## 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 5C

Kali bromatum 5C

(\*\*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\*\*: See product name on front

panel (contains 0.443 mg of the active ingredient per pellet).

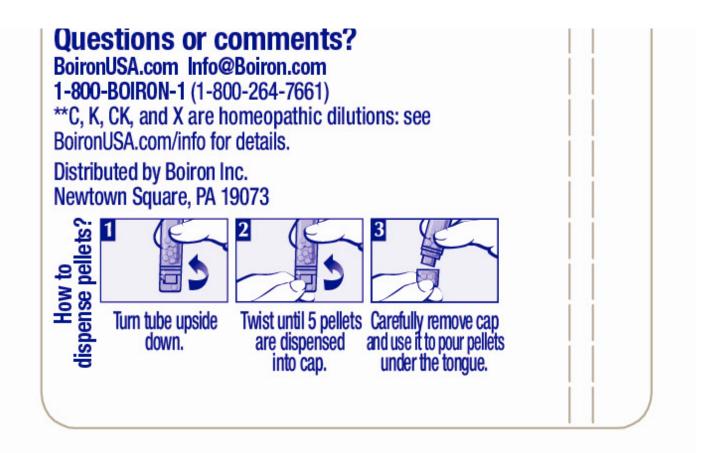
**Uses:** See symptoms on front panel.

**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 Code	(Source)	NDC:0220-2859
Route of Administration	ORAL			
Active Ingredient/Active	мојету			
Ingree	Strength			
POTASSIUM BROMIDE (UNII: OS UNII:952902IX06)	E 5 [hp_C] in 5 [hp_C]			
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
<b>Product Characteristics</b>				
Color	white	Score		
Shape	ROUND	Size		4mm

Flavor		Imprint C	Imprint Code					
С	ontains							
Packaging								
#	ltem Code	Р	ackage Description	Marketing Start Date	Marketing End Date			
1	NDC:0220-2859- 41	5 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983				
Marketing Information								
	Marketing Category		ation Number or Monograp Citation	h Marketing Start Date	Marketing End Date			

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-2859)				

Revised: 11/2023

Boiron