Hard Paraffin, Liquid Paraffin, White Soft Paraffin

Label



1161 Bacitracin Ointment

Label

1162 Bacitracin Ointment

Label

1163 Bacitracin Ointment

BACITRACIN

bacitracin ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67777-116
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BACITRACIN (UNII: 58H6RWO52I) (BACITRACIN - UNII:58H6RWO52I)	BACITRACIN	500 [USP'U] in 1000 mg

Inactive Ingredients	
Ingredient Name	**Strength**
WHITE PETROLATUM (UNII: B6E5W8RQJ4)	
PARAFFIN (UNII: I9O0E3H2ZE)	
MINERAL OIL (UNII: T5L8T28FGP)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:67777-116-20	72 in 1 CASE	09/12/2016	
1	NDC:67777-116-21	14000 mg in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:67777-116-10	1728 in 1 CASE	09/12/2016	
2	NDC:67777-116-11	144 in 1 BOX		
2		900 mg in 1 PACKET; Type 0: Not a Combination Product		
3	NDC:67777-116-30	72 in 1 CASE	09/12/2016	
3	NDC:67777-116-31	1 in 1 BOX		
3		28400 mg 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	09/12/2016	

Labeler - Dynarex Corporation (008124539)

Revised: 11/2022

Dynarex Corporation