MENTHOLATUM ORIGINAL- camphor, menthol ointment The Mentholatum Company

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.

Drug Facts

Active ingredients

Camphor 9%

Menthol 1.3%

Purpose

Camphor - Topical Analgesic

Menthol - Topical Analgesic

Uses

temporarily relieves minor aches and pains of muscles and joints

Warnings

For external use only

When using this product

- do not get into eyes
- do not apply to wounds or to damaged skin
- do not bandage tightly
- do not heat, microwave, or add to hot water as this may cause splattering and result in burns

Stop use and ask a doctor if

- condition worsens
- symptoms persist for more than 7 days or clear up and occur again within a few days

Keep out of reach of children.

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

Directions

• adults and children 2 years and over: apply to affected area not more than 3 to 4 times daily

• children under 2 years: ask a doctor

Inactive ingredients

fragrance, petrolatum, titanium dioxide

Questions?

1-877-636-2677 MON-FRI 9AM to 5PM (EST)

Package/Label Principal Display Panel



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Prod	uct	ıntorr	mation

Due du et Ture	LILIMANI OTC DDLIC	Ham Cada (Causas)	NDC-10742 0002
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10742-0002

Route of Administration TOPICAL

Active Ingredient/Active Moiety				
Ingredient Name	Basis of Strength	Strength		
CAMPHOR (NATURAL) (UNII: N20HL7Q941) (CAMPHOR (NATURAL) - UNII:N20HL7Q941)	CAMPHOR (NATURAL)	90 mg in 1 g		
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL, UNSPECIFIED FORM - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	13 mg in 1 g		

Inactive Ingredients		
Ingredient Name	Strength	
PETROLATUM (UNII: 4T6H12BN9U)		
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742- 0002-1	1 in 1 CARTON	12/01/2014	
1		28 g in 1 JAR; Type 0: Not a Combination Product		
2	NDC:10742- 0002-2	1 in 1 CARTON	12/01/2014	
2		85 g in 1 JAR; Type 0: Not a Combination Product		
3	NDC:10742- 0002-3	1 in 1 CARTON	10/19/2020	
3		34 g in 1 JAR; Type 0: Not a Combination Product		
4	NDC:10742- 0002-4	1 in 1 CARTON	10/19/2020	
4		102.1 g in 1 JAR; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	12/01/2014	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment				
Name	Address	ID/FEI	Business Operations	
The Mentholatum Company		002105757	manufacture(10742-0002)	

Revised: 2/2023 The Mentholatum Company