WESTLAKE LIDOCAINE- lidocaine hcl cream
Westlake Pharma LLC

Westlake Lidocaine HCL 4% Cream

Lidocaine HCl 4%

Topical analgesic

Uses

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

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

Directions

- adults and children 2 years and over: apply to affected area not more than 3 to 4 times daily
- children under 2 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

Maximum Strength

Lidocaine

Pain Relief

Cream

Lidocaine HCl 4% Topical Analgesic

Non Greasy

Compare to aspercreme lidocaine cream

active ingredient

100% Money Back Guarantee

Net wt 1.75 oz (49 g)

WESTLAKE LIDOCAINE

lidocaine hcl cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:82895-717
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)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
WATER (UNII: 059QF0KO0R)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
DICETYL PHOSPHATE (UNII: 2V6E5WN99N)	
DIMETHICONE 350 (UNII: 2Y53S6ATLU)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	
STEARETH-21 (UNII: 53J3F32P58)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82895-717-50	1 in 1 CARTON	10/01/2023	
1		49 g 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	10/01/2023	

Labeler - Westlake Pharma LLC (081176855)

Registrant - Weeks & Leo Company Inc (005290028)

Establishment

Name	Address	ID/FEI	Business Operations
Weeks & Leo Company Inc		005290028	manufacture(82895-717)

Revised: 12/2025

Westlake Pharma LLC