

VISCUM ABIETIS SERIES 1 7ML- viscum abietis series 1 7ml 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.

Viscum Abietis Series 1 7ml

Use: Temporary relief of headache.

FOR ORAL USE ONLY

Directions: Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow. Follow inside instructions to open ampule.

Warnings:

NOT FOR INJECTION

Do not use if allergic to any ingredient. Consult a physician if symptoms worsen. If pregnant or nursing, consult a physician before use.

Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated.

KEEP OUT OF REACH OF CHILDREN.

Active Ingredient: Viscum Abietis (Fir tree mistletoe)

3 amps 1mg = 3X

3 amps 5mg = 1 gm contains: 2X, 500mg

1 amp 10mg = 2X

Inactive Ingredients: Water, Salt, Sodium Hydroxide

Prepared using rhythmical processes.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 shopuriel.com Lot:

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(Fir tree mistletoe)

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Lot:

3 amps 5mg = 1 gm contains 2X, 500mg

Lot:

1 amp 10mg = 2X

Lot:

Inactive Ingredients: Water, Salt, Sodium hydroxide

Prepared using rhythmical processes.

Lot:


Viscum Abietis
Series 1
Homeopathic Ampules
FOR ORAL USE ONLY
net vol. 0.23 fl. oz (7 x 1ml)

Questions? Call 866.642.2858 Uriel, East Troy,
WI 53120 shopuriel.com

Viscum Abietis Series 1

VISCUM ABIETIS SERIES 1 7ML

viscum abietis series 1 7ml liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-9242
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
VISCUM ALBUM FRUITING TOP (UNII: BK9092J5MP) (VISCUM ALBUM FRUITING TOP - UNII:BK9092J5MP)	VISCUM ALBUM FRUITING TOP	2 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-9242-1	7 in 1 BOX	08/01/2014	
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(48951-9242)

Revised: 2/2026

Uriel Pharmacy Inc.