

WALGREENS ICE BLUE GEL- menthol gel
Walgreens Company

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.

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

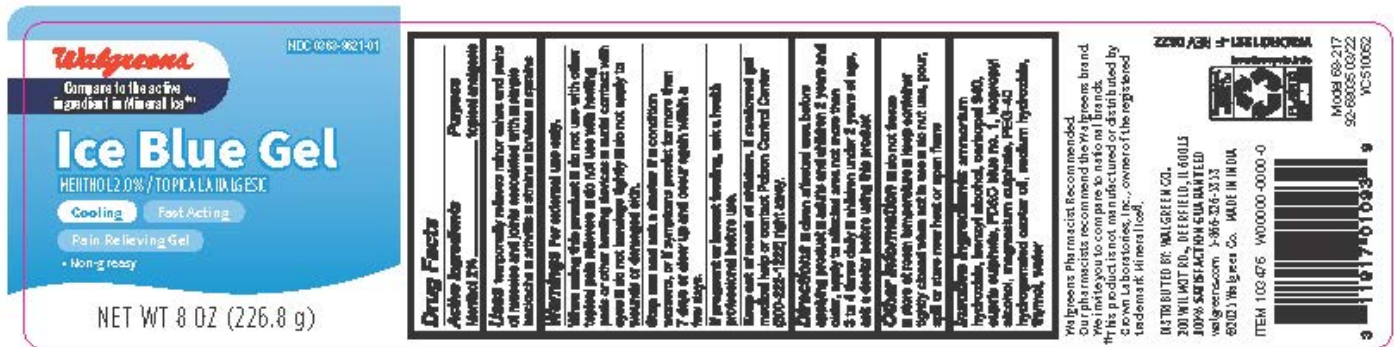
Other Information

- 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			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0363-9621
Route of Administration	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)	
AMMONIA (UNII: 5138Q19F1X)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	

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 not final	part348	04/01/2022	

Labeler - Walgreens Company (008965063)

Establishment

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

Revised: 4/2022

Walgreens Company