

LEADER LIDOCAINE- lidocaine hcl liquid

CARDINAL HEALTH

Leader Lidocaine HCL 4% Roll On

Lidocaine HCl 4%

Topical analgesic

temporarily relieves pain and itching due to:

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

For external use only

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

When using this product do not get into eyes

Stop use and ask a doctor if

- condition gets worse
- symptoms last for more than 7 days
- symptoms clear up and occur again within a few days

Flammable •Keep away from fire or flame

If pregnant or breast-feeding, ask a health professional before use

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

Directions

- adults and children 12 years and over: apply to affected area not more than 3 to 4 times daily
- children under 12 years: ask a doctor

Other information

store between 20 °C and 25 °C (68 °F and 77 °F)

acrylates/C10-30 alkyl acrylate crosspolymer, aloe barbadensis (aloe vera) leaf juice, aminomethyl propanol, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, dicetyl phosphate, dimethicone, edetate disodium, ethylhexylglycerin, glyceryl monostearate, methylparaben, purified water, SD alcohol 40, steareth-21

Leader

NDC 70000-0099-1

Maximum Strength

Lidocaine

Pain Relief

Liquid

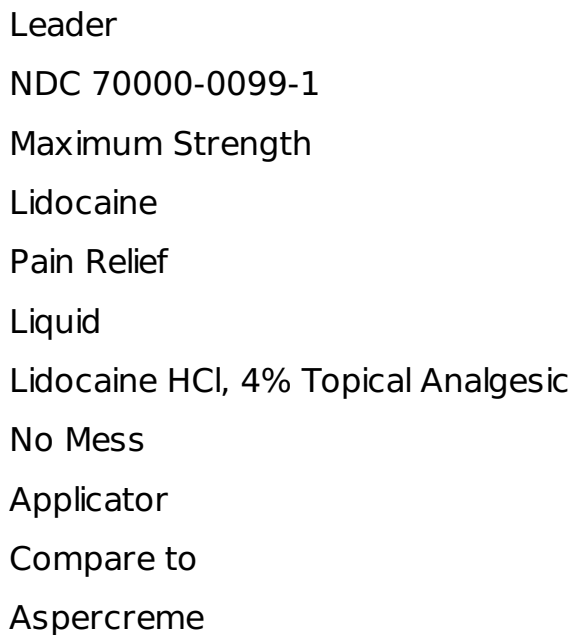
Lidocaine HCl, 4% Topical Analgesic

No Mess

Applicator

Compare to

Aspercreme



Lidocaine liquid

active ingredient

100% Money Back Guarantee

2.5 Fl oz (73 mL)

LEADER LIDOCAINE

lidocaine hcl liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70000-0099
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 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
STEARETH-21 (UNII: 53J3F32P58)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
DICETYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
WATER (UNII: 059QF0KO0R)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70000-0099-1	1 in 1 CARTON	10/20/2021	
1		73 mL in 1 BOTTLE, PLASTIC; 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	10/20/2021	

Labeler - CARDINAL HEALTH (063997360)

Registrant - Weeks & Leo Co., Inc. (005290028)

Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Co., Inc.		005290028	manufacture(70000-0099)

Revised: 12/2025

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