

**WALGREENS ICE BLUE GEL- menthol gel**  
**Walgreens Company**

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**Walgreens Ice Blue Gel**

Menthol 2%

Topical analgesic

**Uses**

Temporarily relieves minor aches and pains of muscles and joints associated with

- simple backache
- arthritis
- strains
- bruises
- sprains

For external use only

- Do not use with other topical pain relievers
- Do not use with heating pads or other heating devices
- Avoid contact with eyes
- Do not bandage tightly
- Do not apply to wounds or damage skin

**Stop use and ask a doctor if:**

Condition worsens, or if symptoms persist for more than 7 day or clear up and occur again withing a few days

**If pregnant or breastfeeding**

Ask a health professional before use

**Keep out of reach of children**

If swallowed get medical help or contact Poison Control Center (800-222-1222) right away

**Directions**

- Clean affected area before applying product
- Adults and children 2 years and older, apply to the affected area not more than 3 to 4 times daily
- Children under 2 years of age, ask a doctor before using this product

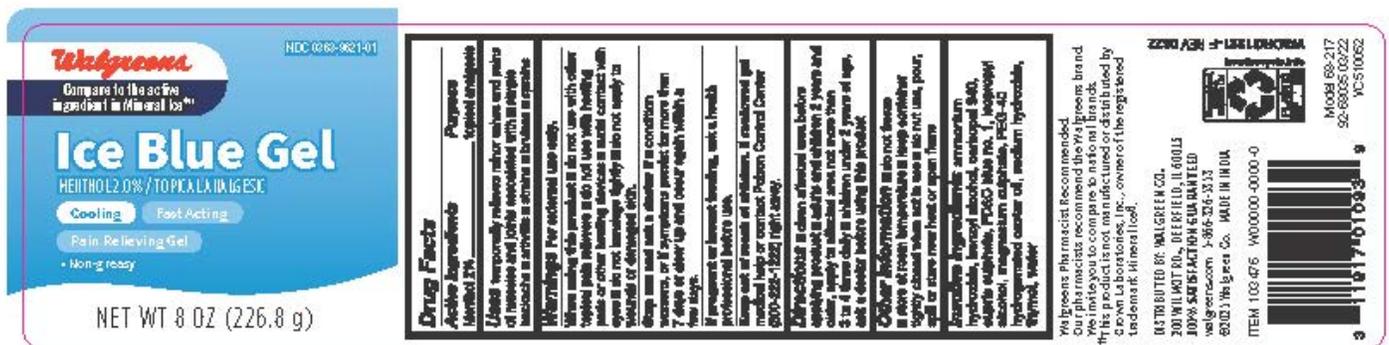
## Other Information

- Do not freeze
- Store at room temperature
- Keep container tightly closed when not in use
- do not use, pour, spill or store near heat or open flame

## Inactive Ingredients

Ammonium hydroxide, benzyl alcohol, carbopol 940, cupric sulphate, FD&C blue no.1, isopropyl alcohol, magnesium sulphate, PEG-40, hydrogenated castor oil, sodium hydroxide, thymol, water

## Walgreens Ice Blue Gel



## WALGREENS ICE BLUE GEL

menthol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:0363-9621
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2 g in 100 g

### Inactive Ingredients

Ingredient Name	Strength
THYMOL (UNII: 3J50XA376E)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CUPRIC SULFATE (UNII: LRX7AJ16DT)	
MAGNESIUM SULFATE, UNSPECIFIED (UNII: DE08037SAB)	
WATER (UNII: 059QF0KO0R)	
AMMONIA (UNII: 5138Q19F1X)	

<b>POLYOXYL 40 HYDROGENATED CASTOR OIL</b> (UNII: 7YC686GQ8F)	
<b>ISOPROPYL ALCOHOL</b> (UNII: ND2M416302)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	
<b>FD&amp;C BLUE NO. 1</b> (UNII: H3R47K3TBD)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0363-9621-01	226.8 g in 1 JAR; Type 0: Not a Combination Product	04/01/2022	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/01/2022	

**Labeler** - Walgreens Company (008965063)

### Establishment

Name	Address	ID/FEI	Business Operations
ANICARE PHARMACEUTICALS PRIVATE LIMITED		916837425	manufacture(0363-9621)

Revised: 12/2024

Walgreens Company