SUNSET COOLING PAIN RELIEF ROLL-ON- menthol gel
SUNSET NOVELTIES, INC

72937-631-03

Menthol 4%

Topical Analgesic

Use

Aid for temporary local relief of minor pain in muscles and joints.

• For external use only.

• Ask a doctor before use if you have redness over affected area.

• Avoid contact with the eyes or mucous membranes.
• Do not apply to wounds or damaged skin.
• Do not apply to the irritated skin or if excessive irritation develops.
• Do not bandage tightly.
• Do not use with heating pad or device.
• Stop use and ask a doctor if condition worsens, or if symptoms persist for 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.

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Adults and Children over 12 years of age.

● Apply a small amount on the affected area.

● Massage in circular motion, let set for a few seconds.

● Repeat as necessary, but no more than 3 to 4 times daily.

Children under 12 years of age: Do not use, consult a doctor

Other information

Store tightly closed in a dry place at controlled room temperature between 59°F-86° F (15°C-30° C).

Water (Aqua), Alcohol Denat, Ceteareth-25, Caprylic/Capric Triglyceride, Glycerin, Calendula Officinalis Extract, Cannabis Sativa Seed Oil, Carbomer, Sodium Hydroxide, Methyl Salicylate, Benzyl Alcohol, Salicylic Acid, Sorbic Acid, FD&C Blue No.1 (CI 42090).

Questions or comments?

Contact us +1 (888) 367-4916

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menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72937-631
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	4 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SORBIC ACID (UNII: X045WJ989B)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
CETEARETH-25 (UNII: 8FA93U5T67)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
ALCOHOL (UNII: 3K9958V90M)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

Product Characteristics			
Color	blue	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-631-03	89 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/01/2025	

Labeler - SUNSET NOVELTIES, INC (067218145)

Revised: 12/2025 SUNSET NOVELTIES, INC