

**BLT 2- lidocaine hcl ointment**  
**CENTURA PHARMACEUTICALS INC**

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**ACTIVE INGREDIENT**

Lidocaine HCL 4%

**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**

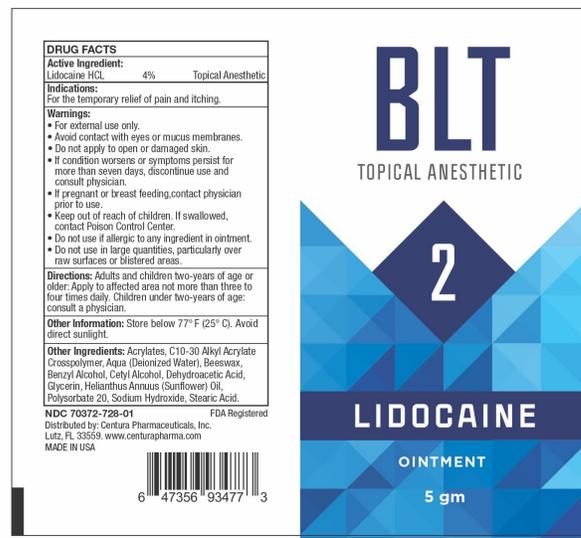
Store below 77° F (25° C). Avoid direct sunlight.

**INACTIVE INGREDIENTS**

Acrylates, C10-30 Alkyl Acrylate Crosspolymer, Aqua (Deionized Water), Beeswax, Benzyl Alcohol, Cetyl Alcohol, Dehydroacetic Acid, Glycerin, Helianthus Anuus (Sunflower) Oil, Polysorbate 20, Sodium Hydroxide, Stearic Acid.

## KEEP OUT OF REACH OF CHILDREN

## PACKAGE LABELING



## BLT 2

lidocaine hcl ointment

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:70372-728
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)		LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g	
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
GLYCERIN (UNII: PDC6A3C00X)				
SUNFLOWER OIL (UNII: 3W1JG795YI)				
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)				
BENZYL ALCOHOL (UNII: LKG8494WBH)				
POLYSORBATE 20 (UNII: 7T1F30V5YH)				
CETYL ALCOHOL (UNII: 936JST6JCN)				
YELLOW WAX (UNII: 2ZA36H0S2V)				
STEARIC ACID (UNII: 4ELV7Z65AP)				
DEHYDROACETIC ACID (UNII: 2KAG279R6R)				
SODIUM HYDROXIDE (UNII: 55X04QC32I)				
WATER (UNII: 059QF0K00R)				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:70372-728-01	5 g in 1 POUCH; Type 0: Not a Combination Product	11/21/2016	
<b>Marketing Information</b>				
<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>	
OTC Monograph Drug	M017	11/21/2016		

**Labeler** - CENTURA PHARMACEUTICALS INC (084921637)

**Registrant** - CENTURA PHARMACEUTICALS INC (084921637)

Revised: 1/2026

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