ULTRA RELIEF- menthol, camphor gel
CHEMCO CORPORATION

49283-578-04

Camphor 3%

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 neccesary, 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, Glycerin, Carbomer, Sodium Hydroxide, Mentha Piperita (Peppermint) Oil, Rosmarinus Officinalis (Rosemary) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Thymus Vulgaris (Thyme) Oil, Cannabis Sativa Seed Oil, Benzyl Alcohol, Salicylic Acid, Sorbic Acid, FD&C Yellow No.5 (CI 19140), FD&C Blue No.1 (CI 42090).

ULTRA RELIEF GEL

ULTRA RELIEF

MENTHOL 6% CAMPHOR 3%

Drug Facts

Active ingredients — ***Purpose***

Camphor 3% ..... Topical Analgesic

Menthol 6%..... Topical Analgesic

Use

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

Warnings

For external use only.

Ask a doctor before use if you have redness over affected area.

When using this product

- Use only as directed.
- Avoid contact with eyes and mucous membranes.
- Do not bandage tightly or use with a heating pad.
- 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 physician.
- Do not apply on large areas of the body or on damaged, broken, or irritated skin.
- Do not expose the area treated with product to heat or direct sunlight.

Stop use and ask a doctor if

- Redness is present.
- Condition worsens.
- Irritation develops.
- Symptoms persist for more than 7 days or clear up and 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 and children over 12 years of age

- Apply a small amount to the affected area.
- Massage in a 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.

Children under 12 years of age:do not use unless directed by doctor.

Other information

Store tightly closed in a dry place at controlled room temperature between 59°-86°F 15° - 30°C

Inactive ingredients

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

ULTRA RELIEF

menthol, camphor gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:49283-578
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)	3 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
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)	

Product Characteristics

Color	turquoise	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-578-04	113 g in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/15/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/15/2026	

Labeler - CHEMCO CORPORATION (032495954)

Revised: 1/2026

CHEMCO CORPORATION