HIGHPOTHECARY THERAPEUTICS CBD ZING- menthol cream
SIBORG LLC

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.

Highpothecary Therapeutics CBD Zing

Drug Facts

Active ingredient

Menthol 1.25%

Purpose

Topical Analgesic

Use

- For the temporary relief of minor aches and pains of muscles and joints.

Warnings

For external use only.

When using this product

- avoid contact with the ey es
- do not apply to wounds or damaged skin
- do not bandage tightly

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.

Directions

- Adults and children 12 years of age and older: Apply to affected area not more than 3 to 4 times daily.
- Children under 12 years of ag e: Consult a doctor

Other information

Protect this product from excessive heat and direct sunlight.

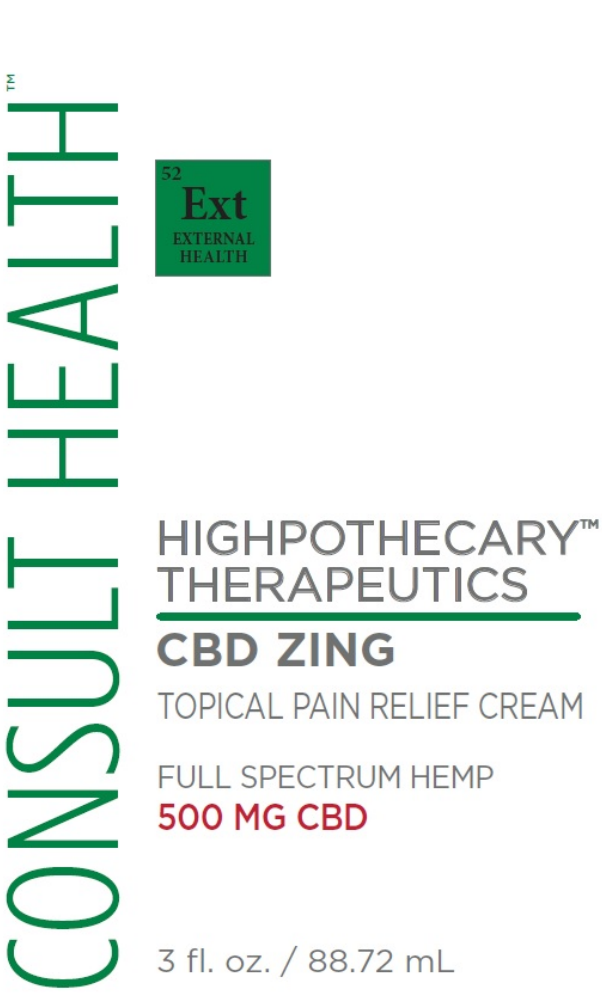
Inactive ingredients

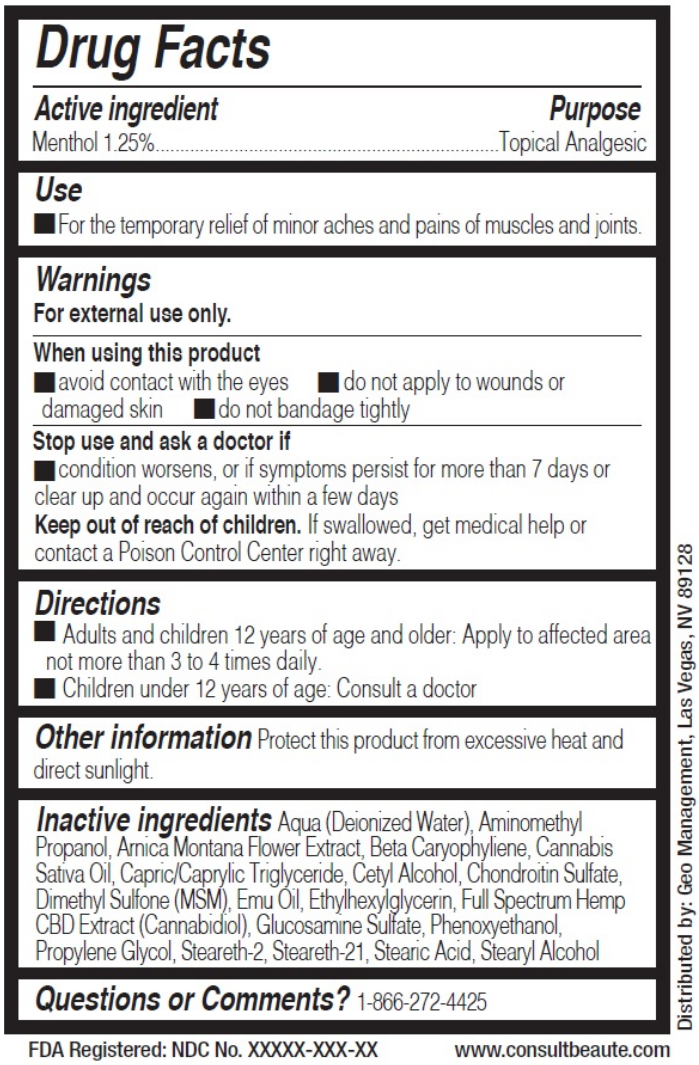
Aqua (Deionized Water), Aminomethyl Propanol, Arnica Montana Flower Extract, Beta Caryophyliene, Cannabis Sativa Oil, Capric/Caprylic Triglyceride, Cetyl Alcohol, Chondroitin Sulfate, Dimethyl Sulfone (MSM), Emu Oil, Ethylhexylglycerin, Full Spectrum Hemp CBD Extract (Cannabidiol), Glucosamine Sulfate, Phenoxyethanol, Propylene Glycol, Steareth-2, Steareth-21, Stearic Acid, Stearyl Alcohol

Questions or Comments?

1-866-272-4425

Package Labeling:





HIGHPOTHECARY THERAPEUTICS CBD ZING
menthol cream

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70803-001
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	12.5 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
AMINOMETHYLPROPANOL (UNII: LU49E6626Q)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
EMU OIL (UNII: 344821WD61)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
GLUCOSAMINE SULFATE (UNII: 1FW7WLR731)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARETH-2 (UNII: V56DFE46J5)	
STEARETH-21 (UNII: 53J3F32P58)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4H1Y)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:70803-001-88	88.72 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	02/07/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/07/2020	

Labeler - SIBORG LLC (102875148)

Registrant - Geo Management Corporation (102875148)