#### ICE COLD ANALGESIC- menthol gel China Ningbo Shangge Cosmetic Technology Corp.

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.

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## Active Ingredient Purpose

Menthol 1.0% ..... Topical Analgesic

**Uses** temporarily relieves:

• minor muscle aches and pains

**Except out of reach of children.** Eln case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

DT for your DailyTouch

Ice Cold Analgesic Gel

Gel Frio Antimuscular

#### **Directions**

For the temporary relief of minor muscle aches and pains.

See important warnings under "When using this product"

- not for use on children under 2 years of age
- adults & children 2 years & older: apply liberally to painful area and massage until gel is absorbed into the skin. Repeat 3 to 4 times daily.

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## **Warnings**

For external use only[]; avoid contact with eyes.

Ask a doctor before use if you have cough associated with

- smoking
- excessive phlegm
- asthma
- emphysema
- persistent or chronic cough

## When using this product, do not:

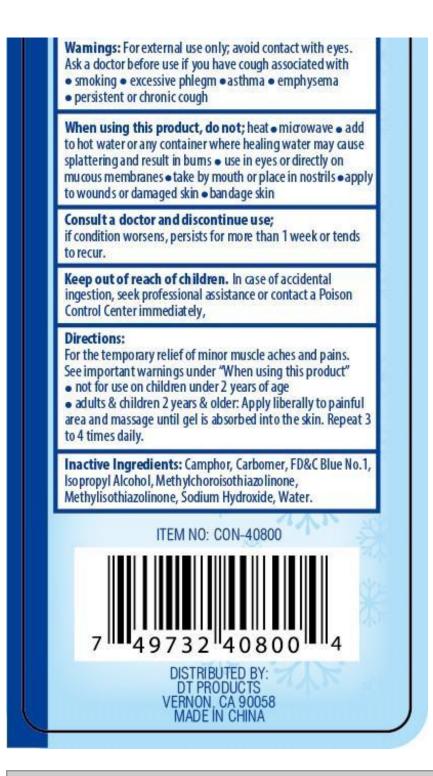
- heat
- microwave
- add to hot water or any container where healing water may cause splattering and result in burns
- use in eyes or directly on mucous membranes
- take my mouth or place in nostrils
- apply to wounds or damaged skin
- bandage skin

**Consult a doctor and discontinue use:** If condition worsens, persists for more than 1 week or tends to recur.

## **Inactive Ingredients**

camphor, carbomer, FD&C blue no.1, isopropyl alcohol, methylchloroisothiazolinone, methylisothiazolinone, sodium hydroxide, water.





# ICE COLD ANALGESIC

 menthol gel

 Product Information

 Product Type
 HUMAN OTC DRUG

 Route of Administration
 TOPICAL

 Active Ingredient/Active Moiety

 Menthol L UNIELTT10EIP3A) (MENTHOL - UNIELTT10EIP3A)

Inactive Ingredients							
	Strength						
CAMPHOR (NATURA	CAMPHOR (NATURAL) (UNII: N20HL7Q941)						
CARBOMER COPOLYMER TYPE A (UNII: 71DD5V995L)							
FD&C BLUE NO.1 (U	FD&C BLUE NO.1 (UNII: H3R47K3TBD)						
ISOPROPYL ALCOHOL (UNII: ND2M416302)							
METHYLCHLORO ISOTHIAZOLINONE (UNII: DEL7T5QRPN)							
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)							
SODIUM HYDROXIDE (UNII: 55X04QC32I)							
WATER (UNII: 059QF0KO0R)							
Packaging							
# Item Code	Package Description	Marketing Start Date	Marketing End Date				
1 NDC:58503-087-01	227 g in 1 PACKAGE; Type 0: Not a Combination Produc	et					
Marketing Information							
Marketing Catego	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date				
OTC monograph not fi	nal part348	07/10/2015					

Labeler - China Ningbo Shangge Cosmetic Technology Corp. (529287434)

Establishment					
Name	Address	ID/FEI	<b>Business Operations</b>		
Volume Distributors, Inc.		002029544	manufacture(58503-087)		

Revised: 7/2015

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China Ningbo Shangge Cosmetic Technology Corp.