HYDROFLUORICUM ACIDUM- hydrofluoric 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.

Hydrofluoricum acidum 6C

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

Intellectual fatigue with loss of hair and itching*

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



HYDROFLUORICUM	ACIDUM			
hydrofluoric acid pellet				
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Product Information				
Product Type	HUMAN OTC DRUG	ltem Code	e (Source)	NDC:0220-2561
Route of Administration	ORAL			
Active Ingredient/Active	e Moiety			
Ingre	Strength			
HYDROFLUORIC ACID (UNII: RGI UNII:Q80VPU4080)	6 [hp_C] in 6 [hp_C]			
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
Product Characteristics	;			
Color	white	Score		
Shape	ROUND	Size		4mm

Flavor		Imprint	Imprint Code						
C	ontains								
Packaging									
#	ltem Code	Ρ	Package Description		Marketing Start Date	Marketing End Date			
1	NDC:0220-2561- 41	6 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	on 0	3/03/1983				
Marketing Information									
	- Marketing Category	Applic	ation Number or Monogra Citation	ph	Marketing Start Date	Marketing End Date			

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-2561)				

Revised: 11/2023

Boiron