**BLT 1- benzocaine ointment**
**CENTURA PHARMACEUTICALS INC**

---

## ACTIVE INGREDIENT

Benzocaine 20%

## PURPOSE

Topical Anesthetic

## USES

For the temporary relief of pain and itching.

## WARNINGS

• For external use only.
• Avoid contact with eyes or mucus membranes.
• Do not apply to open or damaged skin.
• If condition worsens or symptoms persist for more than seven days, discontinue use and consult physician.
• If pregnant or breast feeding,contact physician prior to use.
• Keep out of reach of children. If swallowed, contact Poison Control Center.
• Do not use if allergic to any ingredient in ointment.
• Do not use in large quantities, particularly over raw surfaces or blistered areas.

## DIRECTIONS

Adults and children two-years of age or older: Apply to affected area not more than three to four times daily. Children under two-years of age: consult a physician

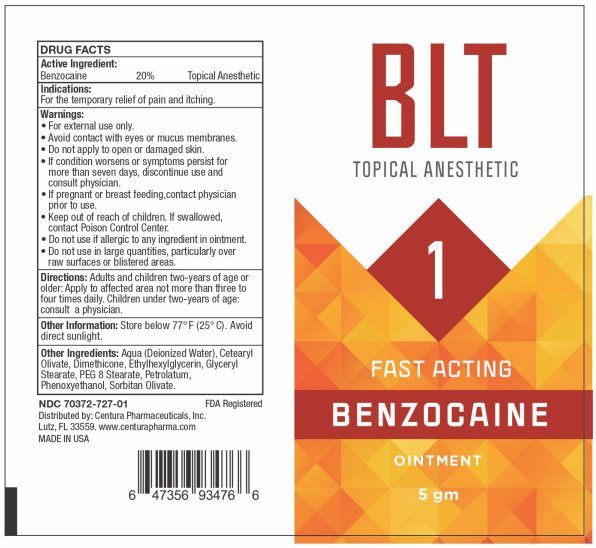
## OTHER INFORMATION

Store below 77° F (25° C). Avoid direct sunlight.

## INACTIVE INGREDIENTS

## KEEP OUT OF REACH OF CHILDREN

## PACKAGE LABELING




# BLT 1

benzocaine ointment

| Product Information | | | |
|---|---|---|---|
| **Product Type** | HUMAN OTC DRUG | **Item Code (Source)** | NDC:70372-727 |
| **Route of Administration** | TOPICAL | | |

| Active Ingredient/Active Moiety | | |
|---|---|---|
| **Ingredient Name** | **Basis of Strength** | **Strength** |
| **BENZOCAINE** (UNII: U3RSY48JW5) (BENZOCAINE - UNII:U3RSY48JW5) | BENZOCAINE | 20 g in 100 g |

| Inactive Ingredients | |
|---|---|
| **Ingredient Name** | **Strength** |

| | |
|---|---|
| **PEG-100 STEARATE** (UNII: YD01N1999R) | |
| **PHENOXYETHANOL** (UNII: HIE492ZZ3T) | |
| **DIMETHICONE** (UNII: 92RU3N3Y1O) | |
| **WATER** (UNII: 059QF0KO0R) | |
| **CETEARYL OLIVATE** (UNII: 58B69Q84JO) | |
| **ETHYLHEXYLGLYCERIN** (UNII: 147D247K3P) | |
| **GLYCERYL STEARATE SE** (UNII: FCZ5MH785I) | |
| **PEG-8 STEARATE** (UNII: 2P9L47VI5E) | |
| **PETROLATUM** (UNII: 4T6H12BN9U) | |
| **SORBITAN OLIVATE** (UNII: MDL271E3GR) | |

## Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|---|---|---|---|
| **1** | NDC:70372-727-01 | 5 g in 1 POUCH; Type 0: Not a Combination Product | 10/15/2016 | |

## Marketing Information

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|---|---|---|---|
| OTC Monograph Drug | M017 | 10/15/2016 | |

## Labeler - CENTURA PHARMACEUTICALS INC (084921637)

## Registrant - CENTURA PHARMACEUTICALS INC (084921637)

Revised: 1/2025 CENTURA PHARMACEUTICALS INC