

KWIK-MOVING- eucalyptus, menthol, methyl salicylate ointment

N. N. IMPEX

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.

Active ingredients

Eucalyptus Oil 2% , Menthol 5% ,Methyl salicylate 15%

Purpose

External Analgesic

Warning

For external use only.

Allergy alert:If prone to allergic reaction from aspirin or salicylates, consult a doctor before use.

Stop use and ask a doctor if condition worsens or if symptoms persist for more than 7 days or clear up and occurs again within a few days. Redness is present. Excessive irritation of the skin develops.

When using this product

Use only as directed. Avoid contact with the eyes or mucous membranes. Do not apply on wounds and damaged skin. Do not bandage tightly. Do not use with heating pad.

If pregnant or breast-feeding, ask a health professional before use.

Uses

For temporary relief of minor aches and pains of muscles and joints due to: simple backaches, bruises, sprains, arthritis

Direction

Adults and children 12 years and older: apply to the affected area and massage gently not more than 3 to 4 times daily.

Children under 12 years of age :consult a physician.

Keep out of reach of children

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

Inactive Ingredient

Light liquid paraffin, microcrystalline wax, paraffin wax, turpentine oil

Other information

Store at room temperature.

Product label



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73640-019
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
EUCALYPTUS OIL (UNII: 2R04ONI662) (EUCALYPTUS OIL - UNII:2R04ONI662)	EUCALYPTUS OIL	2 g in 100 g
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	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
LIGHT MINERAL OIL (UNII: N6K5787QVP)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
PARAFFIN (UNII: I9O0E3H2ZE)	
TURPENTINE OIL (UNII: C5H0QJ6V7F)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73640-019-01	50 g in 1 TUBE; Type 0: Not a Combination Product	02/05/2021	
2	NDC:73640-019-02	100 g in 1 TUBE; Type 0: Not a Combination Product	02/05/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	02/05/2021	

Labeler - N. N. IMPEX (915969592)

Establishment

Name	Address	ID/FEI	Business Operations
ECLAT PHARMA & AEROSOLS PVT LTD		877005000	manufacture(73640-019)

Revised: 2/2021

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