

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



SUNSET
PAIN RELIEF GEL

PAIN RELIEVING & RELAXING GEL
REDUCES ANXIETY, HELPS THE NERVOUS AND CARDIOVASCULAR SYSTEM

+350 HEMP SEED OIL*

ROLL - ON

3 fl oz e (89mL)



Exclusively Distributed by: SUNSET NOVELTIES,
620 SW 12th AVE. POMPAHO BEACH, FL 33069
www.sunsetcbd hemp.com
MANUFACTURED IN USA



Drug Facts

Active ingredients	Purpose
Menthol 4%	Topical Analgesic

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

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

When using this product
• 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.

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
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).

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

Questions or comments?
Contact us +1 (888) 367-4916

* These statements have not been evaluated by the Food and Drug Administration. This product is not intended to diagnose, treat, cure, or prevent any disease.

NDC 72937-632-03

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
CETARETH-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, WTH 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)

Revised: 12/2025

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