

**CBD CRYOTHERAPY PAIN RELIEF- menthol gel**  
**Global Widget, LLC**

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***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**

**NATURE'S**

**SCRIPT**

**CBD Cryotherapy Pain Relief Gel**

**MENTHOL 4%**

1 OZ (28 g)

# PREMIUM CBD COLD THERAPY



## NATURE'S SCRIPT

# CBD CRYOTHERAPY PAIN RELIEF GEL

## 100MG CBD PER BOTTLE

Net Wt. 1oz (28g)

**WARNING:** This product contains a chemical (delta-9 tetrahydrocannabinol) known to the State of California to cause birth defects or other reproductive harm. For more information, go to [www.P65Warnings.ca.gov](http://www.P65Warnings.ca.gov).



**DISTRIBUTED BY:**  
GLOBAL WIDGET  
8419 SUNSTATE ST.  
TAMPA, FL 33634  
(800) 713-6405

### Drug Facts

Active Ingredients	Purpose
Menthol USP 4%	Topical Analgesic
<b>Uses:</b> For the temporary relief of pain	
<b>Warnings:</b> <b>FOR EXTERNAL USE ONLY.</b>	
<b>Do not use:</b> • 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	
<b>When using this product:</b> • 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	
<b>Stop use and ask a doctor if</b> • 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	
<b>If pregnant or breastfeeding:</b> Do not use this product.	
<b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away.	
<b>Directions</b> • Adults 18 years & over rub a thin layer into affected areas up to 4 times daily. • Wash hands after application,	
<b>Other Information:</b> Store in cool, dry place away from direct sunlight Do not use if seal is broken or not present	
<b>Inactive Ingredients</b> Water, Isopropyl Alcohol, Carbomer, Pure CBD Extract, Triethanolamine	

This product contains a total delta-9 tetrahydrocannabinol concentration that does not exceed 0.3% on a dry weight basis.

### PREMIUM • TRUSTED • PURE

This product has not been evaluated by the Food and Drug Administration and is not intended to diagnose, treat, cure or prevent any disease. Warning: Must be 18 and over to purchase or use these products. These products should not be used by nursing mothers or during pregnancy.



NSPFZNF0G0100



73423-002-01

NSLBPF100.2007

[NATURESCRIPT.COM](http://NATURESCRIPT.COM)

## CBD CRYOTHERAPY PAIN RELIEF

menthol gel

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73423-002
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:73423-002-01	12 in 1 PACKAGE	12/17/2020	04/30/2025
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	12/17/2020	04/30/2025

**Labeler** - Global Widget, LLC (089584863)**Establishment**

Name	Address	ID/FEI	Business Operations
Global Widget, LLC		089584863	manufacture(73423-002)

**Establishment**

Name	Address	ID/FEI	Business Operations
Global Widget LLC		118504011	manufacture(73423-002)

Revised: 12/2023

Global Widget, LLC