PURE-AID BLUE ICE GEL- menthol gel
Kareway Product, Inc.`

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.

Pure-Aid Blue Ice Gel 8oz

Menthol 1.25%

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
- Do not use with other topical pain relievers
- Do not use with heating pads or other heating devices

When using this product
- do not use in or near the 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 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 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

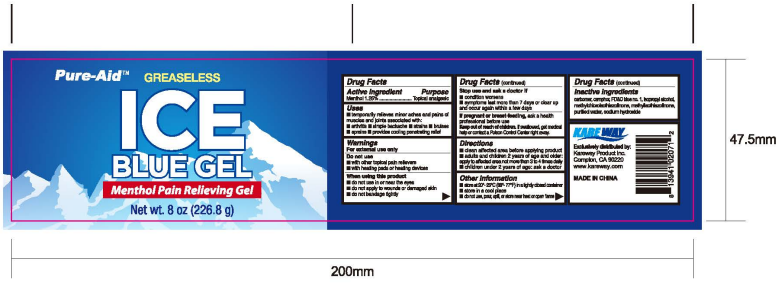
Other Information

- Store at 20ºC-25ºC (68º-77ºF) in a tightly closed container
- store in a cool place
- do not use, pour, spill or store near heat or open flame

Inactive Ingredients

carbomer, camphor, FD&C blue no.1, isopropyl alcohol, methylchoroisothiazolinone, methylsothiazolinone, purified water, sodium hydroxide

Pure-Aid Blue Ice Gel



pure-Aid ICE BLUE GEL			2022.05.18
Item	NO # 2071	Barcode.	8-13941-02071-2
Spec.	SIZE : 200X47.5mm		DIE LINE
CYAN	MAGENTA	YELLOW	BLACK

PURE-AID BLUE ICE GEL

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:67510-0671
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	1.25 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
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:67510-0671-8	226.8 g in 1 JAR; Type 0: Not a Combination Product	07/20/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	07/20/2022	

Labeler - Kareway Product, Inc.` (121840057)

Revised: 7/2022 Kareway Product, Inc.`