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

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

72937-118-02 72937-118-04 72937-118-08 72937-118-16

Methyl Salicylate 18%

Menthol 10%

Topical Analgesic.

USES:

Aid for temporary local relief of minor pain in muscles or joints.

For external use only.

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 doctor.

Do not expose the area treated with product to heat or direct sunlight.

IF PREGNANT OR BREAST - FEEDING:

Ask a health professional before use.

If swallowed, get medical help or contact a Poison Control Center right away.

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.

DIRECTIONS:

Adults an childre over 12 years of age: apply a thin layer to affected area and rub gently no more than 3 to 4 times daily.

Wash hands with soap and water after use.

children undre 12 years of age; do not use unless directed by a doctor/physician.

Store at room temperature 15°C - 30°C.

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 #6 (CI 15985).

KANGAROO CBD HEATING PAIN RELIEF CREAM 2 oz



KANGAROO CBD HEATING PAIN RELIEF CREAM 4 oz



KANGAROO CBD HEATING PAIN RELIEF CREAM 8 oz



KANGAROO CBD HEATING PAIN RELIEF CREAM 16 oz





ils 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.

KANGAROO CBD HEATING PAIN RELIEF

methyl salicylate, menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72937-118
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: 2300U9XXE4)		
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)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
WATER (UNII: 059QF0KO0R)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
PANTHENOL (UNII: WV9CM0O67Z)	
HYDROXYPROLINE (UNII: RMB44W089X)	
HEXYLENE GLYCOL (UNII: KEH0A3F75J)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
DIMETHICONE 1000 (UNII: MCU2324216)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	
TROLAMINE (UNII: 903K93S3TK)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 4690TG57A2)	
MINERAL OIL (UNII: T5L8T28FGP)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

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

ı	Packaging				
7	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
:	NDC:72937-118- 02	60 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/29/2022		
:	NDC:72937-118- 04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/29/2022		
3	NDC:72937-118- 08	237 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/29/2022		
4	NDC:72937-118- 16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	11/29/2022		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	11/29/2022	

Labeler - SUNSET NOVELTIES, INC (067218145)

Revised: 12/2022 SUNSET NOVELTIES, INC