

**CBD PAIN RELIEF CLINICAL STRENGTH MENTHOL FORMULA- menthol,
unspecified form gel
Global Products Group, LLC**

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.

**CBD Pain Relief
Clinical Strength Menthol Formula**

Drug Facts

Active Ingredient

Natural Menthol, USP (8.5%)

Purpose

Topical Analgesic

Uses

Temporarily relieves foot, ankle and leg pain associated with arthritis

- muscle aches
- muscle strains
- muscle sprains
- joint pain.

Warnings

For external use only: Flammable: Keep away from excessive heat or open flame

- Ask a doctor before use if you have sensitive skin or if you are taking any blood thinners
- **When using this product:** Do not use on wounds or irritated skin
- Do not bandage tightly or use with a heating pad
- Wash hands after use with cool water
- **If pregnant or breastfeeding,** ask a health professional before use
- **Keep out of reach of children.** If accidentally swallowed, contact a doctor or poison control center immediately
- **Stop use and ask a doctor:** If condition worsens or if pain persists for more than 7 days, or clears up, then reoccurs within a few days.

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
- Shake well before each use.

Inactive Ingredients

Aloe Barbadensis Leaf Extract, Arnica Montana Flower Extract, Boswellia Serrata Extract, Camphor, Carbomer, Chondroitin Sulfate, Ethylhexylglycerin, Glucosamine Sulfate, Glycerin, Ilex Paraguariensis Extract, Isopropyl Alcohol, Methyl Sulfonylmethane, Peppermint Oil, Phenoxyethanol, Polysorbate 20, Propylene Glycol, Purified Water, Triethanolamine

Other Information

Questions or comments?

Call (877) 383-2334. Store in a cool dry place with the cap tightly closed. Note: Because this product contains natural ingredients, color may vary.

DISTRIBUTED BY

HOTWORX®
24 HOUR INFRARED FITNESS STUDIO

planet beach®
spray & spa

PRINCIPAL DISPLAY PANEL - 88 mL Bottle Label

ROLL ON

CBD
PAIN RELIEF

Clinical Strength Menthol Formula

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HOTWORX®
24 HOUR INFRARED FITNESS STUDIO

planet beach®
spray & spa

150mg CBD 3 fl oz (88 mL)

ROLL ON
CBD
 PAIN RELIEF

Clinical Strength Menthol Formula

DISTRIBUTED BY



150mg CBD 3 fl oz (88 mL)

Pain relief for arthritis, joints and muscles. Contains Glucosamine, Chondroitin, Arnica, Boswellia. THC Free. Shake well before each use. Broad spectrum CBD.

Drug Facts

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Purpose

Uses: Temporarily relieves foot, ankle and leg pain associated with arthritis • muscle aches • muscle strains • muscle sprains • joint pain.

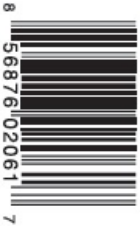
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Manufactured for
 Global Products Group
 13760 Repton Blvd.
 Tampa, FL 33626



CBD PAIN RELIEF CLINICAL STRENGTH MENTHOL FORMULA

menthol, unspecified form gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72593-157
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	85 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA WHOLE (UNII: O80TY208ZW)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CARBOMER INTERPOLYMER TYPE A (ALLYL SUCROSE CROSSLINKED) (UNII: 59TL3WG5CO)	

CHONDROITIN SULFATE (SHARK) (UNII: 2ZAJ1K50XH)
GLUCOSAMINE SULFATE POTASSIUM CHLORIDE (UNII: 15VQ11I66N)
GLYCERIN (UNII: PDC6A3C0OX)
ILEX PARAGUARIENSIS LEAF (UNII: 1Q953B404F)
ISOPROPYL ALCOHOL (UNII: ND2M416302)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)
PEPPERMINT OIL (UNII: AV092KU4JH)
POLYSORBATE 20 (UNII: 7T1F30V5YH)
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)
TROLAMINE (UNII: 9O3K93S3TK)
WATER (UNII: 059QF0KO0R)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72593-157-12	88 mL in 1 BOTTLE, WITH APPLICATOR; Type 0: Not a Combination Product	10/05/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH NOT FINAL	part348	10/05/2020	

Labeler - Global Products Group, LLC (081371764)

Revised: 1/2022

Global Products Group, LLC