## HISTAMINUM HYDROCHLORICUM- histamine dihydrochloride 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.

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## Histaminum hydrochloricum 30X

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

Allergies\*

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**<sup>\*\*</sup>: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

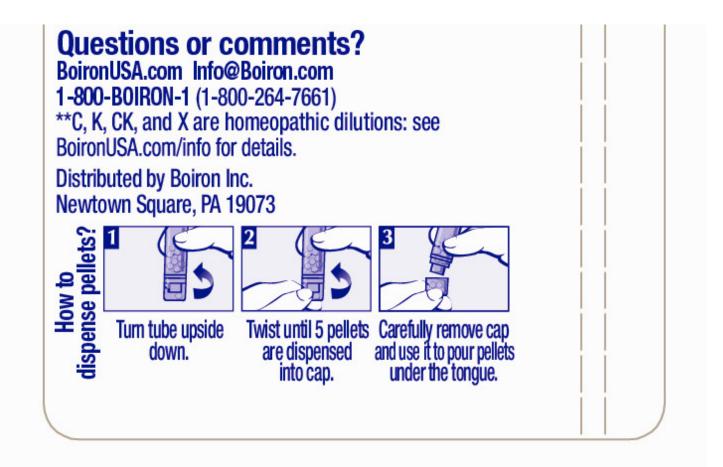
**Uses:** See symptoms on front panel.

**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



HISTAMINUM HYDF	ROCHLORICU	Μ		
histamine dihydrochloride pe	ellet			
Product Information				
Product Type	HUMAN OTC DRUG	Item Co	de (Source)	NDC:0220-2480
Route of Administration	ORAL			
Active Ingredient/Active	e Moiety			
Ingred	lient Name		<b>Basis of Strength</b>	n Strength
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - HISTAMINE UNII:820484N8I3) HISTAMINE		30 [hp_X] in 30 [hp_X]		
Inactive Ingredients				
-	Strength			
SUCROSE (UNII: C151H8M554)				
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)				
Product Characteristics				
Color	white	Score		
Shape	ROUND	Size		4mm
Flavor		Imprint Code		

Co	ntains			
Pa	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing Enc Date
	NDC:0220-2480- 41	30 [hp_X] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
Μ	arketing	Information		
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing Enc Date
	approved meopathic		03/03/1983	
			03/03/1983	

Labeler - Boiron (282560473)

## Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Boiron		282560473	manufacture(0220-2480)				

Revised: 10/2023

Boiron