

FAST RELIEF REUMACETIN- menthol, methyl salicylate cream

Interfarma Corp

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.

FAST RELIEF REUMACETIN

DRUG FACTS

Active Ingredient:	Purpose:
Menthol 2.5%.....	External Analgesic
Methyl Salicylate 15%.....	External Analgesic

Uses

Temporarily relieves minor aches and pains of muscles and joints associated with simple backache, arthritis, strains, bruises, sprains.

Warnings:

- For external use only.
- Do not use this product if safety seal is broken. Store in room temperature.
- Do not use on wounds or damaged skin. Do not apply in large quantities, particularly on raw and irritated or blistered areas.
- When using this product, avoid contact with eyes or mucous membranes. Use only as directed. Do not bandage tightly.
- Stop and ask doctor if condition worsens, if symptoms persist within 7 days or clear up and occur again within a few days. Stop use if excessive skin irritation occurs.
- If pregnant or breast feeding, ask a health professional before use. If swallowed, get medical help or contact a Poison Control Center right away.

Keep out of reach of children to avoid accidental poisoning.

Caution: Discontinue use if excessive irritation of the skin develops. Avoid getting into eyes or mucous membranes. If condition worsens, or symptoms persist for more than 7 days or clear up and occur again within a few days, or redness is present, or in conditions affecting children under 12 years of age, discontinue use and consult a physician immediately.

Directions

Apply to the affected area. Gently massage into the skin for a few seconds. Adults and children over 12 years of age: apply to the affected area not more than 3 to 4 times daily. Children under 12 years of age need to consult a physician.

Inactive Ingredients:

Acrylamide, Amino Acids, Comfrey Leaf, Cetyl Alcohol, Diazolidinyl Urea, Dimethicone, Glucosamine, Methyl Paraben, MSM (Methylsulfonylmethane), Mineral Oil, Panthenol, Propylene Glycol, Propylparaben, Sodium Pyrrolidone Carboxylate, Stearic Acid, Stearyl Alcohol, Trolamine, Water.

Manufactured for:

Interfarma Corp.
Miami, FL 33186.

FAST RELIEF

REUMACETIN

For Temporarily Relieves minor aches and pains soreness, and stiffness associated with simple backaches strains, arthritis and rheumatism.

Net Wt 8 oz (226g)

Made in USA



FAST RELIEF REUMACETIN

menthol, methyl salicylate cream

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69706-0204
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.5 g in 100 g
METHYL SALICYLATE (UNII: LAV5U5022Y) (SALICYLIC ACID - UNII:O414PZ4LPZ)	METHYL SALICYLATE	15 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
ACRYLAMIDE (UNII: 20R035KLCI)	
AMINO ACIDS, WHEAT (UNII: 0370GZL32F)	
CETYL ALCOHOL (UNII: 936JST6JCN)	
COMFREY LEAF (UNII: DG4F8T839X)	
DIAZOLIDINYL UREA (UNII: H5RIZ3MPW4)	
DIMETHICONE (UNII: 92RU3N3Y1O)	
DIMETHYL SULFONE (UNII: 9H4PO4Z4FT)	
GLUCOSAMINE (UNII: N08U5BOQ1K)	
METHYL PARABEN (UNII: A2I8C7HI9T)	
MINERAL OIL (UNII: T5L8T28FGP)	
PANTHENOL (UNII: WV9CM0O67Z)	

PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
PROPYLPARABEN (UNII: Z8IX2SC1OH)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	
STEARIC ACID (UNII: 4ELV7Z65AP)	
STEARYL ALCOHOL (UNII: 2KR89I4HIY)	
TROLAMINE (UNII: 9O3K93S3TK)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69706-0204-9	226 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	11/11/2013	

Labeler - Interfarma Corp (884091703)

Registrant - Interfarma Corp (884091703)

Establishment

Name	Address	ID/FEI	Business Operations
Interfarma Corp		884091703	manufacture(69706-0204)

Revised: 10/2015

Interfarma Corp