BISMUTH MAGNESITE COMP.- bismuth magnesite comp. 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.

Bismuth Magnesite comp. Special Order

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: Bismuth 3X, Magnesite (Magnesium carbonate) 3X, Antimonite (Antimony trisulfide) 6X, Corallium rubrum (Red coral) 6X,

Crocus sativa (Saffron) 6X, Kalium aceticum (Potassium acetate)6X

Inactive Ingredient: Lactose

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. 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
www.urielpharmacy.com

BISMUTH MAGNESITE COMP.

bismuth magnesite comp. powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-8307
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BISMUTH (UNII: U015TT5I8H) (BISMUTH - UNII:U015TT5I8H)	BISMUTH	3 [hp_X]

BISMUTH (UNII: U015T1514I) (BISMUTH - UNII:U015T1514I)	BISMUTH	in 1 g
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	6 [hp_X] in 1 g
SAFFRON (UNII: E849G4X5YJ) (SAFFRON - UNII:E849G4X5YJ)	SAFFRON	6 [hp_X] in 1 g
CORALLIUM RUBRUM EXOSKELETON (UNII: 2CA71K0DLE) (CORALLIUM RUBRUM EXOSKELETON - UNII:2CA71K0DLE)	CORALLIUM RUBRUM EXOSKELETON	6 [hp_X] in 1 g
MAGNESITE (UNII: 0IHC698356) (MAGNESITE - UNII:0IHC698356)	MAGNESITE	3 [hp_X] in 1 g
POTASSIUM ACETATE (UNII: M911911U02) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM ACETATE	6 [hp_X] in 1 g

Inactive Ingredients

Ingredient Name	Strength
LACTOSE (UNII: J2B2A4N98G)	1 [hp_X] in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-8307-4	50 g in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	01/09/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/09/2009	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-8307)

Revised: 4/2018 Uriel Pharmacy Inc.