

HAVASU HEMP SOOTHING PAIN RELIEF RUB- histamine dihydrochloride cream
Havasu Nutrition LLC

HAVASU HEMP CREAM Soothing Pain Relief Rub

Drug Facts:

Active Ingredient

Histamine DHCl 0.03%

Purpose

Topical Analgesic

Uses:

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

Warnings:

- **For external use only.**

Do not

- apply to wounds or damaged skin.
- bandage tightly.

When using this product

- avoid contact with eyes.

Stop use and ask a doctor if

condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days.

Keep out of reach of children.

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

If pregnant or breast feeding,

ask a health professional before use.

Directions:

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




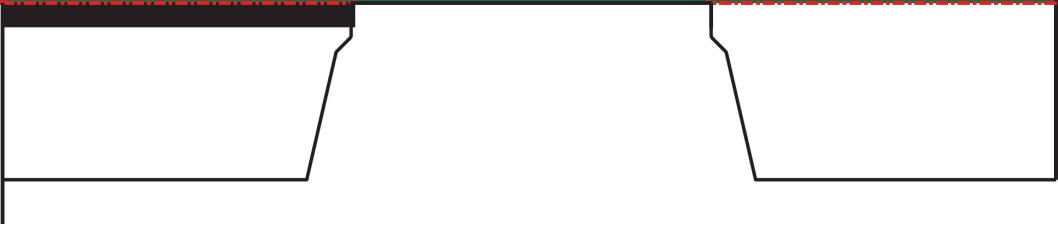
Other Information

- Protect the product in this container from excessive heat and direct sun.

Other Ingredients:

Aloe Barbadosensis Leaf (Aloe Vera Gel) Juice, Aqua (Deionized Water), Arnica Montana Flower Extract, Boswellia Serrata Extract, **Cannabis Sativa (Hemp) Seed Oil, Caprylic/Capric Triglyceride, Cetearyl Olivale, Cetyl Alcohol, Citric Acid, Gluconolactone, Glycerin, Glyceryl Stearate, Glycyrrhiza Glabra (Licorice) Extract, Ilex Paraguariensis (Yerba Mate') Extract, Magnesium Sulfate, Methylsulfonylmethane (MSM), Sodium Benzoate, Sorbitan Olivale, Xanthan Gum, Zemea (Corn) Propanediol.

Package Labeling:

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 <p>X002GV5PLB Hemp Cream (New)</p>			
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HAVASU HEMP SOOTHING PAIN RELIEF RUB

histamine dihydrochloride cream

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:79216-159

Route of Administration	TOPICAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - UNII:820484N8I3)	HISTAMINE DIHYDROCHLORIDE	0.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CETEARYL OLIVATE (UNII: 58B69Q84JO)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
GLYCYRRHIZA GLABRA (UNII: 2788Z9758H)	
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B4O4F)	
MAGNESIUM SULFATE, UNSPECIFIED FORM (UNII: DE08037SAB)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBITAN OLIVATE (UNII: MDL271E3GR)	
XANTHAN GUM (UNII: TTV12P4NEE)	
CORN (UNII: 0N8672707O)	
PROPANEDIOL (UNII: 5965N8W85T)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
WATER (UNII: 059QF0KO0R)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79216-159-01	1 in 1 BOX	05/01/2020	
1		118 mL in 1 JAR; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2020	

