### CURETECH COLD ICE ANALGESIC GEL- menthol gel Curetech Skincare

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.

-----

#### **Curetech Cold Ice Gel**

# **Active Ingredient**

Menthol 2.0%

# **Purpose**

**Topical Analgesic** 

#### Uses:

- for temporary relief of minor aches and pains in muscles and joints associated with:
- simple backaches,
- strains
- sprains
- sports injuries
- arthritis
- bruises

# Warnings

### FOR EXTERNAL USE ONLY

Do not use:

- with other topical pain relievers
- with heating pads or heating devices

# When using this product

- do not use in or near eyes
- do not apply to wounds or damaged skin
- do not bandage tightly

# Stop use and ask a doctor if

- condition worsens
- symptoms last more than 7 days or clear up and occur again within a few days
- redness or irritation develops

### If pregnant or breastfeeding

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**

- clean affected area before applying product
- adults and children 2 years of age and older; apply to affected area not more than 3 to 4 times daily

### **Inactive Ingredients**

ISOPROPYL ALCOHOL (UNII: ND2M416302)

ammonium hydroxide, carbomer, cupric sulfate, FD&C Blue No. 1, isopropyl alcohol, magnesium sulfate, sodium hydroxide, thymol, water



# **CURETECH COLD ICE ANALGESIC GEL** menthol gel **Product Information Product Type** HUMAN OTC DRUG Item Code (Source) NDC:73622-3021 **Route of Administration TOPICAL Active Ingredient/Active Moiety Ingredient Name Basis of Strength** Strength MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A) **MENTHOL** 4.54 g in 227 g **Inactive Ingredients** Strength **Ingredient Name**

CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
CUPRIC SULFATE (UNII: LRX7AJ16DT)	
THYMOL (UNII: 3J50XA376E)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

l	P	Packaging						
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
	1	NDC:73622- 3021-2	227 g in 1 JAR; Type 0: Not a Combination Product	04/08/2019				

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	04/08/2019	

# Labeler - Curetech Skincare (677682180)

Revised: 9/2021 Curetech Skincare