

MYRISTICA ARGENTUM- myristica argentum liquid
Uriel Pharmacy Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Myristica Argentum

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Myristica sebifera (Juice from the bark of Myristica sebifera) 4X, Kalium bichromicum (Potassium dichromate) 6X, Tunica mucosa nasi (Bovine nasal mucosa) 7X, Argentum nitricum (Silver nitrate) 20X

Inactive Ingredients: Water, Salt

Use: Temporary relief of sinus pain.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use.

Questions? Call 866.642.2858

Uriel, East Troy, WI 53120

www.urielpharmacy.com

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Uriel, East Troy, WI 53120
shopuriel.com

Lot:

Uriel

**Myristica
Argentum**

Homeopathic Ampules
net vol. 0.3 fl. oz (10 x 1 ml)

Myristica Argentum

MYRISTICA ARGENTUM

myristica argentum liquid

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:48951-7062

Route of Administration	ORAL
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Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BOS TAURUS NASAL MUCOSA (UNII: 343455G79K) (BOS TAURUS NASAL MUCOSA - UNII:343455G79K)	BOS TAURUS NASAL MUCOSA	7 [hp_X] in 1 mL
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	20 [hp_X] in 1 mL
VIOLA SEBIFERA RESIN (UNII: GHJ5XX5SGS) (VIOLA SEBIFERA RESIN - UNII:GHJ5XX5SGS)	VIOLA SEBIFERA RESIN	4 [hp_X] in 1 mL
DICHROMATE ION (UNII: 9LKY4BFN2V) (DICHROMATE ION - UNII:9LKY4BFN2V)	DICHROMATE ION	6 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-7062-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		09/01/2009	

Labeler - Uriel Pharmacy Inc. (043471163)**Establishment**

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-7062)

Revised: 3/2024

Uriel Pharmacy Inc.