U ICE COLD- menthol ointment
UNIVERSAL DISTRIBUTION CENTER LLC

Drug Facts
Active Ingredients:

Menthol 1 percent.....Topical Analgesic

Camphor 0.5 percent.....Topical Analgesic

Purpose

Topical Analgesic

Uses:

for the temporary relief of minor aches and pains in muscles and joints associated with:

simple backache, strains, sprains, sports injuries, arthritis, bruises

Warnings: External Use Only

For external use only.

Do not use

with other topical pain relievers

with heating pads or heating devices

When using this product:

- Do not use in or near eyes
- Do not apply to wounds or damaged skin
- Do not bandage tightly

Stop use and ask Doctor if:

- Condition worsens, symptoms last more than 7 days or clear up and occur again within a few days
- Redness or irritation develops

If pregnant or breast-feeding,

ask a health professional before use

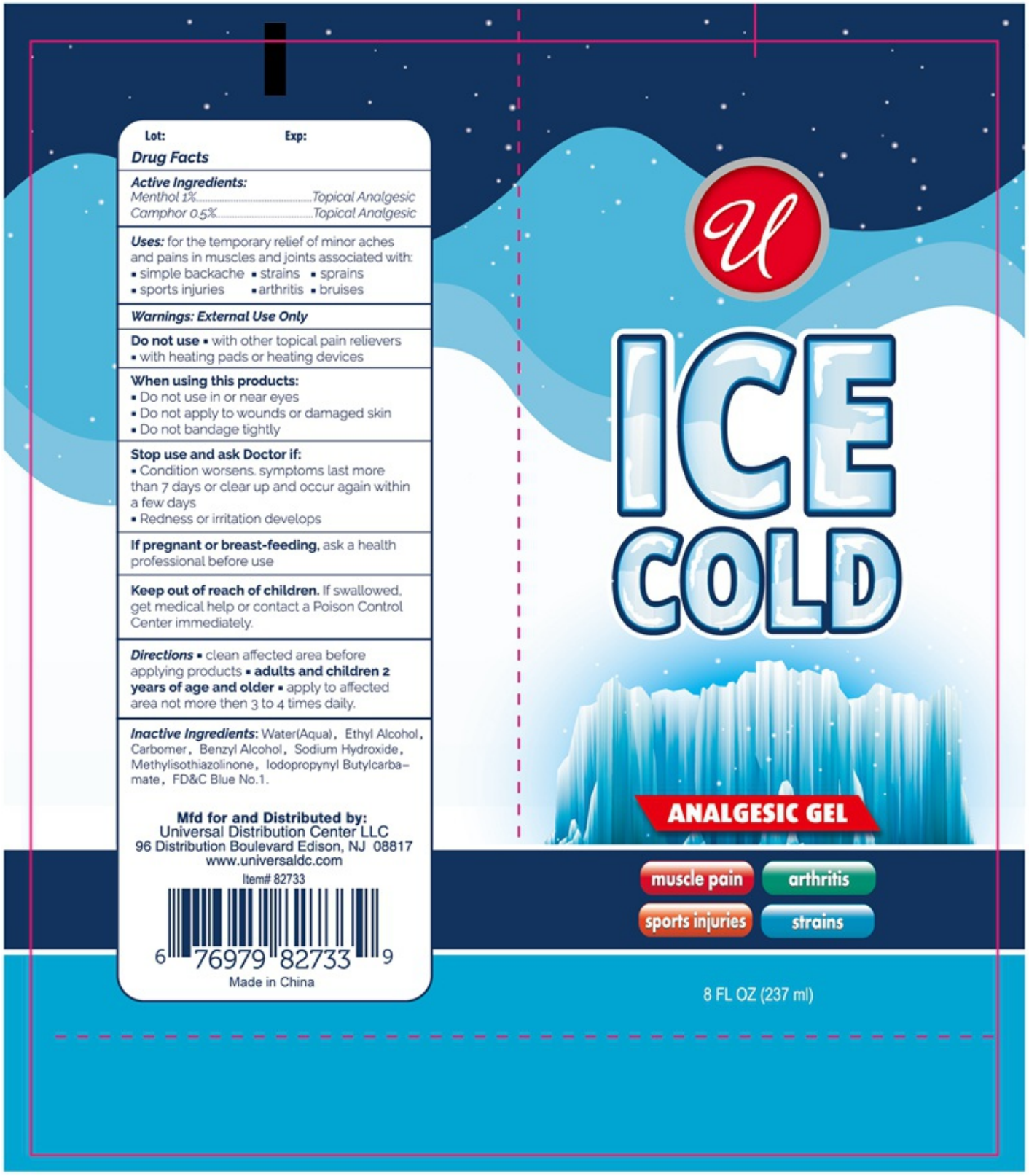
Keep out of reach of children.

If swallowed, get medical help or contact a Posion Control Center immediately.

Directions

- clean affected area before applying products
- **adults and children 2 years of age and older**
- apply to affected area not more than 3 to 4 times daily

Inactive Ingredients:



U ICE COLD

menthol ointment

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-045
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
CAMPHOR, (-)- (UNII: 213N3S8275) (CAMPHOR, (-)- - UNII:213N3S8275)	CAMPHOR, (-)-	0.5 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	1 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CARBOMER 934 (UNII: Z135WT9208)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:52000-045-01	227 g in 1 TUBE; Type 0: Not a Combination Product	08/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	08/01/2018	

Labeler - UNIVERSAL DISTRIBUTION CENTER LLC (019180459)

Revised: 9/2023 UNIVERSAL DISTRIBUTION CENTER LLC