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.

Kali carbonicum 6C

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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 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						
potassium carbonate pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code (Source)		ND	NDC:0220-2809	
Route of Administration	ORAL					
A stine I nave die ut/A stine	Maiatu					
Active Ingredient/Active	моюту			Basis of		
Ingre		Strength				
POTASSIUM CARBONATE (UNII: BQN1B9B9HA) (CARBONATE ION - POTASSIUM CARBONATE) ON - CARBONATE				6 [hp_C] in 6 [hp_C]		
Inactive Ingredients						
Ingredient Name						Strength
SUCROSE (UNII: C151H8M554)						
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						
Product Characteristics						
Color	white	Score	9			
Shape	ROUND	Size				4mm

Flavor		Imprint C	Imprint Code					
C	ontains							
Ρ	ackaging							
#	ltem Code	Р	ackage Description	Marketing Start Date	Marketing End Date			
1	NDC:0220-2809- 41	6 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983				
Marketing Information								
	Marketing Category	Applic	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(0220-2809)					

Revised: 1/2023

Boiron