MURIATICUM ACIDUM- hydrochloric acid 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.

Muriaticum acidum 30C

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

Hives triggered by sun exposure*

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

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



MURIATICUM ACID	UM					
hydrochloric acid pellet						
Product Information						
Product Type	HUMAN OTC DRUG	HUMAN OTC DRUG Item Code (Source)		NDC:0220-3467		
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingre	Strength					
HYDROCHLORIC ACID (UNII: QTT17582CB) (HYDROCHLORIC ACID - HYDROCHLORIC ACID - UNII:QTT17582CB)			30 [hp_C] in 30 [hp_C]			
Inactive Ingredients						
	Strength					
SUCROSE (UNII: C151H8M554)						
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						
Product Characteristics						
Color	white	Score				
Shape	ROUND	Size		4mm		

Flavor		Im	Imprint Code			
ontains						
 						
Packaging						
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
NDC:0220-3467- 41	30 [hp_C] in Product	1 TUBE; Type 0: Not a Co	mbination	03/03/1983		
arkating	Informa	tion				
Marketing information						
Marketing Category	Applic	ation Number or Mo Citation	nograph	Marketing Start Date	Marketing End Date	
approved meopathic				03/03/1983		
	ackaging Item Code NDC:0220-3467- 41	ackaging Item Code NDC:0220-3467- 41 Barketing Informa Marketing Category Applic	Item Code Package Description NDC:0220-3467- 30 [hp_C] in 1 TUBE; Type 0: Not a Co A1 Product Iarketing Category Application Number or Mon Citation	Item Code Package Description NDC:0220-3467- 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product Iarketing Category Application Number or Monograph Citation	Item Code Package Description Marketing Start Date NDC:0220-3467- 41 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 Iterketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-3467)			

Revised: 4/2023

Boiron