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

72937-180-42

Menthol 10%

Methyl Salicylate 18%

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

SUNSET - HEATING PAIN RELIEF CREAM 4 oz TUBE LIMITED EDITION

HEATING PAIN RELIEF

methyl salicylate, menthol cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72937-180
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	9.8 g in 100 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	17.6 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)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
PANTHENOL (UNII: WV9CM0O67Z)	
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:72937-180-42	118 mL 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	05/25/2023	

Labeler - SUNSET NOVELTIES, INC (067218145)

Revised: 2/2024 SUNSET NOVELTIES, INC