

**MENTHOLATUM LIDOCAINE ICE- lidocaine hcl, menthol gel**  
**The Mentholatum Company**

-----  
**Drug Facts - Mentholatum Lidocaine Ice**

**Active ingredients**

Lidocaine HCl 4%

Menthol 1%

**Purpose**

Lidocaine HCl - Topical anesthetic

Menthol - Topical analgesic

**Uses**

temporarily relieves pain

**Warnings**

**For external use only**

**Do not use**

- on large areas of the body or on cut, irritated or blistered skin
- on puncture wounds
- for more than one week without consulting a doctor

**When using this product**

- use only as directed
- do not get into eyes
- do not bandage tightly or apply external heat to the area of use

**Stop use and ask a doctor if**

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days
- redness, rash, or irritation occurs

**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 a thin layer to affected area every 6 to 8 hours, not to exceed 3 applications in a 24 hour period
- children under 12 years: ask a doctor

**Inactive ingredients**

acrylates/C10-30 alkyl acrylate crosspolymer, alcohol, aloe barbadensis leaf juice, aminomethyl propanol, bis-vinyl dimethicone/dimethicone copolymer, C30-45 alkyl cetearyl dimethicone crosspolymer, caprylyl methicone, cetearyl alcohol, ceteth-20 phosphate, chlorphenesin, dicetyl phosphate, dimethicone, edetate disodium, ethoxydiglycol, glycerin, glyceryl monostearate, methoxypropanediol, phenoxyethanol, purified water, steareth-21

**Package/Label Principal Display Panel**



## MENTHOLATUM LIDOCAINE ICE

lidocaine hcl, menthol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:10742-1389
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

<b>LIDOCAINE HYDROCHLORIDE</b> (UNII: V13007Z41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE ANHYDROUS	40 mg in 1 mL
<b>MENTHOL, UNSPECIFIED FORM</b> (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	10 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED)</b> (UNII: 59TL3WG5CO)	
<b>ALCOHOL</b> (UNII: 3K9958V90M)	
<b>ALOE VERA LEAF</b> (UNII: ZY81Z83H0X)	
<b>AMINOMETHYLPROPANOL</b> (UNII: LU49E6626Q)	
<b>DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE)</b> (UNII: 9E4CO0W6C5)	
<b>CAPRYLYL TRISILOXANE</b> (UNII: Q95M2P1KJL)	
<b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)	
<b>CETETH-20 PHOSPHATE</b> (UNII: 921FTA1500)	
<b>CHLORPHENESIN</b> (UNII: I670DAL4SZ)	
<b>DIHEXADECYL PHOSPHATE</b> (UNII: 2V6E5WN99N)	
<b>DIMETHICONE</b> (UNII: 92RU3N3Y1O)	
<b>EDETATE DISODIUM</b> (UNII: 7FLD91C86K)	
<b>DIETHYLENE GLYCOL MONOETHYL ETHER</b> (UNII: A1A18X02B)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>GLYCERYL MONOSTEARATE</b> (UNII: 230OU9XXE4)	
<b>GLYCERIN METHYL ETHER</b> (UNII: 42ESM1DR47)	
<b>PHENOXYETHANOL</b> (UNII: HIE492ZZ3T)	
<b>POLYSORBATE 20</b> (UNII: 7T1F30V5YH)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>STARETH-21</b> (UNII: 53J3F32P58)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-1389-1	1 in 1 CARTON	08/01/2018	
1		80 mL 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	08/01/2018	

**Labeler** - The Mentholatum Company (002105757)

**Registrant** - The Mentholatum Company (002105757)

## Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-1389)

Revised: 12/2024

The Mentholatum Company