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

72937-310-08

Camphor 3%

Menthol 10%

Topical Analgesic

Use

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

Use only as directed. • 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.

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.

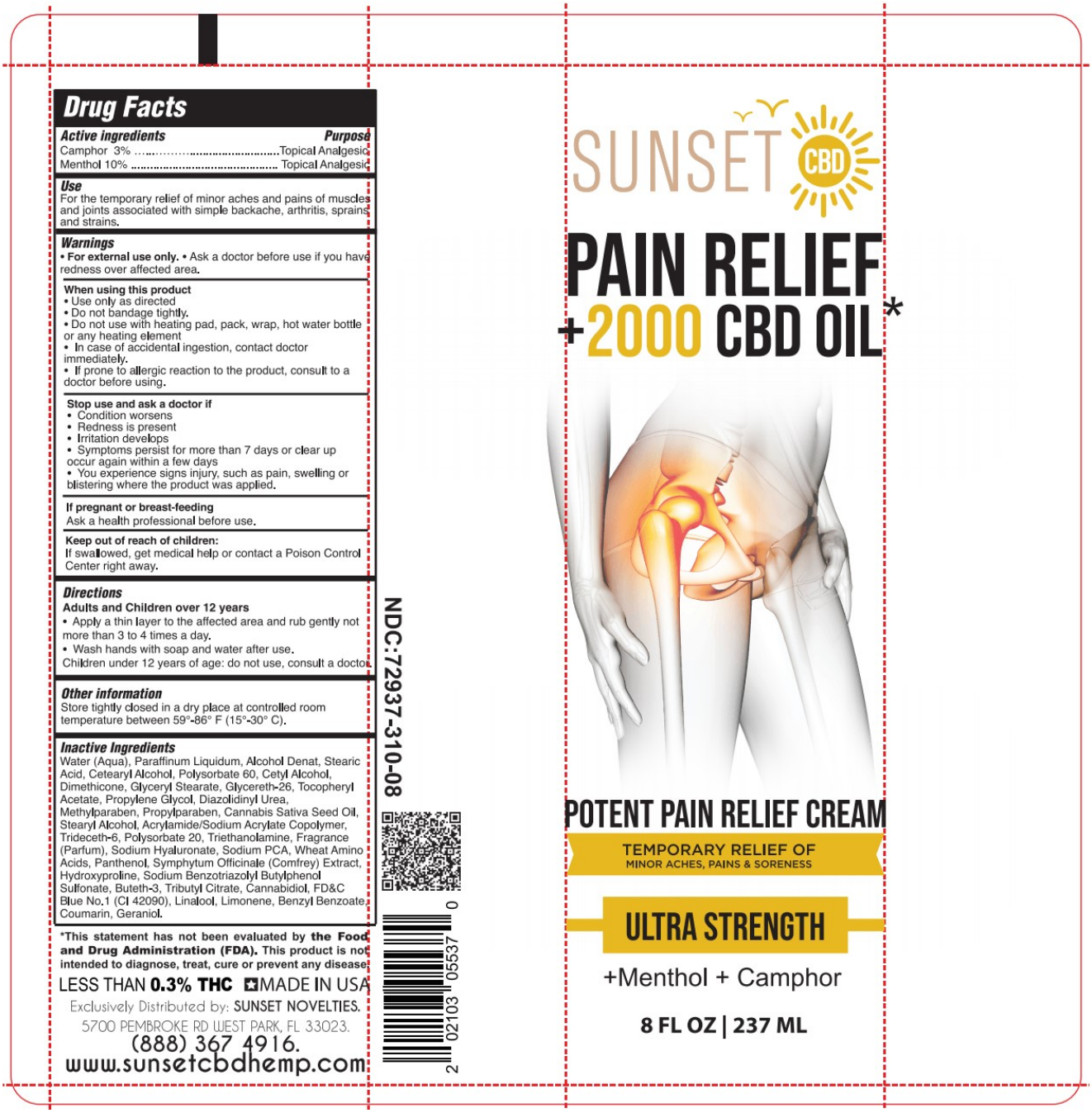
Keep out of reach of children
If swallowed, get medical help, or contact a Poison Control Center right away.

Directions

Adults and Children over 12 years
• Apply a thin layer to the 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 consult a doctor

Water (Aqua), Paraffinum Liquidum, Alcohol Denat, Stearic Acid, Cetearyl Alcohol, Polysorbate 60, Cetyl Alcohol, Dimethicone, Glyceryl Stearate, Glycereth-26, Tocopheryl Acetate, Propylene Glycol, Diazolidinyl Urea, Methylparaben, Propylparaben, Cannabis Sativa Seed Oil, Stearyl Alcohol, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Polysorbate 20, Triethanolamine, Fragrance (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 No.1 (CI 42090), Linalool, Limonene, Benzyl Benzoate, Coumarin, Geraniol.

SUNSET PAIN RELIEF CREAM TUBE 8 FL OZ



MENTHOL, CAMPHOR

menthol, camphor cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72937-310
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 g
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
TRIBUTYL CITRATE (UNII: 827D5B1B6S)	
PANTHENOL (UNII: WV9CM0O67Z)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
ACRYLIC ACID/SODIUM ACRYLATE COPOLYMER (1:1; 600 MPA.S AT 0.2%) (UNII: M4PPW69Y4H)	
GLYCERETH-26 (UNII: NNE56F2N14)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
BUTETH-3 (UNII: OC116GRO69)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
WATER (UNII: 059QF0KO0R)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
SODIUM BENZOTRIAZOLYL BUTYLPHENOL SULFONATE (UNII: 0LA2QC9O3Z)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE 1000 (UNII: MCU2324216)	
ALCOHOL (UNII: 3K9958V90M)	
TROLAMINE (UNII: 9O3K93S3TK)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
COUMARIN (UNII: A4VZ22K1WT)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
BENZYL BENZOATE (UNII: N863NB338G)	
COMFREY (UNII: D05HXK6R3G)	
GERANIOL (UNII: L837108USY)	
MINERAL OIL (UNII: T5L8T28FGP)	

Product Characteristics

Color	green	**Score**	
Shape		**Size**	

Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-310-08	226 g in 1 TUBE; Type 0: Not a Combination Product	03/04/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/04/2021	

Labeler - SUNSET NOVELTIES, INC (067218145)

Revised: 6/2024 SUNSET NOVELTIES, INC