HEMPCIN ADVANCED TOPICAL PAIN RELIEF CREAM- menthol cream
Pedicis Research, LLC

HEMPCIN Advanced Topical Pain Relief Cream

Drug Facts:

Active Ingredient:

Menthol 5%

Purpose

Pain Relieving Cream

Uses:

For the temporary relief of minor aches and pain of muscles and joints

Warnings:

• For External Use only • Avoid contact with the eyes • Do not apply to wounds or damaged skin • Do not bandage tightly • Cap tube tightly and store at room temperature away from heat

When using this product:

• do not use in large quantities particularly over raw surfaces or blistered areas • do not exceed recommended doses unless recommended by a doctor

Stop use and ask a doctor if:

• allergic reaction occurs • condition worsens or does not improve within 7 days • symptoms clear up and do not return within a few days • you notice any unusual effects

Keep out of the reach of children:

If swallowed get medical help or call a Poison Control Center right away

Directions:

• Adults or children 2 years of age and older. Apply externally to affected area not more than 3-4 times daily. • Children under 2 years of age consult a doctor

Other Ingredients:

Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Cannabis Sativa (Hemp) Seed Oil, Cassia Oil, Cetearyl Alcohol, Dimethyl Sulfone (DMSO), Ethylhexylglycerin, Glycerin, Glyceryl Stearate, Ilex Paraguariensis (Yerba Mate') Extract, C13-14 Isoparaffin, Isopropyl Alcohol, Laureth-7, Magnesium Sulfate, Melaleuca Alternifolia (Tea Tree) Oil, Methylsulfonylmethane (MSM), PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Propylene Glycol, Sodium Polyacrylate, Stearic Acid, Triethanolamine

Questions:

Call toll free 1-800-748-6539

Package Labeling:

HEMPCIN ADVANCED TOPICAL PAIN RELIEF CREAM

menthol cream

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:51151-391
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	50 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CHINESE CINNAMON OIL (UNII: A4WO0626T5)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
LAURETH-7 (UNII: Z95S6G8201)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
TEA TREE OIL (UNII: VIF565UC2G)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51151-391-00	1 in 1 BOX	02/13/2023	
1		60 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing	Application Number or Monograph	Marketing Start	Marketing End

Category	Citation	Date	Date
OTC Monograph Drug	M017	02/13/2023	

Labeler - Pedicis Research, LLC (078496974)

Revised: 11/2023 Pedicis Research, LLC