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.

Histaminum hydrochloricum 9C

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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**: 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			
nistamine dihydrochloride pe	ellet				
Product Information					
Product Type	HUMAN OTC DRUG	ltem Cod	e (Source)	NDC:0220-2468	
Route of Administration	ORAL				
Active Ingredient/Active	e Moiety				
Ingred	th Strength				
HISTAMINE DIHYDROCHLORIDE (UNII: 3POA0Q644U) (HISTAMINE - HISTAMINE UNII:820484N8I3) HISTAMINE				9 [hp_C] in 9 [hp_C]	
Inactive Ingredients					
mactive myredients	Ingredient Na			Chuckath	
SUCROSE (UNII: C151H8M554)	Strength				
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)					
	(01111.)202/(111500)				
Product Characteristics					
Color	white	Score			
Shape	ROUND	Size		4mm	
Flavor		Imprint Code			

Co	ntains			
Pa	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
		9 [hp_C] in 1 TUBE; Type 0: Not a Combination		
	NDC:0220-2468- 41	Product	03/03/1983	
			03/03/1983	
•	41		03/03/1983	
	41	Product	05/03/1983	Marketing End Date

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-2468)				

Revised: 6/2023

Boiron