UNIVERSAL ICE COLD ANALGESIC- menthol gel
Universal Distribution Center LLC

UNIVERSAL ICE COLD ANALGESIC GEL

Active Ingredients

Menthol 1.25%

Purpose

Topical Analgesic

Uses

Temporary relieves of minor aches and pains in muscles and joints associated with
- arthritis
- simple backache
- strains
- bruises
- sports injuries
- sprains

Warnings

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 the 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 clean 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 the reach of children

If swallowed, get medical help or contact a Poison 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

Water, Propylene glycol, Isopropyl alcohol, Carbomer, Sodium hydroxide, Benzyl alcohol, Edetate Disodium, Butylated hydroxytoluene, Camphor, Fd&c Blue No. 1

PRINCIPAL DISPLAY PANEL

ICE COLD
ANALGESIC GEL
8 FL.OZ (237 ml)

UNIVERSAL ICE COLD ANALGESIC

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-106
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	
CARBOMER HOMOPOLYMER TYPE C (ALLYL PENTAERYTHRITOL CROSSLINKED) (UNII: 4Q93RCW27E)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-106-01	237 mL in 1 TUBE; Type 0: Not a Combination Product	05/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2020	

Labeler - Universal Distribution Center LLC (019180459)

Revised: 11/2023 Universal Distribution Center LLC