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.

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 develeps

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

Industrial Hemp CBD(200mg), CBG(10mg), CBN(10mg), Isopropyl Alcohol

Package Label

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
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	4 g in 100 mL

Inactive Ingredients	
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
1	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.