CUPRUM STANNUM- cuprum stannum 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.

Cuprum Stannum

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: 100gm contains: 94gm Acidum nitricum (Nitric acid) 2X, 40gm Acidum sulfuricum (Sulfuric acid) 2X, 20gm Stannum met. (Metallic tin) 2X, 2gm Cuprum met. (Copper) 3X; Alumen (Potassium aluminum sulfate) 3X

Inactive Ingredients: Distilled water, Silica

Use: Temporary relief of digestive upset.

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

CUPRUM STANNUM

cuprum stannum liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-3145
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
NITRIC ACID (UNII: 411VRN1TV4) (NITRIC ACID - UNII:411VRN1TV4)	NITRIC ACID	2 [hp_X] in 1 mL
SULFURIC ACID (UNII: O40UQP6WCF) (SULFURIC ACID - UNII:O40UQP6WCF)	SULFURIC ACID	2 [hp_X] in 1 mL
TIN (UNII: 387GMG9FH5) (TIN - UNII:387GMG9FH5)	TIN	2 [hp_X] in 1 mL
COPPER (UNII: 789U1901C5) (COPPER - UNII:789U1901C5)	COPPER	3 [hp_X] in 1 mL
POTASSIUM ALUM (UNII: 1L24V9R23S) (ALUMINUM HYDROXIDE - UNII:5QB0T2IUN0)	POTASSIUM ALUM	3 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SILICON DIOXIDE (UNII: ETJ7Z6XBU4)	

Product Characteristics

Color	white	**Score**	
Shape		**Size**	
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-3145-3	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(48951-3145)