MENTHOL, CAMPHOR- menthol, camphor cream
SUNSET PAIN RELIEF

72937-022-16

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.

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 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: 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), 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, FD&C Blue No.1 (CI 42090), Linalool, Limonene, Benzyl Benzoate, Coumarin, Geraniol.

SUNSET PAIN RELIEF CREAM GOLD 16 OZ

MENTHOL, CAMPHOR

menthol, camphor cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72937-022
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	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**
PANTHENOL (UNII: WV9CM0O67Z)	
ALCOHOL (UNII: 3K9958V90M)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHICONE 1000 (UNII: MCU2324216)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	

TRIDECETH-6 (UNII: 3T5PCR2H0C)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
BENZYL BENZOATE (UNII: N863NB338G)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCERETH-26 (UNII: NNE56F2N14)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
TRIBUTYL CITRATE (UNII: 827D5B1B6S)	
GERANIOL (UNII: L837108USY)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
TROLAMINE (UNII: 9O3K93S3TK)	
BUTETH-3 (UNII: OC116GRO69)	
SODIUM BENZOTRIAZOLYL BUTYLPHENOL SULFONATE (UNII: 0LA2QC9O3Z)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
COUMARIN (UNII: A4VZ22K1WT)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
WATER (UNII: 059QF0KO0R)	
MINERAL OIL (UNII: T5L8T28FGP)	

Product Characteristics

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-022-16	473 g in 1 JAR; Type 0: Not a Combination Product	02/12/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	02/12/2026	

Labeler - SUNSET PAIN RELIEF (067218145)

Revised: 2/2026 SUNSET PAIN RELIEF