

VISCUM PINI SERIES 2 7ML- viscum pini series 2 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 Pini Series 2 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 Pini (Pine tree mistletoe)

2 amps 10mg = 2X

2 amps 20mg = 1 gm contains: 1X, 200mg

3 amps 30mg = 1 gm contains: 1X, 300mg

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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Active Ingredient: Viscum album (Pini) (Pine tree mistletoe)

2 amps 10mg = 2X

2 amps 20mg = 1gm contains: 1X, 200mg

3 amps 30mg = 1gm contains: 1X, 300mg

Inactive Ingredients: Water, Salt, Sodium hydroxide

Prepared using rhythmical processes.

Lot:



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Viscum Pini Series 2

VISCUM PINI SERIES 2 7ML

viscum pini series 2 7ml liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-9257
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	1 [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:48951-9257-1	7 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(48951-9257)

Revised: 2/2026

Uriel Pharmacy Inc.