

LIGHTNING PAIN RELIEF- menthol cream
Clinical Resolution Laboratory, Inc.

Lightning Pain Relief Cream

DRUG FACTS

Active Ingredient

Menthol 4%

Purpose

Topical Analgesic

Uses:

- For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises, and sprains.
- Provides cooling and warming pain relief.

Warnings

- For external use only.
- Avoid contact with the eyes.
- 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.
- Do not apply to wounds or damaged skin.
- Do not bandage tightly.

Keep out of reach of children.

- In case of accidental overdose or ingestion, call a doctor or poison control center immediately.

Do not use this product

if seal is broken or missing.

Directions

- Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age: consult a doctor.

Other Information

- Keep away from direct sunlight or heat.
- Store at room temperature 15°-30° C (59°-86°F).

Inactive Ingredients

Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Aloe Barbadensis Leaf Juice, Arnica Montana Flower Extract, Boswellia Serrata Resin Extract, Butyrospermum Parkii (Shea) Butter, C13-14 Isoparaffin, Cannabis Sativa Seed Oil, Carbomer, Centella Asiatica Leaf Extract, Cinnamomum Camphora (Camphor) Bark Oil, Dimethicone, Dimethyl Isosorbide, Dipotassium Glycyrrhizate, Disodium EDTA, Emu Oil, Ethoxydiglycol, Eucalyptus Globulus Leaf Oil, Glycerin, Glyceryl Stearate, Helianthus Annuus (Sunflower) Seed Oil, Isopropyl Myristate, Laureth-7, Lavandula Angustifolia (Lavender) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methyl Sulfonyl Methane (MSM), PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Polysorbate 20, Pyridoxine HCL, Sodium Benzoate, Tocopheryl Acetate, Triethanolamine, Water

Package Labeling:

 <p><i>Ebabel</i> LIGHTNING PAIN RELIEF TOPICAL ANALGESIC CREAM</p> <p>temporary relief of minor aches and pains of muscles and joints</p> <p>4% MENTHOL</p> <p>Enhanced with inactive ingredients: Arnica, Emu Oil, Vitamin B6 & MSM</p> <p>2.28 oz e 65g</p>	<p>DRUG FACTS</p> <p>Active Ingredient Menthol 4%</p> <p>Purpose Topical Analgesic</p> <p>Uses: ■ For the temporary relief of minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises, and sprains. ■ Provides cooling and warming pain relief.</p> <p>Warnings</p> <p>■ For external use only. ■ Avoid contact with the eyes. ■ 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 the product and consult a doctor. ■ Do not apply to wounds or damaged skin. ■ Do not bandage tightly.</p> <p>Directions Adults and children 2 years of age and older: Apply to affected area not more than 3 to 4 times daily. Children under 2 years of age: consult a doctor.</p>	<p>DRUG FACTS (Continued)</p> <p>Other Information</p> <p>■ Keep away from direct sunlight or heat. ■ Store at room temperature 15°-30°C (59°-86°F). ■ This package is child-resistant. Keep out of reach of children. ■ In case of accidental overdose or ingestion, call a doctor or poison control center immediately. ■ Do not use this product if seal is broken or missing.</p> <p>Inactive Ingredients Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Allantoin, Aloe Barbadensis Leaf Juice, Arnica Montana Flower Extract, Boswellia Serrata Resin Extract, Butyrospermum Parkii (Shea) Butter, C13-14 Isoparaffin, Cannabis Sativa Seed Oil, Carbomer, Centella Asiatica Leaf Extract, Cinnamomum Camphora (Camphor) Bark Oil, Dimethicone, Dimethyl Isosorbide, Dipotassium Glycyrrhizate, Disodium EDTA, Emu Oil, Ethoxydiglycol, Eucalyptus Globulus Leaf Oil, Glycerin, Glyceryl Stearate, Helianthus Annuus (Sunflower) Seed Oil, Isopropyl Myristate, Laureth-7, Lavandula Angustifolia (Lavender) Oil, Melaleuca Alternifolia (Tea Tree) Leaf Oil, Methyl Sulfonyl Methane (MSM), PEG-100 Stearate, Phenoxyethanol, Polyacrylamide, Polysorbate 20, Pyridoxine HCL, Sodium Benzoate, Tocopheryl Acetate, Triethanolamine, Water</p>
	<p>8 50011 66802 3</p> <p>www.shang.com Made in USA</p>	

Package Labeling:101g

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<p>DRUG FACTS</p> <table border="1"> <tr> <td>Active Ingredient</td> <td>Purpose</td> </tr> <tr> <td>Menthol 4%</td> <td>Topical Analgesic</td> </tr> </table>		Active Ingredient	Purpose	Menthol 4%	Topical Analgesic
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Manufactured for Ebanel Laboratories, Inc.
Brea, CA 92821 | 1-844-694-0004
www.ebanel.com | Made in USA

LIGHTNING PAIN RELIEF			
menthol cream			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63742-013
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	40 mg in 1 g
Inactive Ingredients			

Ingredient Name	Strength
ALLANTOIN (UNII: 344S277G0Z)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ARNICA MONTANA FLOWER (UNII: OZ0E5Y15PZ)	
INDIAN FRANKINCENSE (UNII: 4PW41QCO2M)	
SHEA BUTTER (UNII: K49155WL9Y)	
C13-14 ISOPARAFFIN (UNII: E4F12ROE70)	
CANNABIS SATIVA SEED OIL (UNII: 69VJ1LPN1S)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
CENTELLA ASIATICA LEAF (UNII: 6810070TYD)	
CAMPHOR OIL (UNII: 75IZZ8Y727)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DIMETHYL ISOSORBIDE (UNII: SA6A6V432S)	
GLYCYRRHIZINATE DIPOTASSIUM (UNII: CA2Y0FE3FX)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
EMU OIL (UNII: 344821WD61)	
DIETHYLENE GLYCOL MONOETHYL ETHER (UNII: A1A1I8X02B)	
EUCALYPTUS OIL (UNII: 2R04ONI662)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCERYL MONOSTEARATE (UNII: 230OU9XXE4)	
SUNFLOWER OIL (UNII: 3W1JG795YI)	
ISOPROPYL MYRISTATE (UNII: ORE8K4LNJS)	
LAURETH-7 (UNII: Z95S6G8201)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
TEA TREE OIL (UNII: VIF565UC2G)	
PEG-100 STEARATE (UNII: YD01N1999R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
PYRIDOXINE HYDROCHLORIDE (UNII: 68Y4CF58BV)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63742-013-00	65 g in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	
2	NDC:63742-013-01	101 g in 1 BOTTLE; Type 0: Not a Combination Product	05/01/2019	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	05/01/2019	

Labeler - Clinical Resolution Laboratory, Inc. (825047942)

Establishment

Name	Address	ID/FEI	Business Operations
Clinical Resolution Laboratory, Inc.		825047942	manufacture(63742-013)

Revised: 3/2026

Clinical Resolution Laboratory, Inc.