

CALMARA- calmara lotion
Dynarex Corporation

1200 Calmara

Active ingredient

Camphor 0.5%

Purpose

External Analgesic

Uses

For temporary relief of pain and itching associated with

- minor burns
- minor cuts
- insect bites
- sunburn
- scrapes
- minor skin irritations

Warnings

For external use only

When using this product

- Avoid contact with the eyes

Stop use and ask a doctor 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 (1-800-222-1222) right away.

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: do not use, consult a doctor.

Other information

- Store at room temperature 15°-30°C (59°-86°F)
- Avoid excessive heat
- **Tamper evident. Do not use if seal is damaged.**

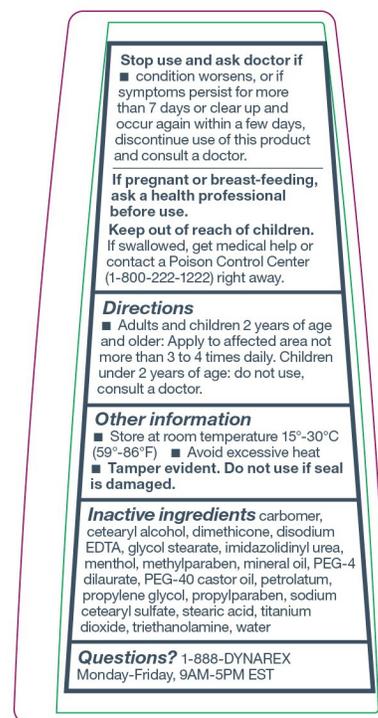
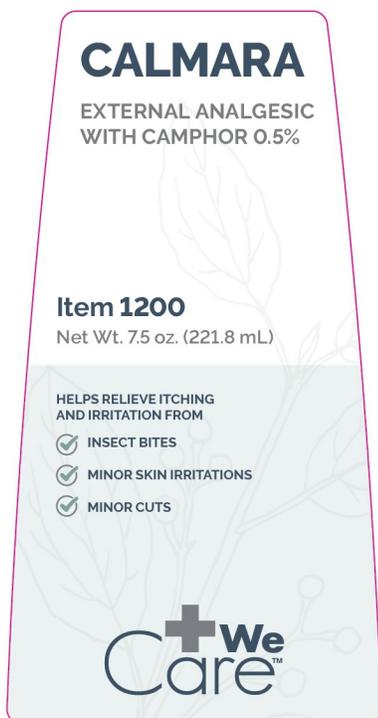
Inactive ingredients

carbomer, ceteryl alcohol, dimethicone, disodium EDTA, glycol stearate, imidazolidinyl urea, menthol, methylparaben, mineral oil, PEG-4 dilaurate, PEG-40 castor oil, petrolatum, propylene glycol, propylparaben, sodium ceteryl sulfate, stearic acid, titanium dioxide, triethanolamine, water

Questions?

1-888-DYNAREX Monday-Friday, 9AM-5PM EST

Label



CALMARA
calmara lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67777-017
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	0.5 mg in 100 mL

Inactive Ingredients

Ingredient Name	Strength
PETROLATUM (UNII: 4T6H12BN9U)	
MINERAL OIL (UNII: T5L8T28FGP)	
PEG-40 CASTOR OIL (UNII: 4ERD2076EF)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
MENTHOL (UNII: L7T10EIP3A)	
WATER (UNII: 059QF0KO0R)	
PEG-4 DILAURATE (UNII: KCR71CW036)	
METHYLPARABEN (UNII: A2I8C7HI9T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
TRIETHANOLAMINE (UNII: 9O3K93S3TK)	
IMIDAZOLIDINYL UREA (UNII: M629807ATL)	
CARBOMER 940 (UNII: 4Q93RCW27E)	
SODIUM CETEARYL SULFATE (UNII: 7ZBS06BH4B)	
CETEARYL ALCOHOL (UNII: 2DMT128M1S)	
DIMETHICONE (UNII: 92RU3N3Y1O)	

Product Characteristics

Color		Score	
Shape	FREEFORM	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:67777-017-02	24 in 1 CASE	01/21/2026	
1	NDC:67777-017-01	221.8 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M017	01/21/2026	

Labeler - Dynarex Corporation (008124539)

Revised: 1/2026

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