## KALI BROMATUM- potassium bromide 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.

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## Kali bromatum 15C

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

Sleeplessness with agitation\*

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





## **Drug Facts**

**Active ingredient**<sup>\*\*</sup>: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

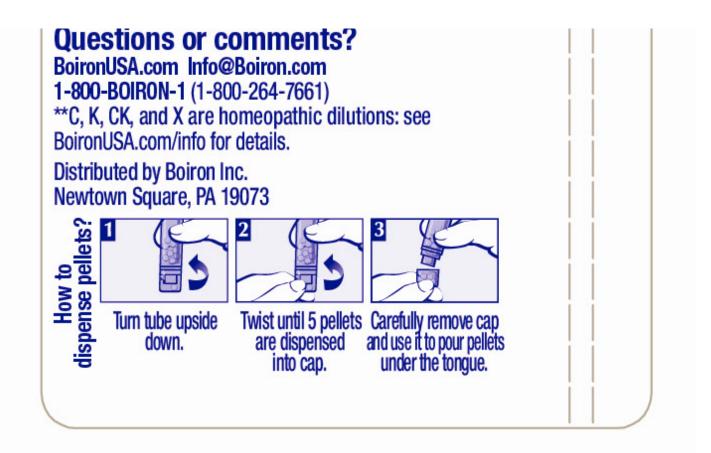
**Uses:** See symptoms on front panel.

**Warnings:** Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children.

**Directions:** 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.

**Other information:** Do not use if pellet dispenser seal is broken.

Drug Facts (continued) Inactive ingredients: lactose, sucrose



KALI BROMATUM				
potassium bromide pellet				
Product Information				
Product Type	HUMAN OTC DRUG	Item Cod	e (Source)	NDC:0220-2822
Route of Administration	ORAL			
Active Ingredient/Active	e Molety		Basis of	
Ingree	Strength			
POTASSIUM BROMIDE (UNII: OSD78555ZM) (BROMIDE ION - UNII:952902IX06)			POTASSIUM BROMIDE	15 [hp_C] in 15 [hp_C]
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)			
SUCROSE (UNII: C151H8M554)				
<b>Product Characteristics</b>	;			
Color	white	Score		
Shape	ROUND	Size		4mm

-	ontains Ackaging		on Mark	keting Start	Marketing End	
-	ackaging		on Mark	keting Start	Marketing End	
-	ackaging		on Mark	keting Start	Marketing End	
-	ackaging		on Marl	keting Start	Marketing End	
Code			on Mark	keting Start	Marketing End	
	ltem Code	Package Descripti		Date	Date	
	NDC:0220-2822 41	15 [hp_C] in 1 TUBE; Type 0: Not a Product	Combination 03/03/1	.983		
tina lı	arkating	Information				
Marketing Information						
etina	Marketing Category	Application Number or I Citation	Monograph Mai	rketing Start Date	Marketing End Date	
	approved		03/03	/1983		
	Catego			ory Citation		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment					
Name	Address	ID/FEI	Business Operations		
Boiron		282560473	manufacture(0220-2822)		

Revised: 10/2023

Boiron