ANTIMONITE BELLADONNA- antimonite belladonna powder
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.

Antimonite Belladonna

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 1/8 teaspoon. Ages 2-11: 1/16 teaspoon. Under age 2: Consult a doctor.

Active Ingredients: Antimonite (Nat. antimony trisulfide) 3X, Atropa belladonna (Nightshade) 4X, Chamomilla (Chamomile) 4X, Bismutum subnitricum (Bismuth) 6X

Inactive Ingredient: Lactose

"prepared using rhythmical processes"

Use: Temporary relief of upset stomach.

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 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 Uriel, East Troy, WI 53120 shopuriel.com

ANTIMONITE BELLADONNA
antimonite belladonna powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-1037
Route of Administration	ORAL		

Active Ingredient/Active Moiety	

Ingredient Name	Basis of Strength	Strength
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	3 [hp_X] in 1 g
ATROPA BELLADONNA (UNII: WQZ3G9PF0H) (ATROPA BELLADONNA - UNII:WQZ3G9PF0H)	ATROPA BELLADONNA	4 [hp_X] in 1 g
MATRICARIA RECUTITA (UNII: G0R4UBI2ZZ) (MATRICARIA RECUTITA - UNII:G0R4UBI2ZZ)	MATRICARIA RECUTITA	4 [hp_X] in 1 g
BISMUTH SUBNITRATE (UNII: H19J064BA5) (BISMUTH CATION - UNII:ZS9CD1I8YE)	BISMUTH SUBNITRATE	6 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-1037-4	50 g 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(48951-1037)

Revised: 1/2024 Uriel Pharmacy Inc.