

HEATING PAIN RELIEF- methyl salicylate, menthol cream
SUNSET NOVELTIES, INC

72937-017-20

Methyl Salicylate 18%

Menthol 10%

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.

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, let set for a few seconds..

Repeat as necessary, but no more than 3 to 4 times daily.

Wash hands with soap and water after use.





Children under 12 years of age consult a doctor.

Other information

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

Water (Aqua), Paraffinum Liquidum, Cetyl Alcohol, Glyceryl Stearate, Stearic Acid, Glycereth-26, Dimethicone, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Caprylyl Glycol, Hexylene Glycol, Stearyl Alcohol, Triethanolamine, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Cannabis Sativa (Hemp) Seed Oil, Phenoxyethanol, FD&C Yellow No.6 (CI 15985).

SUNSET - HEATING PAIN RELIEF CREAM 4 Fl oz TUBE

Drug Facts	<p>*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.</p> <p>Endorsed by: SUNSET NOVELTIES 620 331 12th AVE. POMEROY BEACH, FL 33069 www.sunsetcbd hemp.com MANUFACTURED IN USA</p>	NDC # 72937-017-20
Active ingredients Menthol 10%.....Topical Analgesic Methyl Salicylate 18%.....Topical Analgesic		SUNSET  ULTRA STRENGTH PAIN RELIEF
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 • 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.		HEATING PAIN RELIEF CREAM MENTHOL & AMINO ACIDS TARGETED HEAT RELIEF OF SORE MUSCLES
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.		+250 HEMP SEED OIL*
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. • Wash hands with soap and water after use. Children under 12 years of age consult a doctor	 	2 fl oz e (59 mL)
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), Paraffinum Liquidum, Cetyl Alcohol, Glyceryl Stearate, Stearic Acid, Glycereth-26, Dimethicone, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Caprylyl Glycol, Hexylene Glycol, Stearyl Alcohol, Triethanolamine, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Hemp Seed Oil, Phenoxyethanol, FD&C Yellow No.6 (CI 15985).	
Questions or comments? Contact us +1 (888) 367-4916		

HEATING PAIN RELIEF
menthyl salicylate, menthol cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72937-017
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 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	18 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERETH-26 (UNII: NNE56F2N14)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
COMFREY LEAF (UNII: DG4F8T839X)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
PANTHENOL (UNII: VW9CM0O67Z)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIMETHICONE 1000 (UNII: MCU2324216)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
TROLAMINE (UNII: 9O3K93S3TK)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
MINERAL OIL (UNII: T5L8T28FGP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color	orange (Light Orange)	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72937-017-20	59 mL in 1 TUBE; Type 0: Not a Combination Product	01/29/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/29/2026	

Labeler - SUNSET NOVELTIES, INC (067218145)

Revised: 1/2026

SUNSET NOVELTIES, INC