

## ARGENTUM QUARTZ- argentum quartz 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.*

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### Argentum Quartz

Directions: FOR ORAL USE.

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

Active Ingredients: Argentum met. (Silver) 20X, Quartz (Rock crystal) 30X

Inactive Ingredients: Water, Salt

Use: Temporary relief of ear irritation.

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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www.urielpharmacy.com Lot:



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## ARGENTUM QUARTZ

argentum quartz liquid

### Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-1098
Route of Administration	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SILVER (UNII: 3M4G523W1G) (SILVER - UNII:3M4G523W1G)	SILVER	20 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	30 [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:48951-1098-1	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(48951-1098)