

LIDOGUARD- lidocaine hcl cream
ALAINA HEALTHCARE PRIVATE LIMITED

4% Lidocaine HCl Cream

Drug Facts

Active Ingredient

Purpose

Lidocaine HCl 4%.....Topical Analgesic

Uses

Temporarily relieves pain and itching due to:

- minor cuts
- sunburn
- minor scrapes
- minor burns
- insect bites
- minor skin irritations

Warnings

For external use only.

When using this product

- do not use in or near the eyes
- do not use in large quantities, particularly over raw surfaces or blistered areas

Stop use and ask a doctor if

- Allergic reaction occurs
- condition worsens or does not improve within 7 days
- symptoms clear up and return within a few days
- redness, irritation, swelling, pain or other symptoms begin or increase

Do not apply over large areas of the body

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center.

Directions

- Adults and children 2 years and older: Apply externally to the affected area up to 3-4 times daily.
- Children under 2 years: Consult a doctor.

Other information

- Store at 20° to 25° C (68° to 77° F); excursions permitted to 15° to 30° C (59° to 86° F)

Inactive Ingredients

acrylates/c10-30 alkyl acrylate crosspolymer, aloe barbadensis leaf juice, aminomethyl propanol, butylene glycol, cetearyl alcohol, ceteth-10 phosphate, dicetyl phosphate, dimethicone, disodium EDTA, ethylhexylglycerin, glyceryl stearate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, phenoxyethanol, steareth-21, water

Questions?

Call

PRINCIPAL DISPLAY PANEL

Pain Relief
4% Lidocaine HCl Cream
Pain Relieving Cream
Topical Analgesic
1.05 oz (30g)

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LIDOGUARD

lidocaine hcl cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73492-272
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
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
ALOE VERA LEAF JUICE (UNII: RUE8E6T4NB)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
CETETH-10 PHOSPHATE (UNII: 4E05O5N49G)	
DIHEXADECYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE, UNSPECIFIED (UNII: 92RU3N3Y10)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
STARETH-21 (UNII: 53J3F32P58)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73492-272-30	1 in 1 CARTON	05/12/2025	
1		30 g in 1 TUBE; 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	05/12/2025	

Labeler - ALAINA HEALTHCARE PRIVATE LIMITED (858720927)

Establishment

Name	Address	ID/FEI	Business Operations
ALAINA HEALTHCARE PRIVATE LIMITED		858720927	manufacture(73492-272)

Revised: 12/2025

ALAINA HEALTHCARE PRIVATE LIMITED