KALI SULPHURICUM- potassium sulfate 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 sulphuricum 200CK

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

Colds With Yellow Nasal Discharge*

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 SULPHURICUN	1				
potassium sulfate pellet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code	e (Source)	NDC:0220-2978	
Route of Administration	ORAL				
A . 11 . 1					
Active Ingredient/Active	Molety		Basis of		
Ingre	Strength				
				200 [kp_C] in 200 [kp_C]	
Inactive Ingredients					
	Strength				
LACTOSE, UNSPECIFIED FORM					
SUCROSE (UNII: C151H8M554)					
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	
	NDC:0220- 2978-41	200 [kp_C] in Product	1 TUBE; Type 0: Not a	a Combination	03/03/1983		
Marketing Information							
	U						
	Marketing Category	Applic	ation Number or N Citation	lonograph	Marketing Start Date	Marketing End Date	
	approved meopathic				03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-2978)			

Revised: 1/2023

Boiron