TOPCARE PAIN RELIEVING CREAM- lidocaine hcl 4% cream
Topco Associates, LLC

TopCare Lidocaine Cream

Lidocaine HCl 4%

temporarily relieves minor pain

Topical analgesic

For external use only

In large quantities, particularly over raw surfaces or blistered areas

Stop use

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

do not get into eyes

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

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: do not use, consult a doctor.

Other information

store at controlled room temperature 20°-25°C (68°-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

TopCare Health

Fragrance-free

Pain Relieving Cream

with Lidocaine HCl 4% Topical Analgesic

Maximum Strength

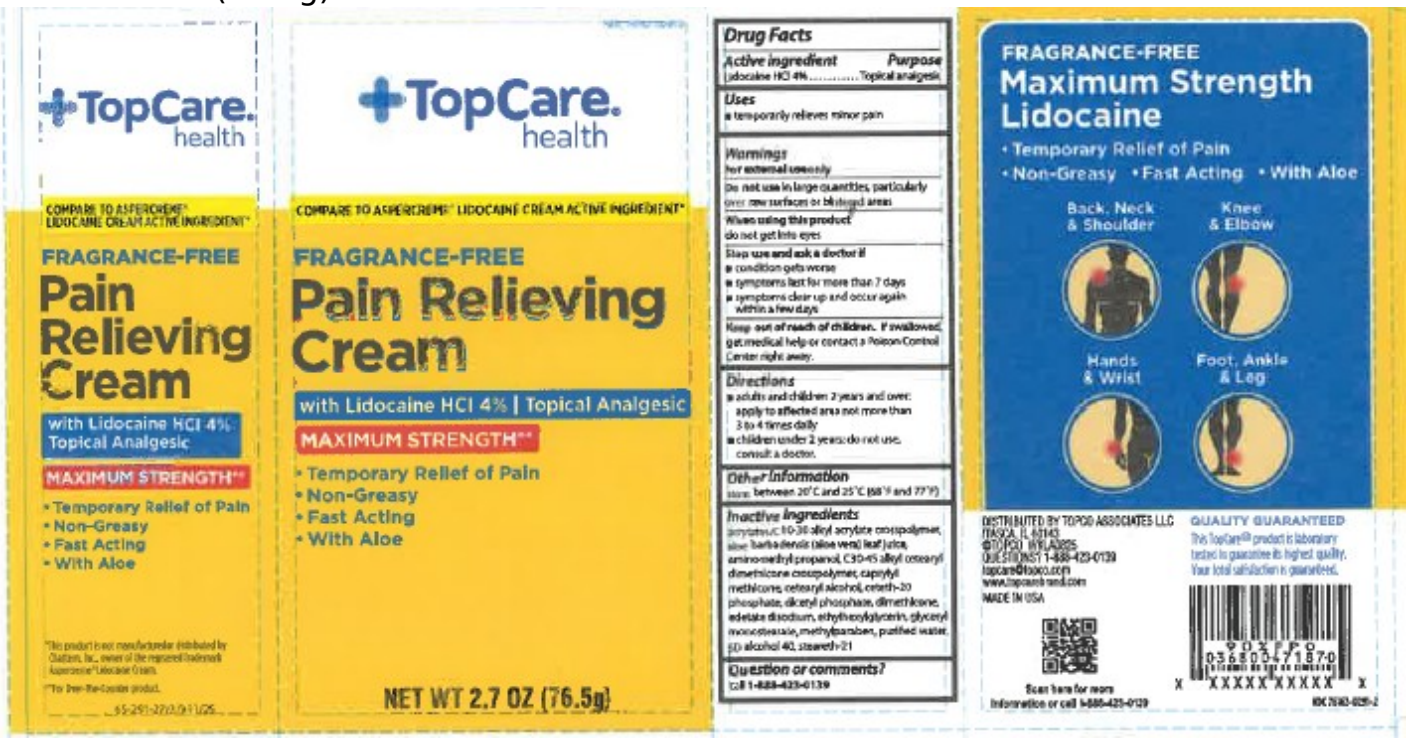
Temporary Relief of Pain

Non-Greasy

Fast Acting

with Aloe

Net Wt 2.7oz (76.5g)



TOPCARE PAIN RELIEVING CREAM

lidocaine hcl 4% cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:76162-291
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 g

Inactive Ingredients	
Ingredient Name	**Strength**
METHYLPARABEN (UNII: A2I8C7HI9T)	
AMINOMETHYL PROPANOL (UNII: LU49E6626Q)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
C30-45 ALKYL CETEARYL DIMETHICONE CROSSPOLYMER (UNII: 4ZK9VP326R)	
DICETYL PHOSPHATE (UNII: 2V6E5WN99N)	

ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
STEARETH-21 (UNII: 53J3F32P58)	
ALCOHOL (UNII: 3K9958V90M)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
WATER (UNII: 059QF0KO0R)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CAPRYLYL METHICONE (UNII: Q95M2P1KJL)	
ACRYLATES/C10-30 ALKYL ACRYLATE CROSSPOLYMER (60000 MPA.S) (UNII: 8Z5ZAL5H3V)	
CETETH-20 PHOSPHATE (UNII: 921FTA1500)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76162-291-02	1 in 1 CARTON	01/30/2026	
1		76.5 g in 1 BOTTLE; 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	01/30/2026	

Labeler - Topco Associates, LLC (006935977)

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

Establishment

Name	Address	ID/FEI	Business Operations
WEEKS & LEO COMPANY, INC.		005290028	manufacture(76162-291)

Revised: 1/2026 Topco Associates, LLC