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

Kalium aceticum comp. 6

Directions: FOR ORAL USE.

Take the contents of one ampule under the tongue and hold for 30 seconds, then swallow.

Active Ingredients: Antimonite (Nat. antimony trisulfide) 6X, Corallium rubrum (Red coral) 6X, Crocus sativa (Saffron) 6X, Kalium aceticum (Potassium acetate) 6X

Inactive Ingredients: Water, Salt, Lactose

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 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.

Questions? Call 866.642.2858 Uriel, East Troy, WI 53120 www.urielpharmacy.com

KALIUM ACETICUM COMP. 6

kalium aceticum comp. 6 liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-6006
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	6 [hp_X] in 1 mL

CORALLIUM RUBRUM EXOSKELETON (UNII: 2CA71K0DLE) (CORALLIUM RUBRUM EXOSKELETON - UNII:2CA71K0DLE)	CORALLIUM RUBRUM EXOSKELETON	6 [hp_X] in 1 mL
SAFFRON (UNII: E849G4X5YJ) (SAFFRON - UNII:E849G4X5YJ)	SAFFRON	6 [hp_X] in 1 mL
POTASSIUM ACETATE (UNII: M911911U02) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM ACETATE	6 [hp_X] in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LACTOSE (UNII: J2B2A4N98G)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-6006-1	10 in 1 BOX	09/01/2009	
1		1 mL in 1 AMPULE; Type 1: Convenience Kit of Co-Package		

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(48951-6006)

Revised: 4/2018 Uriel Pharmacy Inc.