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 12C

X-RAY 12C

(**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

Principle Display Panel - X-ray HPUS 12C





Drug Facts

Active ingredient^{**}: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

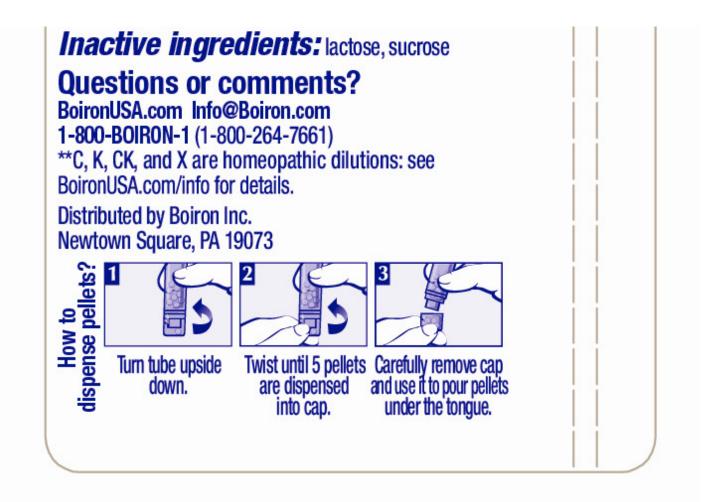
Uses: See symptoms on front panel.

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)



V DAV					
X-RAY					
alcohol, x-ray exposed (1000 r	rad) pellet				
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source) NDC:02		NDC:0220)-5343
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingre	Basis of St	rength	Strength		
ALCOHOL, X-RAY EXPOSED (100 EXPOSED (1000 RAD) - UNII:6PRJ936	ALCOHOL, X-RAY EXPOSED (1000 RAD)		12 [hp_C]		
Inactive Ingredients					
Ing	Strength				
SUCROSE (UNII: C151H8M554)					
LACTOSE (UNII: J2B2A4N98G)					
Product Characteristics					

Color		white	Score			
Shape		ROUND	Size		4mm	
Flavor			Imprint Cod			
Contains						
Packaging						
# Item Code	Pa	ackage Descriptio	on Marketing Start Date		Marketing End Date	
NDC:0220-5343-	80 in 1 TUBE; Type 0: Not a Combination Product			03/03/1983		
41	Product			03/03/190	5	
41	Product			03/03/190	5	
41	Product			03/03/190	3	
Marketing I		tion		03/03/190	5	
41	nforma	tion ation Number or N Citation	Monograpi		rketing Start Date	Marketing End Date

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment					
Name	Address	ID/FEI	Business Operations		
Boiron		282560473	manufacture(0220-5343)		

Revised: 11/2023

Boiron