MMM ICE GEL- menthol gel
Southern Sales & Services, Inc

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

MMM Therapeutic Ice Gel
Topical Analgesic

Active Ingredients Menthol 2.0% Purpose: Topical Analgesic

Topical Analgesic

For the temporary relief of minor aches and pains of muscles and joints associated with
- Simple backache
- Arthritis
- Strains
- Bruises
- Sprains

For external use only. Avoid contact with eyes.

• do not bandage tightly
• do not apply to wounds or damaged skin

Stop use and ask a doctor if

• condition worsens • symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away

If pregnant or breast-feeding, ask a health professional before use.

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: consult a doctor.

Other information

• do not freeze
• Keep lid tightly closed

Inactive ingredients carbomer, propylene glycol, methylparaben, isopropyl alcohol, triethanol- amine, FD&C Blue no. 1, purified water.

227 g NDC: 69822-013-80

MMM Therapeutic Ice Gel

Topical Analgesic

Net Wt. 8 oz

MMM ICE GEL

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69822-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	2 g in 100 g

Inactive Ingredients	
Ingredient Name	**Strength**
TROLAMINE (UNII: 9O3K93S3TK)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
CARBOMER HOMOPOLYMER TYPE C (UNII: 4Q93RCW27E)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:69822-013-80	227 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	01/01/2021	

Labeler - Southern Sales & Services, Inc (013114906)

Registrant - Southern Sales & Services, Inc (013114906)

Establishment

Name	Address	ID/FEI	Business Operations
Southern Sales & Services, Inc		013114906	label(69822-013)

Revised: 1/2021

Southern Sales & Services, Inc