

**GREEN LEAF CBD ULTRA RELIEF- menthol, camphor gel**  
**SURF LINE, 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.*

-----  
**83345-602-04**

Menthol 6%

Camphor 2%

Topical Analgesic.

Pain Relieving.

**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.

Ask a health professional before use.

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

**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.

**DIRECTIONS:**

Adults and Children over 12 years:

Apply a small amount on the area to be.

Massage in circular motion until absorbed.

Repeat as needed, but no more than 3 to 4 times per day.  
Store tightly closed in a dry place at room temperature between 59°-86° F (15°-30° C).  
Wash hands with soap and water after use.

Aqua, Alcohol Denat, Propylene Glycol, Polysorbate 20, Salicylic Acid, Glycerin, Carbomer, Sodium Hydroxide, Mentha Arvensis Herb 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. 1 Powder (CI 42090), FD&C Yellow. 5 (CI 19140), Limonene.

## GREEN LEAF CBD ULTRA RELIEF GEL

**DRUG FACTS**

**ACTIVE INGREDIENTS:**  
Ibuprofen ..... 6% Topical Analgesic  
Camphor ..... 2% Pain Relieving

**USERS:** Relief for temporary local relief in muscle or joint.

**WARNINGS:** For external use only.  
• Do not use if 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.

**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.

**STOP USE AND ASK A DOCTOR IF:**  
• Condition worsens.  
• Redness is present.  
• Irritation develops.  
• Symptoms persist for more than 7 days or flare up occur again within a few days.  
• You experience signs of injury, such as pain, swelling, or blistering where the product was applied.

**DIRECTIONS:**  
**Adults and Children over 12 years:**  
• Apply a small amount on the area to be treated.  
• Massage in circular motion until absorbed.  
• Repeat as needed, but no more than 3 to 4 times per day.  
• Store tightly closed in a dry place at room temperature between 59°-86° F (15°-30° C).  
• Wash hands with soap and water after use.

**INACTIVE INGREDIENTS:** Aqua, Alcohol Denat., Propylene Glycol, Polysorbate 20, Salicylic Acid, Glycerin, Carbomer, Sodium Hydroxide, Mentha Arvensis Herb Oil, Rosmarinus Officinalis (Rosemary) Oil, Thymus Vulgaris (Thyme) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Cannabidiol, Delta-8-Tetrahydrocannabinol, Benzyl Alcohol, Sorbic Acid, FD&C Blue. 1 Powder (CI 42090), FD&C Yellow. 5 (CI 19140), Limonene.

NDC# 83345-602-04

**ULTRA RELIEF GEL**

With MENTHOL

**1000 CBD + 100 THC**

**TOPICAL**

**NET WT. 4oz / 118g**

**INDUSTRIAL HEMP**

**Green Leaf CBD**  
Ultra Relief Gel is formulated with CBD extracted from USA-grown hemp. It works to **alleviate** discomfort in areas that require it most.

**\*CAUTION:** This product is for external use only. Do not use it on damaged skin, wounds, eyes, or mucous membranes. Do not exceed the recommended amount. It is intended for use by healthy adults who are 18 years or older. Keep it out of reach of children. Cannabidiol (CBD) use during pregnancy, while breastfeeding, while taking any medication, or if you have any medical condition may be harmful. Prior to use, consult a healthcare professional. Discontinue use immediately if you experience any adverse reactions and consult your physician. This product has not been evaluated for safety or efficacy by the FDA. There is no standardized CBD testing methodology to verify CBD content. Our third-party lab product test reports, which are the basis for the CBD amount, are available on our website and by scanning the QR code on the package, but the test results may vary depending on the individual laboratory and/or test method. The amount of mg/CBD may differ depending on the dosage and the package. This product contains a total delta-9 THC that does not exceed 0.3%. The sensitivity of drug tests varies, and CBD products may cause a positive THC result. Natural color variations may occur. May cause drowsiness.

**\*The Food and Drug Administration (FDA) has not evaluated these statements. This product is not intended to diagnose, treat, cure, or prevent any disease.\***

GreenLeafCBD.com  
Less Than 0.3% THC. #MADE IN USA

Distributed by:  
**GreenLeafCBD**  
3004 N. Kings Hwy,  
Harris Ranch, NC 28577  
(843) 663-0437

## GREEN LEAF CBD ULTRA RELIEF

menthol, camphor gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83345-602
<b>Route of Administration</b>	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:83345-602-04	118 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	04/17/2023	
Marketing Information				
Marketing Category		Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final		part348	04/17/2023	

**Labeler** - SURF LINE, INC (091171162)

Revised: 4/2023

SURF LINE, INC