

ICE GEL ICE COLD ANALGESIC- menthol gel
Jell Pharmaceuticals Pvt Ltd

Ice Gel

Active ingredient

Menthol 1.25%

Purpose

Topical Analgesic

Uses

■ temporarily relieves minor aches and pains of muscles and joints associated with:
■ arthritis ■ simple backache ■ strains ■ bruises ■ sports injuries ■ sprains ■
provides cooling penetrating relief

Warnings

For external use only

Do not use

■ with other topical 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 a doctor if

■ condition worsens ■ symptoms last more than 7 days or clear up and occur again within a few days ■ redness or irritation develops

Keep out of reach of children

If pregnant or breast-feeding, ask a health professional before use. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ clean affected area before applying product ■ adults and children 2 years of age and older: apply to affected area not more than 3 to 4 times daily ■ children under 2 years of age: ask a doctor

Other information

■ store at controlled room temperature 20 to 25 °C (68 to 77°F) in a tightly closed container ■ do not use, pour, spill or store near heat or open flame

Inactive Ingredients

benzyl alcohol, BHT, camphor, carbopol, disodium EDTA, FD&C blue no. 1, isopropyl alcohol, PEG-40 hydrogenated castor oi, propylene glycol, sodium hydroxide, water

Package Label



ICE GEL ICE COLD ANALGESIC

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:30400-110
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
BUTYLATED HYDROXYTOLUENE (UNII: 1P9D0Z171K)	

CARBOXPOLYMETHYLENE (UNII: 0A5MM307FC)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
POLYOXYL 40 HYDROGENATED CASTOR OIL (UNII: 7YC686GQ8F)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
WATER (UNII: 059QF0KO0R)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:30400-110-01	227 g in 1 JAR; Type 0: Not a Combination Product	04/02/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	04/02/2018	

Labeler - Jell Pharmaceuticals Pvt Ltd (726025211)

Establishment

Name	Address	ID/FEI	Business Operations
Jell Pharmaceuticals Pvt Ltd		726025211	manufacture(30400-110)

Revised: 2/2026

Jell Pharmaceuticals Pvt Ltd