CBD PAIN FREEZE SHRINK- menthol gel
Global Widget, LLC

Drug Facts

Active Ingredients

Menthol USP 4%

Purpose

Topical Analgesic

Uses

For the temporary relief of pain

Warnings

FOR EXTERNAL USE ONLY

Do not use:

- On eyes or on mucous membranes
- On wounds, damaged or irritated skin
- If you are allergic to Menthol or any of the ingredients listed below

When using this product:

- Use only as directed
- Do not bandage or cover with any type of wrap except clothing
- Do not use with heating pad or devices, or apply external heat

Stop use and ask a doctor if

- Localized skin reactions occur, such as rash, itching, redness, irritation, pain, swelling and blistering
- Conditions worsen
- Symptoms persist for more than 7 days or clear up and occur again within a few days

If pregnant or breastfeeding: Do not use this product.

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

Directions

• Adults 18 years & over rub a thin layer into affected areas up to 4 times daily.

• ***Wash hands after application.***

Other Information:

Store in cool, dry place away from direct sunlight

Do not use if seal is broken or not present

Inactive Ingredients

Water, Isopropyl Alcohol, Carbomer, Pure CBD Extract, Triethanolamine

Distributed by

Global Widget, LLC

8419 Sunstate Street, Tampa, FL 33634

Principal Display Panel

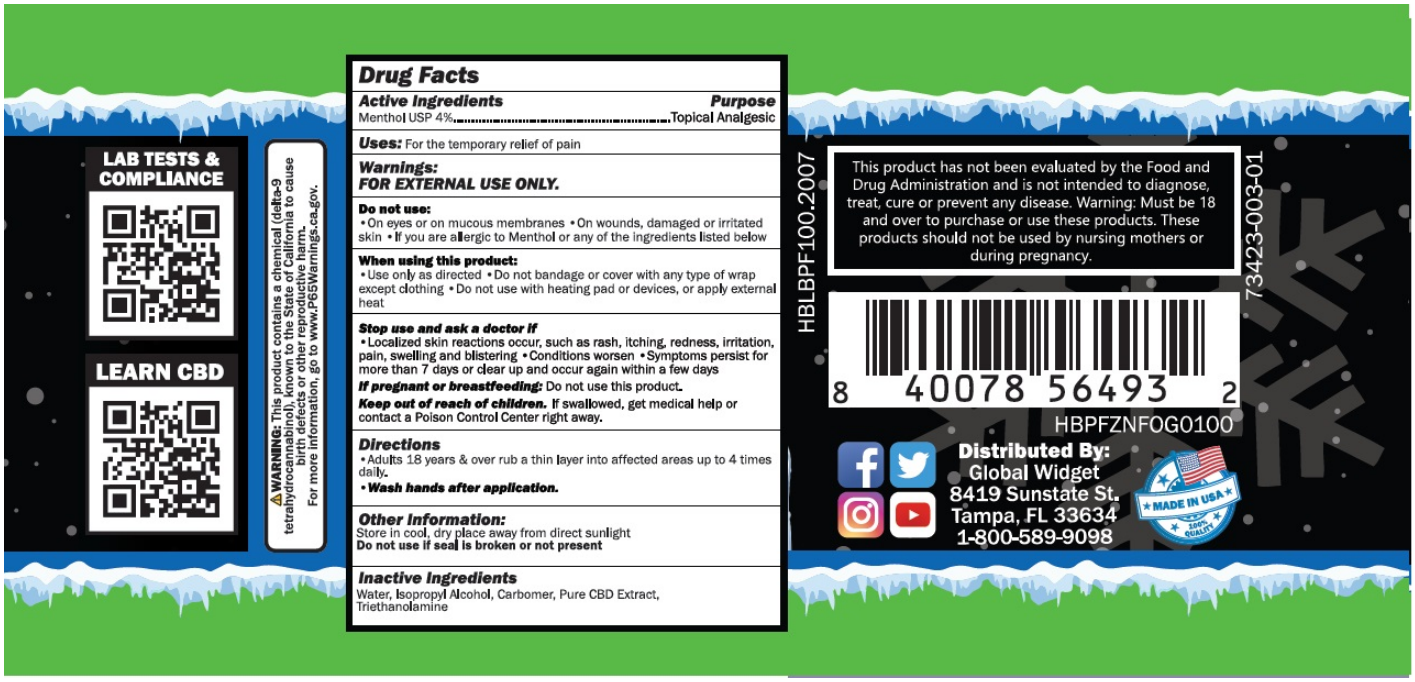
HEMP

BOMBS

CBD Pain Freeze Shrink

MENTHOL 4%

1 OZ (28 g)



CBD PAIN FREEZE SHRINK

menthol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:73423-003
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	40 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73423-003-01	12 in 1 PACKAGE	12/15/2020	09/30/2025
1		28 g in 1 BOTTLE; 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	12/15/2020	09/30/2025

Labeler - Global Widget, LLC (089584863)

Establishment

Name	Address	ID/FEI	Business Operations
Global Widget, LLC		089584863	manufacture(73423-003)

Establishment

Name	Address	ID/FEI	Business Operations
Global Widget LLC		118504011	manufacture(73423-003)

Revised: 12/2023 Global Widget, LLC