## KALI CHLORICUM- potassium chlorate 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 chloricum 30C

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

Pain from canker sores\*

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).

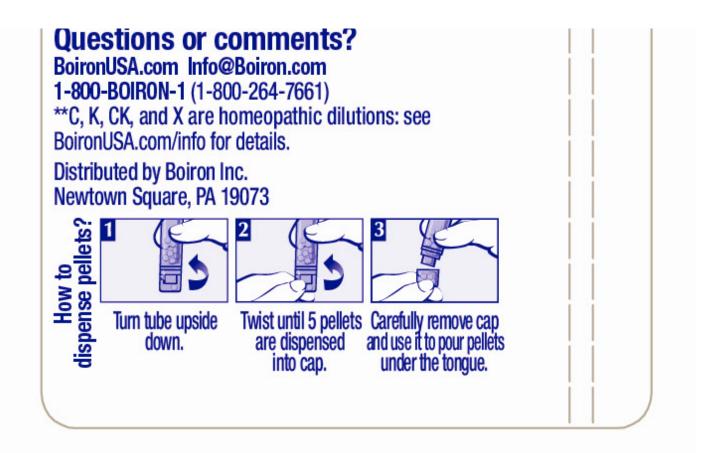
**Uses:** See symptoms on front panel.

**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 CHLORICUM						
potassium chlorate pellet						
Product Information						
Product Type	HUMAN OTC DRUG	HUMAN OTC DRUG Item Code (Source)			NDC:0220-2871	
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingre	Strength					
POTASSIUM CHLORATE (UNII: H35KS68EE7) (POTASSIUM CATION - POTASSIUM CHLORATE UNII:295053K152)					30 [hp_C] in 30 [hp_C]	
Inactive Ingredients						
	Strength					
SUCROSE (UNII: C151H8M554)						
LACTOSE, UNSPECIFIED FORM						
Product Characteristics						
Color	white	Score				
Shape	ROUND	Size			4mm	

Flavor		Imprint Code					
Contains							
Packaging							
#	ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
1	NDC:0220-2871- 41	30 [hp_C] in Product	_C] in 1 TUBE; Type 0: Not a Combination t		03/03/1983		
м	larkoting	Informa	tion				
Marketing Information							
	Marketing Category	Applic	ication Number or Monograph Citation		Marketing Start Date	Marketing End Date	
	approved meopathic				03/03/1983		
					03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Boiron		282560473	manufacture(0220-2871)				

Revised: 11/2023

Boiron