

ARTHRIMED-PLUS- menthol,eucalyptus oil,clove oil spray

Multi Service

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.

ArthriMed-Plus Spray

Active Ingredients

Menthol 2.5%

Encalyptus oil (Eucalyptus globutus) (leaf) 3%

Clove Oil (Eugenia caryophyllatus) (leaf) 0.2%

Purpose

Topical Analgesic

Uses

For the temporary relief of aches and pains of muscles and joints associated with backache, lumbago, strains, bruises, sprains and arthritic or rheumatic pain, pain of tendons and ligaments.

Warnings

For external use only. Do not apply to wounds or damaged skin. Do not inhale. Avoid contact with eyes and mucous membranes. Discontinue use if rash or irritation occurs. The application of external heat, such as an electric heating pad, may result in excessive skin irritation or skin burn. Do not bandage. Keep away from heat, sparks and open flames.

Keep out of reach of children.

Directions

For adults and children 2 years of age and older: Shake well before use. Apply to the affected area not more than 3-4 times daily. Allow absorption to occur naturally for maximum penetration. 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 health care practitioner.

Ingredient Name	Basis of Strength	Strength
MENTHOL (UNII: L7T10EIP3A) (MENTHOL - UNII:L7T10EIP3A)	MENTHOL	2.5 g in 100 g
EUCALYPTUS OIL (UNII: 2R04ONI662) (EUCALYPTUS OIL - UNII:2R04ONI662)	EUCALYPTUS OIL	3 g in 100 g
CLOVE OIL (UNII: 578389D6D0) (CLOVE OIL - UNII:578389D6D0)	CLOVE OIL	0.2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
CAMPHOR (SYNTHETIC) (UNII: 5TJD82A1ET)	
ISOPROPYL ALCOHOL (UNII: ND2M416302)	
PEPPERMINT OIL (UNII: AV092KU4JH)	
WATER (UNII: 059QF0KO0R)	
WEST INDIAN LEMONGRASS OIL (UNII: 5BIA40E9ED)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:82760-004-01	120 g in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	08/02/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part348	08/02/2022	

Labeler - Multi Service (208803005)

Establishment

Name	Address	ID/FEI	Business Operations
Multi Service		208803005	manufacture(82760-004)

Revised: 11/2022

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