

JUST CBD - CBD AND THC ULTRA RELIEF- menthol, camphor gel
Just Brands LLC

73647-062-04

73647-026-04

Camphor 2%

Menthol 6%

Topical Analgesic.

USES:

Aid for temporary local relief of minor pain in muscles or joints.

For external use only.

Use only as directed.

Do not bandage tightly or use with a heating pad.

Avoid contact with eyes and mucous membranes.

Do not apply to wounds or damaged, broken, or irritated skin.

A transient burning sensation or redness may occur upon application but generally disappears in several days.

If you experience an allergic reaction, discontinue use, and consult a doctor.

Do not expose the area treated with product to heat or direct sunlight.

STOP USE AND ASK A DOCTOR IF:

Condition worsens.

Redness is present.

Irritation develops.

Symptoms persist for more than 7 days or clear up occur again within a few days.

You experience signs injury, such as pain, swelling or blistering where the product was applied.

Ask a health professional before use.

If swallowed, get medical help or contact a Poison Control Center right away.

DIRECTIONS:

Adults over 21 years:

Apply a small amount on the affected area.

Massage in circular motion, let set for a few seconds.

Repeat as necessary, but no more than 3 to 4 times daily

Wash hands with soap and water after use.

Store tightly closed in a dry place at controlled room temperature between 59°-86° f

(15°-30° c). This product is intended for use by healthy adults aged 21 years & older. consult a healthcare professional prior to use of full spectrum THC. Full spectrum THC may be harmful if you are pregnant, nursing or are taking any medication or have a medical condition.

Water (Aqua), Alcohol Denat, Propylene Glycol, Polysorbate 20, Salicylic Acid, Glycerin, Carbomer, Sodium Hydroxide, Mentha Piperita (Peppermint) Oil, Cannabidiol, Rosmarinus Officinalis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Delta-8-Tetrahydrocannabinol, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Benzyl Alcohol, Sorbic Acid, FD&C Blue No.1 (CI 42090), FD&C Yellow No.5 (CI 19140), Limonene.

JUST CBD - CBD+THC ULTRA RELIEF GEL +1000

Drug Facts

Active ingredient	Purpose
Camphor 2%	Topical Analgesic
Menthol 6%	Topical Analgesic

Use
Aid for temporary local relief of minor pain in muscles or joints.

Warnings
For external use only.

When using this product

- Use only as directed. • Do not bandage tightly or use with a heating pad. • Avoid contact with eyes and mucous membranes.
- Do not apply to wounds or damaged, broken, or irritated skin. • A transient burning sensation or redness may occur upon application but generally disappears in several days. • If you experience an allergic reaction, discontinue use, and consult a doctor. • Do not expose the area treated with product to heat or direct sunlight.

Stop use and ask a doctor if

- Condition worsens. • Redness is present. • Irritation develops. • Symptoms persist for more than 7 days or clear up occur again within a few days. • You experience signs injury, such as pain, swelling or blistering where the product was applied.

If pregnant or breast – feeding
Ask a health professional before use.

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

Directions
Adults over 21 years

- Apply a small amount on the affected area.
- Massage in circular motion, let set for a few seconds. • Repeat as necessary, but no more than 3 to 4 times daily. • Wash hands with soap and water after use.

Other information
Store tightly closed in a dry place at controlled room temperature between 59°-86° F (15°-30° C). This product is intended for use by healthy adults aged 21 years & older, consult a healthcare professional prior to use of full spectrum THC. Full spectrum THC may be harmful if you are pregnant, nursing or are taking any medication or have a medical condition.

Inactive Ingredients
Water (Aqua), Alcohol Denat, Propylene Glycol, Polysorbate 20, Salicylic Acid, Glycerin, Carbomer, Sodium Hydroxide, Mentha Piperita (Peppermint) Oil, Cannabidiol, Rosmarinus Officinalis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Delta-8-Tetrahydrocannabinol, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Benzyl Alcohol, Sorbic Acid, FD&C Blue No.1 (CI 42090), FD&C Yellow No.5 (CI 19140), Limonene.

JUSTCBD™ Full Spectrum (CBD & THC) Ultra Relief Gel is infused with USA grown Hemp derived CBD. It helps ease areas of discomfort that need it most.

- The FDA has not evaluated this product for safety or efficacy, there is no current standardized methodology for verifying full spectrum THC content as required by Florida state law. Full spectrum THC potency is verified by an independent third-party lab. The results of the independent third-party lab tests are available to you on our website and by scanning the QR code on the package. Test results may vary by individual laboratory and/or test method. This product contains less than 0.3% total delta-9 tetrahydrocannabinol (delta-9 THC), drug tests vary in sensitivity, and you may test positive for delta-9 THC when taking full spectrum THC products. Over time color variations may occur in this product, the amount (mg) of full spectrum THC may vary by dose and by packaging. If full spectrum THC is used beyond the expiration date, potency may decrease. May cause drowsiness, do not drive, or operate heavy machinery while taking full spectrum THC.

• This product is not intended to diagnose, treat, cure, or prevent any disease.

JUSTCBDSTORE.COM

Distributed by:
JUST BRANDS LLC
3406 SW 25TH Ter Ste 1
Fort Lauderdale, FL 33312
1-833-458-7822
NDC# 73647-062-04

JUST CBD - CBD+THC ULTRA RELIEF GEL +5000

INDUSTRIAL HEMP

Drug Facts

Active ingredient Purpose
 Camphor 2% Topical Analgesic
 Menthol 6% Topical Analgesic

Use
 Aid for temporary local relief of minor pain in muscles or joints.

Warnings
 For external use only.

When using this product
 • Use only as directed. • Do not bandage tightly or use with a heating pad. • Avoid contact with eyes and mucous membranes. • Do not apply to wounds or damaged, broken, or irritated skin. • A transient burning sensation or redness may occur upon application but generally disappears in several days. • If you experience an allergic reaction, discontinue use, and consult a doctor. • Do not expose the area treated with product to heat or direct sunlight.

Stop use and ask a doctor if
 • Condition worsens. • Redness is present. • Irritation develops.
 • Symptoms persist for more than 7 days or clear up occur again within a few days. • You experience signs injury, such as pain, swelling or blistering where the product was applied.

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

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

Directions
Adults over 21 years
 • Apply a small amount on the affected area.
 • Massage in circular motion. Let set for a few seconds. • Repeat as necessary, but no more than 3 to 4 times daily. • Wash hands with soap and water after use.

Other information
 Store tightly closed in a dry place at controlled room temperature between 59°-86° F (15°-30° C). This product is intended only for use by healthy adults aged 21 years & older. Consult a healthcare professional prior to use of full spectrum THC. Full spectrum THC may be harmful if you are pregnant, nursing or are taking any medication or have a medical condition.

Inactive Ingredients
 Water (Aqua) - Alcohol Denat., Propylene Glycol, Polysorbate 20, Salicylic Acid, Glycerin, Carbomer, Sodium Hydroxide, Mentha Piperita (Peppermint) - Oil, Cannabidiol, Rosmarinus Officinalis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Delta-8-Tetrahydrocannabinol, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Benzyl Alcohol, Sorbic Acid, FD&C Blue No.1 (CI 42090), FD&C Yellow No.5 (CI 19140), Limonene.

relief

JUSTCBD™ Full Spectrum (CBD & THC) Ultra Relief Gel is infused with USA grown Hemp derived CBD. It helps ease areas of discomfort that need it most.

• The FDA has not evaluated this product for safety or efficacy, there is no current standardized methodology for verifying full spectrum THC content as required by Florida state law. Full spectrum THC potency is verified by an independent third-party lab. The results of the independent third-party lab tests are available to you on our website and by scanning the QR code on the package. Test results may vary by individual laboratory and/or test method. This product contains less than 0.3% total delta 9 tetrahydrocannabinol (delta-9 THC), drug tests vary in sensitivity, and you may test positive for delta 9 THC when taking full spectrum THC products. Over time color variations may occur in this product, the amount (mg) of full spectrum THC may vary by dose and by packaging. If full spectrum THC is used beyond the expiration date, potency may decrease. May cause drowsiness, do not drive, or operate heavy machinery while taking full spectrum THC.

• This product is not intended to diagnose, treat, cure, or prevent any disease.

JUSTCBDSTORE.COM

Scan for 3rd Party Lab Test Results

Distributed by:
 JUST BRANDS LLC.
 3406 SW 26TH Ter Ste 1
 Fort Lauderdale, FL 33312
 1-833-458-7822
 NDC# 73647-026-04

TOPICAL **4 OZ (113g)**

JUST CBD - CBD AND THC ULTRA RELIEF

menthol, camphor gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73647-062
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 g in 100 g
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
PEPPERMINT OIL (UNII: AV092KU4JH)	

ROSEMARY OIL (UNII: 8LGU7VM393)	
THYME OIL (UNII: 2UK410MY6B)	
TEA TREE OIL (UNII: VIF565UC2G)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SORBIC ACID (UNII: X045WJ989B)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
.DELTA.8-TETRAHYDROCANNABINOL (UNII: B49D0HH807)	

Product Characteristics

Color	turquoise	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73647-062-04	113 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/14/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/14/2023	

JUST CBD - CBD AND THC ULTRA RELIEF

menthol, camphor gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73647-026
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
-----------------	-------------------	----------

MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	6 g in 100 g
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET) (CAMPHOR (SYNTHETIC) - UNII:5TJD82A1ET)	CAMPHOR (SYNTHETIC)	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CARBOMER 940 (UNII: 4Q93RCW27E)	
WATER (UNII: 059QF0KO0R)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MENTHA ARVENSIS LEAF OIL (UNII: 1AEY1M553N)	
ROSEMARY OIL (UNII: 8LGU7VM393)	
THYME OIL (UNII: 2UK410MY6B)	
TEA TREE OIL (UNII: VIF565UC2G)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
SORBIC ACID (UNII: X045WJ989B)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
ALCOHOL (UNII: 3K9958V90M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SALICYLIC ACID (UNII: O414PZ4LPZ)	
GLYCERIN (UNII: PDC6A3C0OX)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
CANNABIDIOL (UNII: 19GBJ60SN5)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
.DELTA.8-TETRAHYDROCANNABINOL (UNII: B49D0HH807)	

Product Characteristics

Color	turquoise	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73647-026-04	113 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	03/14/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	03/14/2023	

