KALIUM ACETICUM COMP. 6- kalium aceticum comp. 6 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.

Kalium aceticum comp. 6

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) 6X, Corallium rubrum (Red coral) 6X, Crocus sativa (Saffron) 6X, Kalium aceticum (Potassium acetate) 6X

Inactive Ingredient: Lactose

"prepared using rhythmical processes"

Use: Temporary relief of skin rash.

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

KALIUM ACETICUM COMP. 6

kalium aceticum comp. 6 powder

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-6004
Route of Administration	ORAL		

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

CORALLIUM RUBRUM EXOSKELETON (UNII: 2CA71K0DLE) (CORALLIUM RUBRUM EXOSKELETON - UNII:2CA71K0DLE)	CORALLIUM RUBRUM EXOSKELETON	6 [hp_X] in 1 g
SAFFRON (UNII: E849G4X5YJ) (SAFFRON - UNII:E849G4X5YJ)	SAFFRON	6 [hp_X] in 1 g
POTASSIUM ACETATE (UNII: M911911U02) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM ACETATE	6 [hp_X] in 1 g
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	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:48951-6004-4	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(48951-6004)

Revised: 1/2024 Uriel Pharmacy Inc.