CRYOFREEZE ROLL-ON EXTRA STRENGTH- menthol, unspecified form gel Omax Health, 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.

Cryofreeze[™] Roll-On Extra Strength

Drug Facts

Active Ingredients

Natural Menthol, USP (8.5%)

Purpose

Topical Analgesic

Uses

Temporary relief from minor aches and pains from sore muscles and joints, backache, arthritis.

Warnings

For external use only.

Flammable: Keep out of reach of children.

Contact a doctor before using if have sensitive skin or are pregnant, breastfeeding or on any medications. If swallowed get medical help or call Poison control center immediately.

Directions

Use only as directed. Do not use on children under 12 years of age. Roll onto affected area no more than four times daily.

When using this product

Do not use with heating pad. Do not bandage. Wash hands with cold water immediately after use and do not touch eyes or mucous membranes. **Stop use and ask a doctor** if redness or irritation occurs, if condition worsens, or if pain persists for more than 7 days or clears up, then reoccurs.

Inactive Ingredients

Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Serrata

Extract, Camphor, Industrial Hemp Extract, Carbomer, Chondroitin Sulfate, Ethylhexylglycerin, Glucosamine Sulfate, Glycerin, Ilex Paraguariensis Extract, Isopropyl Alcohol, Methyl Sulfonylmethane, Peppermint Oil, Phenoxyethanol, Polysorbate 20, Propylene Glycol, Purified Water, Triethanolamine. **No Parabens.**

Other Information

Questions call 1-800-765-6691. Store in a cool dry place with the cap tightly closed. Note: Because this product contains natural ingredients, colors may vary.

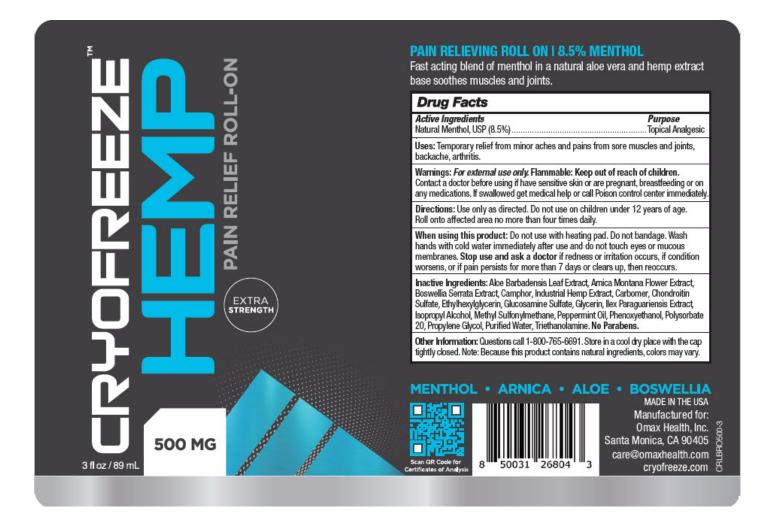
PRINCIPAL DISPLAY PANEL - 89 mL Bottle Label

CRYOFREEZE™ HEMP PAIN RELIEF ROLL-ON

EXTRA STRENGTH

3 fl oz / 89 mL

500 MG



Product Info	ormation						
Product Type		HUMAN OTC DRUG	ltem Code	(Source)	N	DC:7303	36-004
		TOPICAL		(Source)			
Route of Admi	Inistration	TOPICAL					
Active Ingre	dient/Active	Moiety					
	Ingr	edient Name		Bas	sis of Stro	ength	Strengt
MENTHOL, UNSI FORM - UNII:L7T1((UNII: L7T10EIP3A) (MENTH(DL, UNSPECIFI		THOL, PECIFIED FC	RM	85 mg in 1 mL
Inactive Ing	redients						
		Ingredient Nam	e				Strength
ALOE VERA LEA	F (UNII: ZY81Z83H	10X)					
ARNICA MONTA	NA WHOLE (UNII:	080TY208ZW)					
	ICENSE (UNII: 4P)						
CAMPHOR (SYN	THETIC) (UNII: 5T	JD82A1ET)					
HEMP (UNII: TD1I	MUT01Q7)						
CARBOMER INTI	ERPOLYMER TYP	E A (ALLYL SUCROSE CR	OSSLINKED)	(UNII: 59TL	3WG5CO)		
CHONDROITIN S	ULFATE (SHARK) (UNII: 2ZAJ1K50XH)					
GLUCOSAMINE	SULFATE POTAS	SIUM CHLORIDE (UNII: 15)	VQ11I66N)				
GLYCERIN (UNII:	PDC6A3C0OX)						
	IENSIS LEAF (UNI	· ·					
ISOPROPYL ALC	OHOL (UNII: ND2)	4416302)					
	IOL (UNII: HIE4922	- /					
	YCERIN (UNII: 147						
	ONE (UNII: 9H4PC						
	L (UNII: AV092KU4	•					
	20 (UNII: 7T1F30V						
	COL (UNII: 6DC90	(167V3)					
TROLAMINE (UNII: 903K93S3TK)							
WATER (UNII: 05	9QF0KO0R)						
Packaging							
# Item Code	Р	ackage Description			ng Start ate		eting End Date
1 NDC:73036- 004-01	89 mL in 1 BOTT Combination Proc	LE, WTH APPLICATOR; Type duct	0: Not a	03/16/202	2		
Markoting	lnformat	ion					
Marketing	,						

final	part348	03/16/2022	
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Labeler - Omax Health, Inc (965730778)

Revised: 3/2022

Omax Health, Inc