X-RAY- alcohol, x-ray exposed (1000 rad) 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.

X-RAY 200CK

X-RAY 200CK

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

Itching or skin rash aggravated at night and in bed*

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).

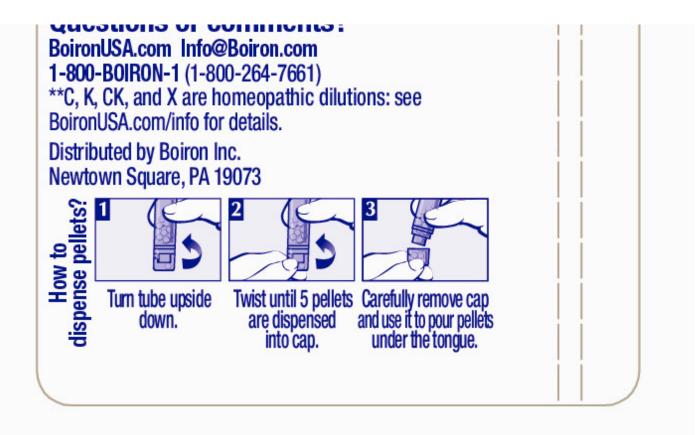
Uses: See symptoms on front panel.

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 Questions or comments?



X-RAY								
alcohol, x-ray exposed (1000 rad) pellet								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:0220-5345				
Route of Administration	ORAL							
Active Ingredient/Active	Moiety							
Ingredient Name Basis of Streng				gth	Strength			
ALCOHOL, X-RAY EXPOSED (1000 RAD) (UNII: 6PRJ93602P) (ALCOHOL, X- RAY EXPOSED (1000 RAD) - UNII:6PRJ93602P) ALCOHOL, X-RAY EXPOSED (1000 RAD) - UNII:6PRJ93602P)			D)	200 [kp_C] in 200 [kp_C]				
Inactive Ingredients								
Ingredient Name					Strength			
SUCROSE (UNII: C151H8M554)								
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)								
Product Characteristics								
Color	vhite	Score						
Shape F	ROUND	Size 4mm						
Flavor		Imprint Code						

Co	ontains							
Packaging								
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0220- 5345-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product	on 03/03/1983					
Marketing Information								
	Marketing Category	Application Number or Monograp Citation	h 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(0220-5345)				

Revised: 11/2023

Boiron