ARCTIC ICE ANALGESIC- menthol gel
Blue Cross Labs

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.

Arctic Ice Analgesic Gel

Active Ingredient: Purpose

Menthol, 1% ..... Topical analgesic

Uses: temporarily relieves:

- minor muscle aches and pains

Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately

Consult a doctor and discontinue use;

if conditon worsens, persists for more than 1 week or tends to recur

Warnings: For external use only; avoid contact with eyes. Ask a doctor before use if you have associated with

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

When using this product, do; heat

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

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 year & older: Apply liberally to painful area and massage until gel is absorbed into the skin. Repeat 3 to 4 time daily.

Blue 1, Camphor, Carbomer, Isopropyl Alcohol, Methylchoroisothiazolinone, Methylisothiazolinone, Sodium Hydroxide, Water.

Arctic Ice

Analgesic Gel

NET WT. 8 OZ. (227)

ARCTIC ICE ANALGESIC

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:22431-600
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
METHYLCHLOROISOTHIAZOLINONE (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:22431-600-01	227 g in 1 JAR; Type 0: Not a Combination Product	09/16/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part341	09/16/2015	

Labeler - Blue Cross Labs (008298879)

Registrant - Blue Cross Labs (008298879)

Establishment

Name	Address	ID/FEI	Business Operations
Blue Cross Labs		008298879	manufacture(22431-600)

Revised: 1/2017

Blue Cross Labs