

**AURUM 21X- aurum 21x 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.*

-----  
**Aurum 21X**

Directions: FOR ORAL USE.

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

Active Ingredient: Aurum metallicum (Metallic gold) 21X

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

Made by 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: Aurum metallicum (Metallic gold) 21X

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  
Made by Uriel, East Troy, WI 53120  
shopuriel.com

Lot:



Aurum 21X

**AURUM 21X**

aurum 21x liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-1318
<b>Route of Administration</b>	ORAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>GOLD</b> (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	21 [hp_X] in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)	
<b>WATER</b> (UNII: 059QF0KO0R)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1318-1	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(48951-1318)

Revised: 1/2025

Uriel Pharmacy Inc.