

AURUM HYPERICUM STIBIUM- aurum hypericum stibium 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 Hypericum Stibium

Directions: FOR ORAL USE

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

Active Ingredients: Aurum met. (Metallic gold) 10X, Hypericum (St. Johns wort) 10X, Stibium met. (Antimony) 10X

Inactive Ingredients: Water, Salt

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:

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Lot:



Aurum Hypericum
Stibium

AURUM HYPERICUM STIBIUM

aurum hypericum stibium liquid

Product Information

| | | | |
|--------------------------------|----------------|---------------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:48951-1340 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|---|-------------------|-------------------|
| ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3) | ANTIMONY | 10 [hp_X] in 1 mL |
| GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO) | GOLD | 10 [hp_X] in 1 mL |
| ST. JOHN'S WORT (UNII: UFH8805FKA) (ST. JOHN'S WORT - UNII:UFH8805FKA) | ST. JOHN'S WORT | 10 [hp_X] in 1 mL |

Inactive Ingredients

| Ingredient Name | Strength |
|---|----------|
| SODIUM CHLORIDE (UNII: 451W47IQ8X) | |
| WATER (UNII: 059QF0KO0R) | |

Packaging

| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
|---|------------------|---|----------------------|--------------------|
| 1 | NDC:48951-1340-1 | 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(48951-1340) |

Revised: 5/2022

Uriel Pharmacy Inc.