MENTHOL- menthol gel
SUNSET PAIN RELIEF

72937-632-03

Menthol 4%

Topical Analgesic

USE

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

For external use only. · Ask a doctor before use if you have redness over affected area

Use only as directed.
Do not bandage tightly or use with a heating pad.
Avoid contact with eyes and mucous membranes.
Do not apply to wounds or damaged, broken or irritated skin.
If you experience an allergic reaction, discontinue use and consult a doctor.

Do not expose the area treated with product to heat or direct sunlight.

Stop use and ask a doctor if

Condition worsens.
Redness is present.
Irritation develops.
Symptoms persist for more than 7 days or clear up occur again within a few days.

You experience signs injury, such as pain, swelling or blistering where the product was applied.

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

Apply a small amount to 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.

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

Water (Aqua), Alcohol Denat, Ceteareth-25, Glycerin, Calendula Officinalis Extract, Caprylic/Capric Triglyceride, Carbomer, Sodium Hydroxide, Cannabis Sativa Seed Oil, Methyl Salicylate, Benzyl Alcohol, Salicylic Acid, Sorbic Acid.

SUNSET PAIN RELIEF +350 HEMP ROLL-ON

MENTHOL

menthol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72937-632
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	4 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
CETEARETH-25 (UNII: 8FA93U5T67)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	
ALCOHOL (UNII: 3K9958V90M)	

SALICYLIC ACID (UNII: O414PZ4LPZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
SORBIC ACID (UNII: X045WJ989B)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CALENDULA OFFICINALIS FLOWER (UNII: P0M7O4Y7YD)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-632-03	90 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	12/20/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/20/2025	

Labeler - SUNSET PAIN RELIEF (067218145)