KALIUM ACETICUM COMP 6- kalium aceticum comp 6 pellet
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 6X

Directions: FOR ORAL USE ONLY.

Dissolve pellets under the tongue 3-4 times daily. Ages 12 and older: 10 pellets. Ages 2-11: 5 pellets. Under age 2: Consult
a doctor.

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

Inactive Ingredient: Organic sucrose

"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. Contains sugar. Diabetics and persons intolerant of sucrose (sugar): Consult a doctor before use. Contains traces of lactose. Do not use if allergic to any ingredient. 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 Lot:

KALIUM ACETICUM COMP 6

kalium aceticum comp 6 pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:48951-6079
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SAFFRON (UNII: E849G4X5YJ) (SAFFRON - UNII:E849G4X5YJ)	SAFFRON	6 [hp_X]
POTASSIUM ACETATE (UNII: M911911U02) (POTASSIUM CATION - UNII:295O53K152)	POTASSIUM ACETATE	6 [hp_X]
ANTIMONY TRISULFIDE (UNII: F79059A38U) (ANTIMONY TRISULFIDE - UNII:F79059A38U)	ANTIMONY TRISULFIDE	6 [hp_X]
CORALLIUM RUBRUM EXOSKELETON (UNII: 2CA71K0DLE) (CORALLIUM RUBRUM EXOSKELETON - UNII:2CA71K0DLE)	CORALLIUM RUBRUM EXOSKELETON	6 [hp_X]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:48951-6079-2	1350 in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	08/19/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		08/19/2021	

Labeler - Uriel Pharmacy Inc. (043471163)

Establishment

Name	Address	ID/FEI	Business Operations
Uriel Pharmacy Inc.		043471163	manufacture(48951-6079)

Revised: 1/2024 Uriel Pharmacy Inc.