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 12C

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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).

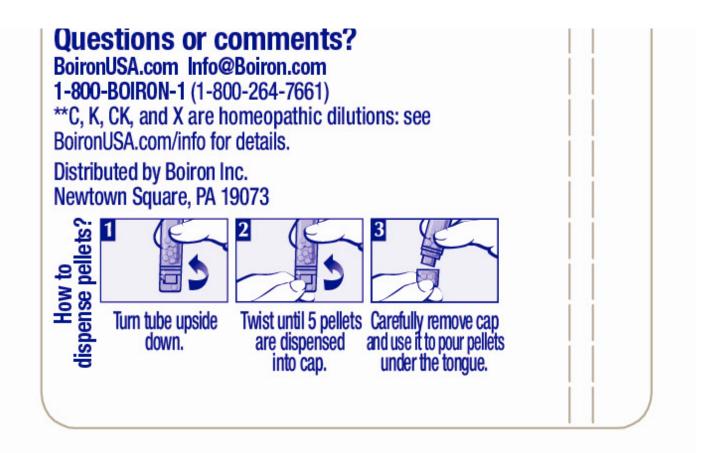
Uses: See symptoms on front panel.

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 SULPHURICU	М			
potassium sulfate pellet				
Product Information				
Product Type	HUMAN OTC DRUG	ltem Code	(Source)	NDC:0220-2975
Route of Administration	ORAL			
Active Ingredient/Active	e molety		Basis of	
Ingre	Strength			
POTASSIUM SULFATE (UNII: 1K573LC5TV) (POTASSIUM CATION - UNII:295053K152)			POTASSIUM SULFATE	12 [hp_C] in 12 [hp_C]
Inactive Ingredients				
	Strength			
SUCROSE (UNII: C151H8M554)				
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)			
Product Characteristics	5			
Color	white	Score		
Shape	ROUND	Size		4mm

Flavor		Imprint Code					
ntains							
Packaging							
Item Code	P	Package Description		Marketing Start Date	Marketing End Date		
	12 [hp_C] in Product	1 TUBE; Type 0: Not a	Combination	03/03/1983			
	Informe						
Marketing Information							
Marketing Category	Applic	ation Number or M Citation	lonograph	Marketing Start Date	Marketing End Date		
approved meopathic				03/03/1983			
	NDC:0220-2975- 41 arketing Category approved	Ackaging Item Code NDC:0220-2975- 41 Product Product Product Product Applic Category approved	Ackaging Package Description Item Code Package Description NDC:0220-2975- 41 12 [hp_C] in 1 TUBE; Type 0: Not a Product Iarketing Information Marketing Category Application Number or N Citation approved	Ackaging Package Description Item Code Package Description NDC:0220-2975- 41 12 [hp_C] in 1 TUBE; Type 0: Not a Combination Product Iarketing Application Number or Monograph Citation Marketing category Application Number or Monograph Citation	Ackaging Marketing Start Date Item Code Package Description Marketing Start Date NDC:0220-2975- 41 12 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 Iarketing Information Marketing Start Date 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(0220-2975)			

Revised: 11/2023

Boiron