

MINERAL ICE PAIN RELIEVING- menthol gel

Crown Laboratories

Mineral Ice

Active ingredient

Menthol 2%

Purpose

Topical analgesic

Uses

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

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 a doctor if

- condition worsens
- symptoms last more than 7 days or clear up 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 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 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?

call **1-833-279-6522**

Additional Information Listed on Other Panels

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Principal Display

NDC 0316-0226-08

ORIGINAL THERAPEUTIC

Mineral Ice®

Menthol Pain Relieving Gel

Greaseless with DEEPCOLD® Pain Reliever

Net wt. 8 oz (226.8 g)

P11529.00



MINERAL ICE PAIN RELIEVING

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0316-0226
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
AMMONIA (UNII: 5138Q19F1X)	
CARBOMER HOMOPOLYMER TYPE B (UNII: HHT01ZNK31)	
CUPRIC SULFATE (UNII: LRX7AJ16DT)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
MAGNESIUM SULFATE HEPTAHYDRATE (UNII: SK47B8698T)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
THYMOL (UNII: 3J50XA376E)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0226-35	99.2 g in 1 JAR; Type 0: Not a Combination Product	12/01/2018	01/31/2025
2	NDC:0316-0226-08	226.8 g in 1 JAR; Type 0: Not a Combination Product	12/01/2018	
3	NDC:0316-0226-16	453.6 g in 1 JAR; Type 0: Not a Combination Product	12/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	11/01/2011	

Labeler - Crown Laboratories (119508400)

Revised: 1/2026

Crown Laboratories