MENTHOL, CAMPHOR- menthol, camphor gel SUNSET NOVELTIES, INC

72937-006-42

Camphor 2%

Menthol 6%

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

A transient burning sensation or redness may occur upon application but generally disappears in several days

If you experience an allergic reaction, discontinue use, and consult a physician

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.

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

DIRECTIONS

Adults and Children over 12 years

Apply a small amount on the affected area.

Massage in circular motion until absorbed.

Repeat as needed, but no more than 3 to 4 times per day.

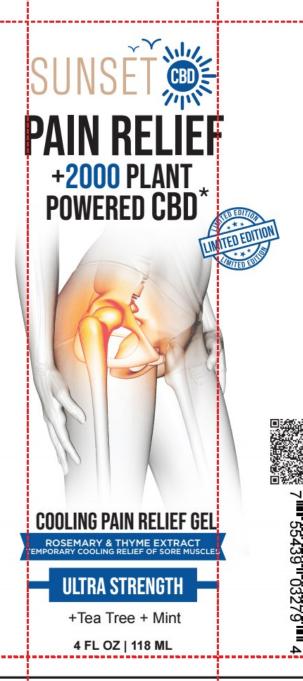
Wash hands with soap and water after use.

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, Propylene Glycol, Polysorbate 20, Carbomer, Sodium Hydroxide, Salicylic Acid, Glycerin, Mentha Piperita (Peppermint) Oil, Rosmarinus Officinalis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Cannabis Sativa Seed Oil, Benzyl Alcohol, Sorbic Acid, Cannabidiol, FD&C Blue No. 1 (Cl 42090), FD&C Yellow No.5 (Cl 19140).

SUNSET - COOLING PAIN RELIEF GEL 402 TUBE LIMITED EDITION



Drug Facts

Active ingredients

Purpose

Menthol 6%

Topical Analgesic

NDC#

72937-006-42

Aid for temporary local relief of minor pain in muscles or

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
- A transient burning sensation or redness may occur upon application but generally disappears in several days
 If you experience an allergic reaction, discontinue use, and
- Try of experience an analysic reaction, discommus use, consult a physician
 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 Contro Center right away.

Adults and Children over 12 years:

- Apply a small amount on desired area
- Massage in circular motions until absorbed.
- Repeat as needed but no more than 3 to 4 times per day. Store tightly closed in a dry place at room temperature between 59°-86° F (15°-30° C)

 • Wash hands with soap and water after use

Children under 12 years of age: do not use, co

Other information

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

Inactive Ingredients

Water (Aqua), Alcohol Denat, Propylene Glycol, Polysorba 20, Carbomer, Sodium Hydroxide, Salicylic Acid, Glycerin, 20, Carbonier, Soulini Hydroxide, Salcoyia Acab, Gylochia Mentha Piperita (Peppermint) Oil, Rosmarinus Officinallis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Cannabis Sativa Seed Oil, Benzyl Alcohol, Sorbic Acid, Cannabidiol, FD&C Blue No. 1 (Cl 42090), FD&C Yellow No.5 (Cl 19140).

*This statement has not been evaluated by the Food and Drug Administration (FDA). This product is not intended to diagnose, treat, cure or prevent any

ESS THAN 0.3% THC MADE IN USA

Exclusively Distributed by: SUNSET NOVELTIES.

5700 PEMBROKE RD WEST PARK, FL 33023.

(888) 367 4916. www.sunsetcbdhemp.com

MENTHOL, CAMPHOR

menthol, camphor gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:72937-006

TOPICAL Route of Administration

Active Ingredient/Active Moiety

Ingredient Name Basis of Strength Strength

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 g in 100 g
CAMPHOR, (-)- (UNII: 213N3S8275) (CAMPHOR, (-) UNII:213N3S8275)	CAMPHOR, (-)-	2 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
PEPPERMINT OIL (UNII: AV092KU4JH)		
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)		
CARBOMER 940 (UNII: 4Q93RCW27E)		
GLYCERIN (UNII: PDC6A3C0OX)		
SALICYLIC ACID (UNII: O414PZ4LPZ)		
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)		
POLYSORBATE 20 (UNII: 7T1F30V5YH)		
ROSEMARY OIL (UNII: 8LGU7VM393)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
SORBIC ACID (UNII: X045WJ989B)		
WATER (UNII: 059QF0KO0R)		
BENZYL ALCOHOL (UNII: LKG8494WBH)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		
THYME OIL (UNII: 2UK410MY6B)		
MELALEUCA ALTERNIFOLIA (TEA TREE) LEAF OIL (UNII: VIF565UC2G)		
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)		
ALCOHOL (UNII: 3K9958V90M)		
CANNABIDIOL (UNII: 19GBJ60SN5)		

Product Characteristics		
Color	white (TURQUOISE)	Score
Shape		Size
Flavor		Imprint Code
Contains		

Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
		NDC:72937-006- 42	113 g in 1 TUBE; Type 0: Not a Combination Product	02/07/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	09/22/2020	