

**MENTHOL, CAMPHOR- menthol, camphor cream**  
**SUNSET NOVELTIES, INC**

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**72937-103-04**

Menthol 10%

Camphor 3%

External Analgesic

Pain Relieving

**USES:**

For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, sprains and strains.

For external use only.

Do not bandage tightly. Do not use with heating pad, pack, wrap, hot water bottle or any heating element.

In case of accidental ingestion, contact doctor immediately. If prone to allergic reaction to the product, consult to a doctor before using.

Avoid contact with eyes, mucous membranes, rashesm wounds or damaged skin.

Do nor apply on nose or genital area.

If rash, redness or itchiness results; discontinue use and consult a doctor.

**DIRECTIONS:**

Adults and children over 12 years of age; apply a thin layer to affected area and rub gently not more than 3 to 4 times a day.

Wash hands with soap and water after use.

Children under 12 years of age; do not use unless directed by doctor/physician

Keep away from children. Package not child resistant.

**OTHER INFORMATION:**

Store at room temperature 15° - 30°C (59° - 86°F)

Aqua, Paraffinum Liquidum, Alcohol Denat, Stearic Acid, Cetearyl Alcohol, Polysorbate 60, Cetyl Alcohol, Dimethicone, Glyceryl Stearate, Glycereth-26, Propylene Glycol, Diazolidinyl Urea, Methylparaben, Propylparaben, Tocopheryl Acetate, Cannabis Sativa Seed Oil, Stearyl Alcohol, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Polysorbate 20, Triethanolamine, Parfum, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Sodium Benzotriazolyl Butylphenol Sulfonate, Buteth-3, Tributyl Citrate, Cannabidiol, FD&C Blue #1 Powder (CI 42090), Linalool, Limonene, Benzyl Benzoate, Coumarin,

Geraniol.

## GREEN FARM PAIN RELIEF CREAM 4oz

**DRUG FACTS**

**ACTIVE INGREDIENTS**  
Menthol 10%, Camphor 2%

**PURPOSE**  
Topical Analgesic Pain Relievers

**USES:** For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, sprains and strains.

**WARNING:**  
• Must be 18 or older to purchase this product.  
• For external use only.  
• Do not massage topically. Do not use with heating pad, pack, wrap, hot water bottle or any heating device.  
• Keep away from children. Package not child resistant.  
• In case of accidental ingestion, contact doctor immediately. Be prone to allergic reaction to the product. Consult a doctor before using.

**WHEN USING THIS PRODUCT:** Avoid contact with eyes, mucous membranes, nostrils, mouth or damaged skin. Do not apply on more and painful areas. If rash, redness or further trouble, discontinue use and consult a doctor.

**LEAD TIME KJ'S INC.** Exclusively Distributed by: **SHADE BY USA**  
SUNSET HONEY WAX, 1000 WOODLAND RD, SUITE 100, WILKESBORO, NC 27651  
WWW.SUNSETHONEY.COM

**4 FL. OZ.**

**GREEN FARM**  
INFUSED CBD PRODUCTS

**PAIN RELIEF CREAM +1000 STRENGTH**

**ULTRA STRENGTH**

**DIRECTIONS**  
Adults and children over 12 years of age:  
• Apply a small amount on desired area.  
• Massage in circular motions until absorbed.  
• Repeat as needed but not more than 3 to 4 times a day.  
• Wash hands with soap and water after use.

**OTHER INFORMATION:**  
• Store at room temperature 15° - 30°C (59° - 86°F)

**Inactive Ingredients:** Aqua, Paraffinum Liquidum, Alcohol Denat, Stearic Acid, Cetylmyristyl Alcohol, Polysorbate 80, Cetyl Alcohol, Dimethylsiloxane, Glyceryl Stearate, Olefin-20, Propylene Glycol, Cholesterol, Urea, Menthylacetate, Propylparaben, Tocopherol Acetate, Camphor Saliva Sweet Oil, Stearyl Alcohol, Argemone/Sodium Argemone Copolymer, Tocopherol, Polysorbate 20, Triethanolamine, Parfum, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symptol, Ethylhexyl Glycerin, Hydroxypropyl, Sodium Benzoate/ethyl Benzoate, Salicylate, Butyl-2, Triethyl Citrate, Citronellol, FDAC Blue #1 Powder (CI 42090), Limonol, Limonene, Benzyl Benzoate, Coumarin, Geraniol.

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 disease.

## MENTHOL, CAMPHOR

menthol, camphor cream

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:72937-103
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>MENTHOL</b> (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 g
<b>CAMPHOR (NATURAL)</b> (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	3 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
TRIBUTYL CITRATE (UNII: 827D5B1B6S)	
PANTHENOL (UNII: WW9CM0O67Z)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
MINERAL OIL (UNII: T5L8T28FGP)	
BUTETH-3 (UNII: OC116GRO69)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALCOHOL (UNII: 3K9958V90M)	
ACRYLIC ACID/SODIUM ACRYLATE COPOLYMER (1:1; 600 MPA.S AT 0.2%) (UNII: M4PPW69Y4H)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERETH-26 (UNII: NNE56F2N14)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
BASIC BLUE 1 (UNII: 92N74OA24D)	
WATER (UNII: 059QF0KO0R)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM BENZOTRIAZOLYL BUTYLPHENOL SULFONATE (UNII: 0LA2QC9O3Z)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE 1000 (UNII: MCU2324216)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
TROLAMINE (UNII: 9O3K93S3TK)	

## Product Characteristics

Color	green	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-103-04	113 g in 1 BOTTLE; Type 0: Not a Combination Product	10/05/2020	03/12/2026

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug M017

10/05/2020

03/12/2026

**Labeler** - SUNSET NOVELTIES, INC (067218145)

Revised: 11/2025

SUNSET NOVELTIES, INC