## KALI CARBONICUM- potassium carbonate 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 carbonicum 10M

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

Pain and feeling of weakness in the lower back\*

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



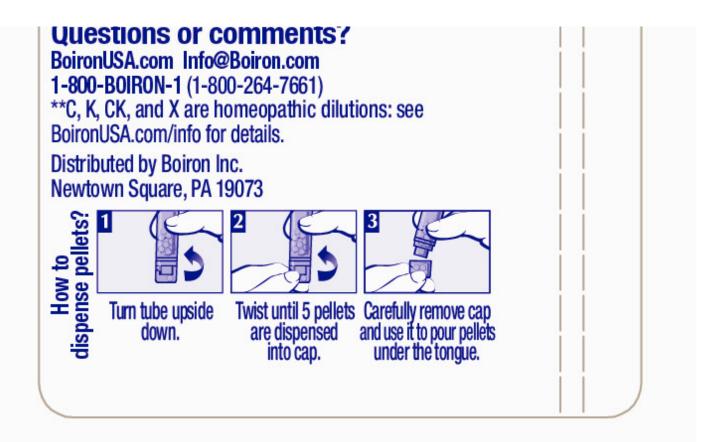


## Drug Facts Active ingredient\*\*: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before use. Keep out of reach of children. Directions: Adults and children: At the onset of cumptoms of persist for persist for persist for persist of children.

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 CARBONICUM				
ootassium carbonate pellet				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:0220-2816
Route of Administration	ORAL			
Active Ingredient/Active	Molety		Basis of	
Ingree	Strength			
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - UNII:7UJQ50PE7D)			POTASSIUM CARBONATE	10 [hp_M] in 10 [hp_M]
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
<b>Product Characteristics</b>				
Color	white	Score		
Shape	ROUND	Size		4mm

Flavor		Imprint Code					
ntains							
- <b>I</b>							
Packaging							
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date		
	10 [hp_M] in Product			03/03/1983			
o rikotin a	Informa	tion					
Marketing Information							
Marketing Category	Applic	ation Number or I Citation	Monograph	Marketing Start Date	Marketing End Date		
approved meopathic				03/03/1983			
	Item Code NDC:0220-2816- 41  arketing Marketing Category approved	ackaging         Item Code       P         NDC:0220-2816-       10 [hp_M] in 2         41       Product         arketing       Information         Marketing       Applic         Category       Applic         approved       Applic	Ackaging       Package Descripti         Item Code       Package Descripti         NDC:0220-2816-       10 [hp_M] in 1 TUBE; Type 0: Not a         41       Product         Image: State of the state	Ackaging       Package Description         Item Code       Package Description         NDC:0220-2816- 41       10 [hp_M] in 1 TUBE; Type 0: Not a Combination Product         Image: Comparison of the second sec	Ackaging       Marketing Start Date         Item Code       Package Description       Marketing Start Date         NDC:0220-2816- 41       10 [hp_M] in 1 TUBE; Type 0: Not a Combination Product       03/03/1983         Intersecting Information       Marketing Start Date       03/03/1983         Marketing Category       Application Number or Monograph Citation       Marketing Start Date         approved       03/03/1983       03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-2816)			

Revised: 11/2023

Boiron