MENTHOLATUM PAIN RELIEVING- menthol, methyl salicylate lotion
The Mentholatum Company

Drug Facts - Mentholatum Pain Relieving

Active ingredients

Menthol 6%

Methyl salicylate 20%

Purpose

Menthol - Topical analgesic

Methyl salicylate - Topical analgesic

Uses

temporarily relieves minor aches and pains of muscles and joints due to
- arthritis
- simple backache
- strains
- sprains

Warnings

For external use only

When using this product
- use only as directed
- do not get into eyes or on mucous membranes
- do not apply to wounds or to damaged skin
- do not bandage tightly
- do not use with a heating pad, other heat sources, or right after a shower/bath
- do not use in combination with other external analgesic products

Stop use and ask a doctor if
- condition worsens
- excessive irritation, burning or discomfort of the skin develops
- symptoms persist for more than 7 days or clear up and occur again within a few days

If pregnant or breast-feeding,

ask a healthcare professional before use.

Keep Out of Reach of Children.

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

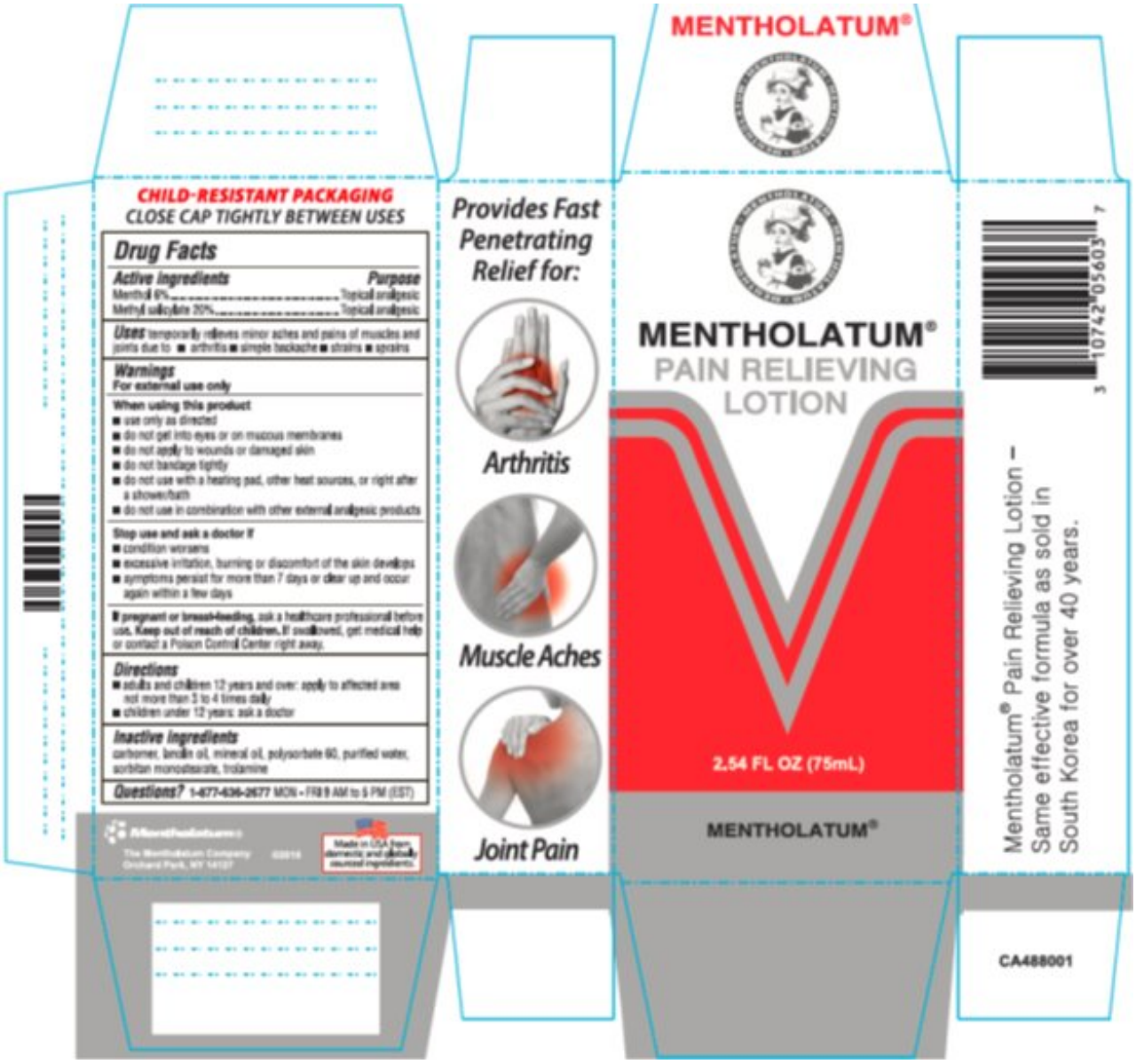
Directions

- adults and children 12 years and over: apply to affected area not more than 3 to 4 times daily
- children under 12 years: ask a doctor

Inactive ingredients

carbomer, lanolin oil, polysorbate 60, purified water, sorbitan monosterate, trolamine

Package/Label Principal Display Panel



MENTHOLATUM PAIN RELIEVING

menthol, methyl salicylate lotion

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:10742-8169
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL, UNSPECIFIED FORM (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL, UNSPECIFIED FORM	60 mg in 1 mL
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	200 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
LANOLIN OIL (UNII: OVV5IIJ58F)	
MINERAL OIL (UNII: T5L8T28FGP)	
POLYSORBATE 60 (UNII: CAL22UVI4M)	
WATER (UNII: 059QF0KO0R)	
SORBITAN MONOSTEARATE (UNII: NVZ4I0H58X)	
TROLAMINE (UNII: 9O3K93S3TK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:10742-8169-1	1 in 1 CARTON	11/01/2018	
1		100 mL in 1 BOTTLE; Type 0: Not a Combination Product		
2	NDC:10742-8169-2	1 in 1 CARTON	11/01/2018	
2		75 mL in 1 BOTTLE; 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	11/01/2018	

Labeler - The Mentholatum Company (002105757)

Registrant - The Mentholatum Company (002105757)

Establishment

Name	Address	ID/FEI	Business Operations
The Mentholatum Company		002105757	manufacture(10742-8169)

Revised: 12/2024 The Mentholatum Company