

MENTHOL, CAMPHOR- menthol, camphor cream
Chemco Corporation

49283-576-02

Menthol 10%
Camphor 3%

External Analgesic
Pain Relieving

Uses

For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, sprains and strains.

Use only as directed. For external use only. Do not bandage tightly. Do not use with heating pad, pack, wrap, hot water bottle or any heating element. Keep away from children. Package not child resistant. In case of accidental ingestion, contact doctor immediately. If prone to allergic reaction to the product, consult to a doctor before using.

Avoid contact with eyes, mucous membranes, rashes, wounds or damaged skin. Do not apply on nose and genital area. If rash, redness or itchiness results; discontinue use and consult a doctor.

Keep away from children. Package not child resistant. In case of accidental ingestion, contact doctor immediately.

Directions

Adults and children over 12 years of age;

- apply a thin layer to affected area and rub gently not more than 3 to 4 times a day.
- Wash hands with soap and water after use.

Children under 12 years of age; do not use unless directed by doctor/physician

Other information

Store at room temperature 15° - 30°C (59° - 86°F)

Inactive ingredients

Water (Aqua), Alcohol Denat, Stearic Acid, Paraffinum Liquidum, Cetearyl Alcohol, Polysorbate 60, Cetyl Alcohol, Dimethicone, Glyceryl Stearate, Acrylamide/Sodium Acrylate Copolymer, Trideceth-6, Glycereth-26, Tocopheryl Acetate, Caprylic/Capric

Triglyceride, Propylene Glycol, Diazolidinyl Urea, Stearyl Alcohol, Polysorbate 20, Triethanolamine, Cannabis Sativa Seed Oil, Sodium Hyaluronate, Sodium PCA, Wheat Amino Acids, Panthenol, Symphytum Officinale (Comfrey) Extract, Hydroxyproline, Benzotriazolyl Dodecyl p-Cresol, Methylparaben, Propylparaben, FD&C YELLOW No.5 (CI 19140), FD&C Blue No.1 (CI 42090).

PAIN RELIEF CREAM

CAMPHOR MENTHOL

Drug Facts
Active ingredients
Purpose
Camphor 3% External Analgesic
Menthol 10% External Analgesic
Uses
For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, sprains and strains.
Warnings
<ul style="list-style-type: none">For external use only.Use only as directed.Do not bandage tightly.Do not use with heating pad, pack, wrap, hot water bottle or any heating element.
Ask a doctor before use if you have redness over affected area.
When using this product
<ul style="list-style-type: none">Avoid contact with eyes, mucous membranes, rashes.Do not apply to wounds or damaged skin.If condition worsens, or if symptoms persist for more than 7 days or clear up and occur again within a few days, discontinue use of this product, and consult a doctor.
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 <ul style="list-style-type: none">apply a thin layer to the affected area and rub gently not 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
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MENTHOL, CAMPHOR

menthol, camphor cream

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49283-576
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	10 g in 100 g

CAMPBOR (NATURAL) (UNII: N20HL7Q941) (CAMPBOR (NATURAL) - UNII:N20HL7Q941)

CAMPBOR (NATURAL)

3 g
in 100 g

Inactive Ingredients

Ingredient Name	Strength
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
SODIUM COCOYL WHEAT AMINO ACIDS (UNII: JW3VT57I11)	
WATER (UNII: 059QF0KO0R)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
GLYCERETH-26 (UNII: NNE56F2N14)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
TRIBUTYL CITRATE (UNII: 827D5B1B6S)	
BUTETH-3 (UNII: OC116GRO69)	
PANTHENOL (UNII: WW9CM0O67Z)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
HYALURONATE SODIUM (UNII: YSE9PPT4TH)	
DIMETHICONE 1000 (UNII: MCU2324216)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYPROLINE (UNII: RMB44WO89X)	
SYMPHYTUM OFFICINALE WHOLE (UNII: H8FJJ6KX5Y)	
TRIDECETH-6 (UNII: 3T5PCR2H0C)	
SODIUM BENZOTRIAZOLYL BUTYLPHENOL SULFONATE (UNII: 0LA2QC9O3Z)	
CETOSTEARYL ALCOHOL (UNII: 2DMT128M1S)	
GLYCERYL STEARATE SE (UNII: FCZ5MH785I)	
ALCOHOL 95% (UNII: 7528N5H79B)	
ACRYLAMIDE (UNII: 20R035KLCI)	
MINERAL OIL (UNII: T5L8T28FGP)	
.ALPHA.-TOCOPHEROL ACETATE, D- (UNII: A7E6112E4N)	
CAPRYLIC/CAPRIC/LAURIC TRIGLYCERIDE (UNII: FJ1H6M2JG9)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	

Product Characteristics

Color	green	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49283-576-02	57 g in 1 JAR; Type 0: Not a Combination Product	12/20/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	12/20/2025	

Labeler - Chemco Corporation (032495954)

Revised: 12/2025

Chemco Corporation