AURUM STIBIUM HYOSCYAMUS- aurum stibium hyoscyamus 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 Stibium Hyoscyamus

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredients: Onopordon (Cotton thistle) 3X, Primula (Cowslip) 3X, Hyoscyamus (Henbane) 4X, Stibium metallicum (Antimony) 8X, Aurum metallicum (Metallic gold) 10X

Inactive Ingredients: Distilled water, Organic cane alcohol

"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. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858
Made with care by Uriel, East Troy, WI 53120
shopuriel.com

AURUM STIBIUM HYOSCYAMUS

aurum stibium hyoscyamus liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1206

Route of Administration	ORAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ONOPORDUM ACANTHIUM FLOWER (UNII: AP97AUF88E) (ONOPORDUM ACANTHIUM FLOWER - UNII:AP97AUF88E)	ONOPORDUM ACANTHIUM FLOWER	3 [hp_X]
PRIMULA VERIS FLOWER (UNII: W5BET37294) (PRIMULA VERIS FLOWER - UNII:W5BET37294)	PRIMULA VERIS FLOWER	3 [hp_X]
HYOSCYAMUS NIGER LEAF (UNII: 32IT7G8BAW) (HYOSCYAMUS NIGER LEAF - UNII:32IT7G8BAW)	HYOSCYAMUS NIGER LEAF	4 [hp_X]
ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3)	ANTIMONY	8 [hp_X]
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	10 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1206-3	60 in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	09/01/2009	

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(48951-1206)