

STROPHANTHUS AURUM- strophanthus aurum 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.

Strophanthus Aurum

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, Strophanthus (Strophantus) 5X, Stibium met. (Antimony) 8X, Aurum met. (Metallic gold) 10X

Inactive Ingredients: Distilled water, 20% Organic cane alcohol

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
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STROPHANTHUS AURUM

strophanthus aurum liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:48951-8403
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] in 1 mL
PRIMULA VERIS FLOWER (UNII: W5BET37294) (PRIMULA VERIS FLOWER - UNII:W5BET37294)	PRIMULA VERIS FLOWER	3 [hp_X] in 1 mL
HYOSCYAMUS NIGER LEAF (UNII: 32IT7G8BAW) (HYOSCYAMUS NIGER LEAF - UNII:32IT7G8BAW)	HYOSCYAMUS NIGER LEAF	4 [hp_X] in 1 mL
STROPHANTHUS KOMBE WHOLE (UNII: Q1O0N901S4) (STROPHANTHUS KOMBE WHOLE - UNII:Q1O0N901S4)	STROPHANTHUS KOMBE WHOLE	5 [hp_X] in 1 mL
ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3)	ANTIMONY	8 [hp_X] in 1 mL
GOLD (UNII: 79Y1949PYO) (GOLD - UNII:79Y1949PYO)	GOLD	10 [hp_X] in 1 mL

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8403-3	60 mL 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(48951-8403)

Revised: 11/2023

Uriel Pharmacy Inc.