

DG HEALTH MAXIMUM STRENGTH COLD N HOT- lidocaine hcl, menthol cream DOLGENCORP, LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Drug Facts

Active ingredients

Lidocaine HCl 4%

Menthol 1%

Purpose

Lidocaine HCl - Topical anesthetic

Menthol – Topical analgesic

Uses

temporarily relieves minor pain

Warnings

For external use only

Do not use

- on large areas of the body or on cut, irritated, swollen, 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 (such as a heating pad) 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, 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, glycerin, glyceryl stearate SE, phenoxyethanol, purified water, SD alcohol 40, steareth-21

Package/Label Principal Display Panel



DG HEALTH MAXIMUM STRENGTH COLD N HOT

lidocaine hcl, menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55910-760
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
LIDOCAINE HYDROCHLORIDE (UNII: V13007Z 41A) (LIDOCAINE - UNII:98PI200987)	LIDOCAINE HYDROCHLORIDE	40 mg in 1 g

LEVOMENTHOL (UNII: BZ1R15MTK7) (LEVOMENTHOL - UNII:BZ1R15MTK7)			LEVOMENTHOL	10 mg in 1 g
Inactive Ingredients				
Ingredient Name				Strength
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)				
ALOE VERA LEAF (UNII: ZY81Z83H0X)				
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)				
DIMETHICONE/VINYL DIMETHICONE CROSSPOLYMER (SOFT PARTICLE) (UNII: 9E4CO0W6C5)				
C30-45 ALKYL METHICONE (UNII: NFX970DSI2)				
CAPRYLYL TRISILOXANE (UNII: Q95M2P1KJL)				
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)				
CETETH-20 PHOSPHATE (UNII: 921FTA1500)				
CHLORPHENESIN (UNII: I670DAL4SZ)				
DIHEXADECYL PHOSPHATE (UNII: 2V6E5VN99N)				
DIMETHICONE (UNII: 92RU3N3Y1O)				
EDETATE DISODIUM (UNII: 7FLD91C86K)				
GLYCERIN (UNII: PDC6A3C0OX)				
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)				
PHENOXYETHANOL (UNII: HIE492ZZ3T)				
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				
STEARETH-21 (UNII: 53J3F32P58)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55910-760-01	1 in 1 CARTON	03/30/2020	
1		49.6 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 not final		part348	03/30/2020	

Labeler - DOLGENCORP, LLC (068331990)