KALI ARSENICOSUM- potassium arsenite anhydrous pellet
Boiron

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.

Kali arsenicosum 200CK

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(**contains 0.443 mg of the active ingredient per pellet)

Dry skin rash with itching worsened by heat*

Stop use and ask a doctor if symptoms persist for more than 3 days or worsen

If pregnant or breast-feeding ask a health professional before use

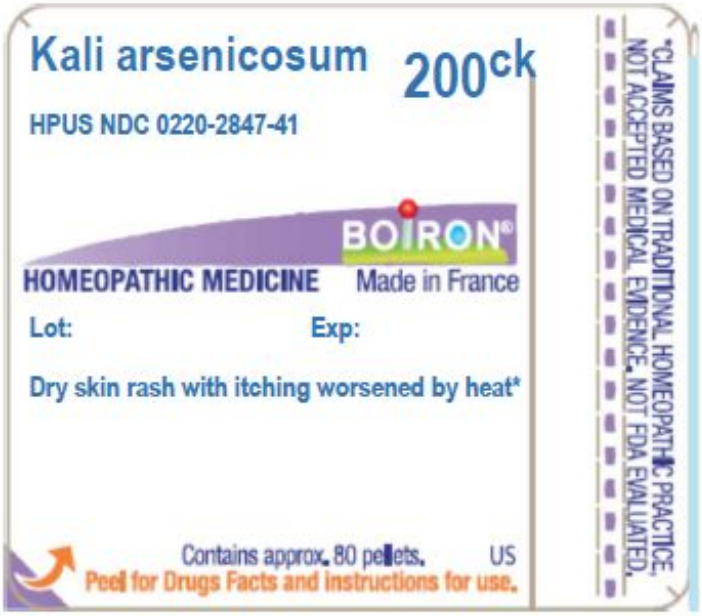
Keep out of reach of children

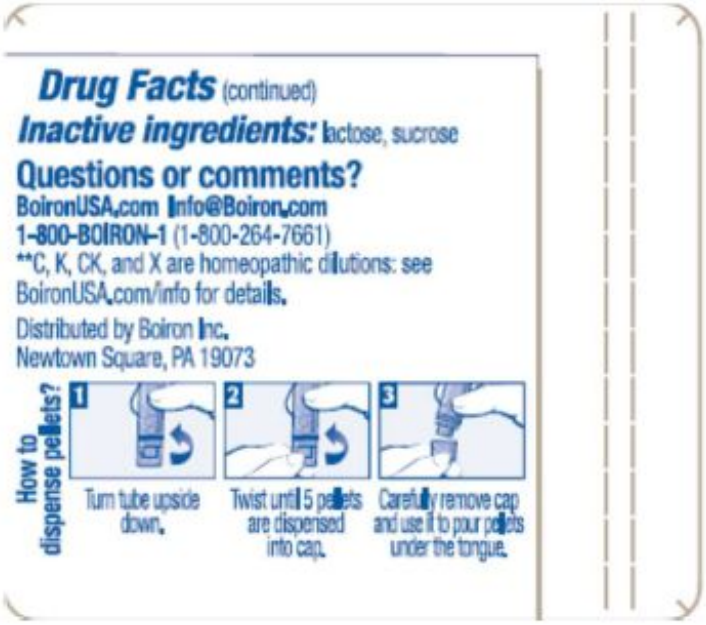
Do not use if pellet dispenser seal is broken.
Contains approx 80 pellets.
How to dispense pellets? Turn tube upside down. Twist until 5 pellets are dispensed into cap. Carefully remove the cap and use it to pour pellets under the tongue.
*CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.
*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

lactose, sucrose

Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

1-800-BOIRON-1 (1-800-264-7661),
BoironUSA.com Info@boiron.com
Distributed by Boiron, Inc. Newtown Square, PA 19073





KALI ARSENICOSUM

potassium arsenite anhydrous pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0220-2847
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
POTASSIUM ARSENITE ANHYDROUS (UNII: BM2U42PAKI) (ARSENITE ION - UNII:N5509X556J)	POTASSIUM ARSENITE ANHYDROUS	200 [kp_C] in 200 [kp_C]

Inactive Ingredients	
Ingredient Name	**Strength**
SUCROSE (UNII: C151H8M554)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	

Product Characteristics			
Color	white	**Score**	
Shape	ROUND	**Size**	4mm
Flavor		**Imprint Code**	
Contains			

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**

1	NDC:0220-2847-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		03/03/1983	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment

Name	Address	ID/FEI	Business Operations
Boiron		282560473	manufacture(0220-2847)

Revised: 10/2023 Boiron