# CBD GLOBAL FREEZE- methyl salicylate, menthol, camphor spray Cosmetic Specialty Labs, 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.

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#### **Active Incredients**

Methyl Salicylate 30%
Menthol 10%
Camphor Oil 4%

# **Purpose**

Topical analgesi

#### Uses

temporarily relieves minor pain associated with: • arthritis • simple backache • muscle strains • sprains • bruises • cramps

# **Warnings**

for external use only

When using this prodcut • use only as directed • avoid contact with eyes or mucous membranes • do not apply to wounds or damaged skin •do not bandage tightly

**Stop use and ask a doctor if** • condition worsens • symptoms persist for more than 7 days or clear up and occur again within a few days • redness is present • irritation develops

If pregnant or breast-feeding, ask a health professional before use.

**Keep out of reach of children.** If swallowed, get medical help or contact Poison Control Center right away.

### **Directions:**

**Adults and children over 2 years:** • apply generously to affected area • massage into painful area until throughly absorbed into skin • repeat as necessary, but no more than 4 time daily

Children 2 years or younger: ask a doctor

# Inactive ingredients

## Package Label















220MG True Cannabinoid Content™

Spray
Net Wt. 0.8 floz / 25ml

# Drug Facts

Active Ingredients Purpose
Methyl Salicylate 30% ...Topical analgesic
Menthol 10% ...........Topical analgesic
Camphor Oil 4% .........Topical analgesic

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*Inactive Ingredients:* Industrial Hemp CBD (200mg), CB3(+0mg) ,CBH(+0mg), Isopropyl Alcohol.

NDC #58133-954-01 Varufacured for C3D G dba . LCC Golden, CD LSA Queetione ? Call 1-720-524-6369

### **CBD GLOBAL FREEZE**

methyl salicylate, menthol, camphor spray

### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58133-954

Route of Administration TOPICAL

# **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength	
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	10 g in 100 mL	
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	30 g in 100 mL	
<b>CAMPHOR (SYNTHETIC)</b> (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 mL	

<b>Inactive Ingredients</b>
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Ingredient Name	Strength	
CANNABIDIOL (UNII: 19GBJ60SN5)		

ISOPROPYL ALCOHOL (UNII: ND2M416302)

CANNABINOL (UNII: 7UYP6MC9GH)	
CANNABIGEROL (UNII: J1K406072N)	

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	NDC:58133- 954-01	25 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	09/28/2022	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	09/28/2022	

# **Labeler -** Cosmetic Specialty Labs, Inc. (032973000)

Establishment			
Name	Address	ID/FEI	Business Operations
Cosmetic Specialty Labs, Inc.		032973000	manufacture(58133-954), pack(58133-954), label(58133-954)

Revised: 9/2022 Cosmetic Specialty Labs, Inc.