

ARGENTUM 8- argentum 8 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.

Argentum 8

Directions: FOR ORAL USE.

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

Active Ingredient: Argentum met. (Silver) 8X

Inactive Ingredients: Water, Salt

"prepared using rhythmical processes"

Use: Temporary relief of headache.

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

shopuriel.com Lot:

Directions: FOR ORAL USE.
Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredient: Argentum met. (Silver) 8X

Inactive Ingredients: Water, Salt, Lactose

"prepared using rhythmical processes"

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN

Warning: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Contains traces of lactose. 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
shopuriel.com

Lot:

Argentum 8X

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

ARGENTUM 8

argentum 8 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1087	
Route of Administration	ORAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)		SILVER	8 [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:48951-1087-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 0: Not a Combination Product		
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(48951-1087)

Revised: 1/2024

Uriel Pharmacy Inc.