BLT 2- lidocaine hcl ointment
CENTURA PHARMACEUTICALS INC

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

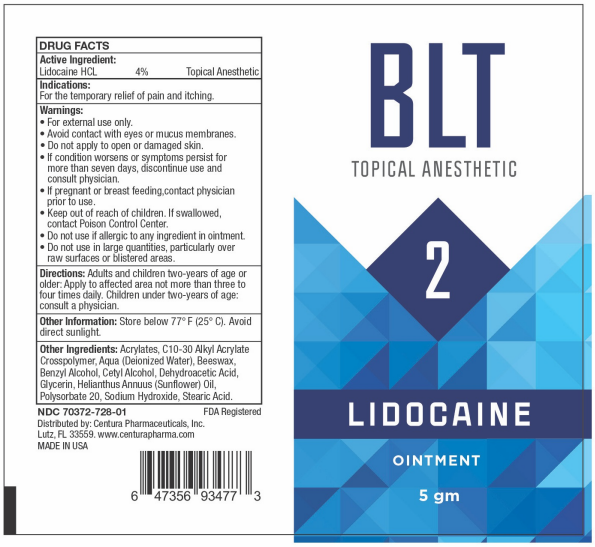
Store below 77° F (25° C). Avoid direct sunlight.

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			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70372-728
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	4 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
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: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70372-728-01	5 g in 1 POUCH; Type 0: Not a Combination Product	11/21/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/21/2016	

Labeler - CENTURA PHARMACEUTICALS INC (084921637)

Registrant - CENTURA PHARMACEUTICALS INC (084921637)

Revised: 1/2025 CENTURA PHARMACEUTICALS INC