VITIS STIBIUM- vitis stibium tablet
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.

Vitis Stibium

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 1-2 tablets. Ages 2-11: 1 tablet. Under age 2: Consult a doctor.

Active Ingredients: Fragaria (Wild strawberry) 1X double strength, Vitis (Grape vine) 1X double strength, Calcarea Formicica 3X, Stibium (Antimony) 6X

Inactive Ingredients: Glucose monohydrate, Lactose, Magnesium stearate

"prepared using rhythmical processes"

Use: Supports normal liver function.

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.

Questions? Call 866.642.2858
Made by Uriel, East Troy, WI 53120
shopuriel.com Lot:

VITIS STIBIUM
vitis stibium tablet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-9377
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
FRAGARIA VESCA WHOLE (UNII: 5J340B7H82) (FRAGARIA VESCA WHOLE - UNII:5J340B7H82)	FRAGARIA VESCA WHOLE	1 [hp_X]
VITIS VINIFERA LEAF (UNII: R1H893D80E) (VITIS VINIFERA LEAF - UNII:R1H893D80E)	VITIS VINIFERA LEAF	1 [hp_X]
CALCIUM FORMATE (UNII: NP3JD65NPY) (CALCIUM CATION - UNII:2M83C4R6ZB)	CALCIUM CATION	3 [hp_X]
ANTIMONY (UNII: 9IT35J3UV3) (ANTIMONY - UNII:9IT35J3UV3)	ANTIMONY	6 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
DEXTRATES (UNII: G263MI44RU)	
MAGNESIUM STEARATE (UNII: 70097M6I30)	

Product Characteristics

Color	brown	**Score**	no score
Shape	ROUND	**Size**	10mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-9377-4	50 in 1 BOTTLE, GLASS; 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(48951-9377)

Revised: 1/2024 Uriel Pharmacy, Inc.