

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.

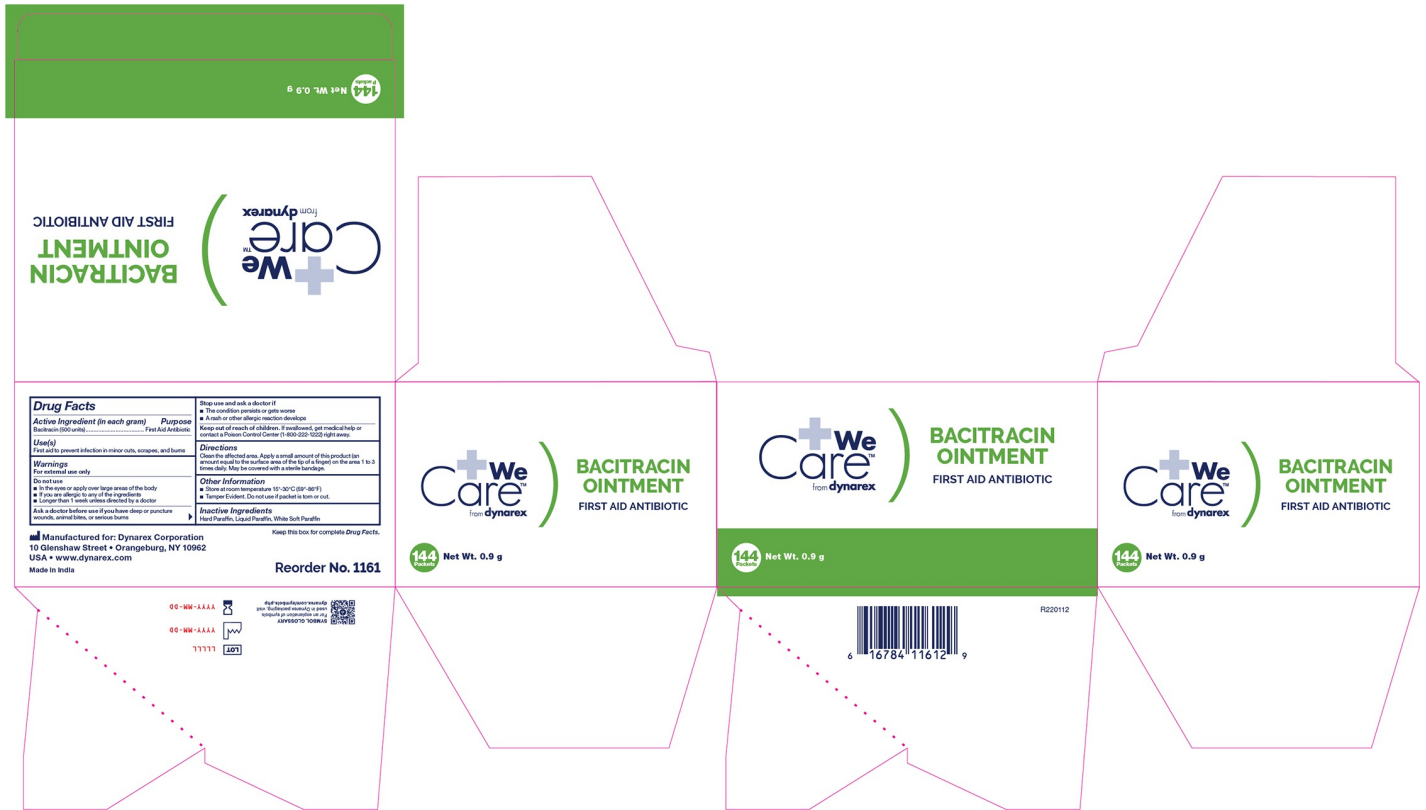
Other information

Store at room temperature

Inactive ingredients

Hard Paraffin, Liquid Paraffin, White Soft Paraffin

Label



1161 Bacitracin Ointment

Label

Drug Facts

Active Ingredient
Bacitracin (500 units in each gram).....**Purpose**
First Aid Antibiotic

Use(s)
■ First aid to help prevent infection in minor cuts, scrapes, and burns

Warnings For external use only

Do not use

- In the eyes or 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.

Stop use and ask a doctor if the condition persists or gets worse, or if a rash or other allergic reaction develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Drug Facts (continued)

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) onto the area 1 to 3 times daily. May be covered with a sterile bandage.

Other Information

- Store at room temperature 15°-30°C (59°-86°F)
- Tamper evident. Do not use if seal is damaged.

Inactive Ingredients

Hard Paraffin, Liquid Paraffin, White Soft Paraffin

**BACITRACIN
OINTMENT USP**
FIRST AID ANTIBIOTIC



YYYY-MM-DD
YYYY-MM-DD
LLLLL



BACITRACIN OINTMENT USP

FIRST AID ANTIBIOTIC

Net Wt. 0.5 oz. (14.2 g)



SYMBOL GLOSSARY
For an explanation of symbols used in Dynarex packaging, visit dynarex.com/symbols.php



6 16784 11622 8

Manufactured for:
Dynarex Corporation
10 Glenshaw Street
Orangeburg, NY 10962
USA • www.dynarex.com
Made In India R220126

Keep this box for complete **Drug Facts**

Reorder No. 1162

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: 58H6RW052I) (BACITRACIN - UNII:58H6RW052I) | | 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