WISH ULTRA WINTER ICE- menthol gel
LJ Pharma

Active Ingredient	**Purpose**
Menthol 2%	Topical analgesic

Uses

Temporarily relieves minor aches and pains of muscles and joints associated with:

- Arthritis
- Simple backache
- Strains
- Bruises
- Sprains
- Provides cooling relief

Warnings

For external use only

Do not use:

- With other topical pain relievers
- With heating pads or heating devices

Stop use and ask a doctor if:

- Condition worsens
- Symptoms last more than 7 days and/or subside and occur again within a few days

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 Poison Control Center right away.

Directions

- Clean affected area before applying product
- Adults and children 2 years and older: Apply to affected area no more than 3 to 4 times daily
- Children under 2 years of age: Ask a doctor

Other information

- store at 20° - 25°C (68° - 77° F) [see USP Controlled Room Temperature], in a tightly closed container
- store in a cool place
- do not use, pour, spill or store near heat or open flame

Inactive Ingredients

Ammonium hydroxide, carbomer, cupric sulfate, FD&C blue no. 1, isopropyl alcohol, magnesium sulfate, purified water, sodium hydroxide, thymol

Questions or Comments? 1-888-402-0184

Product label

WISH ULTRA WINTER ICE

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:84809-008
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	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
AMMONIA (UNII: 5138Q19F1X)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	

CUPRIC SULFATE (UNII: LRX7AJ16DT)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
THYMOL (UNII: 3J50XA376E)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84809-008-01	200 g in 1 BOTTLE; Type 0: Not a Combination Product	07/25/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	07/25/2025	

Labeler - LJ Pharma (914078633)

Establishment

Name	Address	ID/FEI	Business Operations
LJ Pharma		914078633	manufacture(84809-008)

Revised: 8/2025 LJ Pharma