

LIDOZALL- lidocaine hydrochloride cream
V2 Pharma LLC

Lidozall

Lidozall

Rapid-onset topical analgesic

Lidozall 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.

<p>72835-004-01</p> <h1>LIDOZALL</h1> <p>Lidocaine 4% Cream (2 grams)</p>	<h2>Drug Facts</h2> <table border="1"><thead><tr><th>Active Ingredients</th><th>Purpose</th></tr></thead><tbody><tr><td>Lidocaine 4.0%</td><td>External Analgesic</td></tr></tbody></table> <p>Uses For temporary relief of pain and itching.</p> <p>Warnings For external use only. Do not use on wounds or damaged skin, in large quantities, or if you are allergic to any ingredients of this product. When using this product use only as directed. Avoid contact with the eyes, rashes, or mucous membranes. Stop use and ask doctor if condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days. Keep out of reach of children and pets If swallowed get medical help or contact a Poison Control Center right away.</p> <p>Directions Adults and children 12 years of age and over: Clean and dry affected area, apply to affected area not more than 3 to 4 times daily. Children 12 years of age or younger: ask a doctor.</p> <p>Other Information Protect this product from excessive heat and direct sun.</p> <p>Inactive Ingredients Aqua (Deionized Water), Arnica Montana Flower Extract, C13-14 Isoparaffin, Chondroitin Sulfate, Emu Oil, Ethoxydiglycol, Ethylhexylglycerin, Glucosamine Sulfate, Isopropyl Palmitate, Laureth-7, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methylsulfonylmethane (MSM), Phenoxyethanol, Polyacrylamide, Propylene Glycol, Stearic Acid, Triethanolamine</p> <p> 3 72835 00401 1</p> <p>Manufactured for: V2 Pharma, LLC Hudson, MI 49247 www.V2Pharma.com +1 800.519.8334</p>	Active Ingredients	Purpose	Lidocaine 4.0%	External Analgesic
Active Ingredients	Purpose				
Lidocaine 4.0%	External Analgesic				
<p>Rapid-onset topical analgesic</p>					
<p>Lidozall should be administered under the supervision of a licensed medical practitioner.</p> <p>V2 PHARMA</p>					



Lidozall Box of 20 Count, 2 Gram Packets
NDC 72835-004-20



LIDOZALL

lidocaine hydrochloride cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72835-004
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (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: A1A18X02B)	
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:72835-004-20	20 in 1 CARTON	12/16/2024	
1	NDC:72835-004-01	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	11/21/2024	

Labeler - V2 Pharma LLC (102457346)

Establishment

Name	Address	ID/FEI	Business Operations
Southeast Holdings Corp		080504027	manufacture(72835-004)

Revised: 11/2025

V2 Pharma LLC