LIDOZALL PLUS- lidocaine hydrochloride cream
V2 Pharma LLC

Lidozall Plus

Lidozall Plus

Rapid-onset topical analgesic

Lidozall Plus should be administered under the supervision of a licensed medical practitioner.

Active Ingredients

Lidocaine HCL 4.0% w/w

Purpose

External analgesic

Uses

For temporary relief of pain and itching due to minor skin irritations.

Warnings

- For external use only.
- Avoid contact with eyes.

- Do not use in large quantities, particularly over raw surfaces or blistered areas.

- Stop use and ask a doctor if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days. Discontinue use.

- Keep out of reach of children. If product is swallowed, get medical help or contact a Poison Control Center right away.

Directions

For adults and children two-years or older: Apply 1 packet (2 grams) topically to the affected area(s) up to 4 times daily as needed for pain. Do not use more than 4 packets (8 grams) per day.

Inactive Ingredients

Aqua (Deionized Water), Arnica Montana Floweri Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Oil,

Methylsulfonylmenthane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol, Stearic Acid, Triethanolamine.

Other Information

Protect this product from excessive heat and direct sun.

Product Label

Packaging for Lidozall Plus is shown below:

LIDOZALL PLUS

lidocaine hydrochloride cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72835-114
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
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
WATER (UNII: 059QF0KO0R)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
SODIUM CHONDROITIN SULFATE (PORCINE; 5500 MW) (UNII: H5BJH23Z9A)	
EMU OIL (UNII: 344821WD61)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
ISOPROPYL PALMITATE (UNII: 8CRQ2TH63M)	
MELALEUCA ALTERNIFOLIA LEAF (UNII: G43C57162K)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72835-114-15	15 in 1 CARTON	02/11/2025	
1		2 g in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/11/2025	

Labeler - V2 Pharma LLC (102457346)

Establishment

Name	Address	ID/FEI	Business Operations
V2 Pharma LLC		102457346	label(72835-114) , manufacture(72835-114) , pack(72835-114)