BADASS BOTANICALS HEMP PAIN RELIEF- menthol cream Frontier Sales, LLC

BADASS BOTANICALS Hemp Pain Relief Cream

Drug Facts

Active Ingredient:

Menthol

Topical Analgesic

Indications:

For the temporary relief of minor aches and pains of the muscles and joints associated with arthritis, simple backache, sprains, bruises and strains.

Warnings:

- For external use only.
- Avoid contact with eyes.
- If symptoms persist for more than seven days, discontinue use and consult a physician.

Keep out of reach of children

- If swallowed, consult physician.
- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

If pregnant or breast feeding,

contact physician prior to use.

Directions:

- Adults and children two-years of age or older: Apply to affected area not more than three to four times daily.
- Children under two-years of age: consult a physician.

Additional Information:

Store at room temperature.

Other Ingredients:

Aloe Barbadensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, Butyrospermum Parkii (Shea) Butter, *Cannabis Sativa (Hemp) Oil, Cetyl Esters, Ethylhexylglycerin, Eucalyptus Globulus Oil, Glucosamine Sulfate, Glycerin, Glyceryl Stearate, Glycyrrhiza Glabra (Licorice) Extract, Helianthus Annuus (Sunflower) Oil, Isoproply Alcohol, Methyl Gluceth-20, Methyl Salicylate, Methylsulfonylmethane (MSM), PEG-100 Stearate, Phenoxyethanol, Steareth-20, Tocopheryl Acetate (Vitamin E), Triethanolamine, Zemea (Corn) Propanediol.

Package Labeling:



BADASS BOTANICALS HEMP PAIN RELIEF

menthol cream

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:73200-191

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)

MENTHOL

25 mg in 1 mL

Inactive Ingredients

Ingredient Name
Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)

ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)

INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)

SHEA BUTTER (UNII: K49155WL9Y)

HEMP (UNII: TD1MUT01Q7)

WATER (UNII: 059QF0KO0R)

CETYL ESTERS WAX (UNII: D072FFP9GU)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)

EUCALYPTUS OIL (UNII: 2R040NI662)

GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)

GLYCERIN (UNII: PDC6A3C0OX)

GLYCERYL MONOSTEARATE (UNII: 2300U9XXE4)

GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)

HELIANTHUS ANNUUS FLOWERING TOP (UNII: BKJ0J3D1BP)

ISOPROPYL ALCOHOL (UNII: ND2M416302)	
METHYL GLUCETH-20 (UNII: J3QD0LD11P)	
METHYL SALICYLATE (UNII: LAV5U5022Y)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
STEARETH-20 (UNII: L0Q8IK9E08)	
.ALPHATOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 903K93S3TK)	
CORN (UNII: 0N86727070)	

F	Packaging					
#	tem Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:73200-191- 02	60 mL in 1 JAR; Type 0: Not a Combination Product	11/30/2019			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M017	11/30/2019				

Labeler - Frontier Sales, LLC (081265948)

Revised: 11/2023 Frontier Sales, LLC