FORMICA STIBIUM COMP. SPECIAL ORDER- formica stibium comp. special order 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.

Formica Stibium comp. Special Order

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 Ingredient: Pulmo (Bovine lung) 6X, Stibium met. (Antimony) 8X, Nontronite (Nat. Iron silicate) 12X, Formica (Red wood ant) 15X

Inactive Ingredient: Distilled water

Use: Temporary relief of cough.

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. Contains traces of lactose. 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.

REFRIGERATE AFTER OPENING.
BEST WHEN USED WITHIN 90 DAYS OF OPENING.

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

FORMICA STIBIUM COMP. SPECIAL ORDER

formica stibium comp. special order liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-4102
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**

SUS SCROFA LUNG (UNII: 7GL3G1COB3) (SUS SCROFA LUNG - UNII:7GL3G1COB3)	SUS SCROFA LUNG	6 [hp_X] in 1 mL
ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3)	ANTIMONY	8 [hp_X] in 1 mL
SILICON DIOXIDE (UNII: ETJ7Z6XBU4) (SILICON DIOXIDE - UNII:ETJ7Z6XBU4)	SILICON DIOXIDE	12 [hp_X] in 1 mL
FORMICA RUFA (UNII: 55H0W83JO5) (FORMICA RUFA - UNII:55H0W83JO5)	FORMICA RUFA	15 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-4102-3	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(48951-4102)

Revised: 4/2018 Uriel Pharmacy Inc.