AURUM MURIATICUM NATRONATUM- sodium tetrachloroaurate 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.

Aurum muriaticum natronatum 9C

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

Warts, hemorrhoids.*

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



AURUM MURIATICU	JM NATRON	ATUI	М			
odium tetrachloroaurate pe	ellet					
Product Information						
Product Type	HUMAN OTC DRUG		Item Code (Source)		NDC:0220-0649	
Route of Administration	ORAL					
Active Ingredient/Active	e Moiety					
Ingredient Name Basis of Stren						Strength
SODIUM TETRACHLOROAURATE (UNII: 7FT6QUT299)SODIUM(TETRACHLOROAURATE ION - UNII:ZNL6IP5PJX)TETRACHLOROAUR					RATE	9 [hp_C] in 9 [hp_C]
Inactive Ingredients						
Ingredient Name						Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						
SUCROSE (UNII: C151H8M554)						
Due duet Chausete vistige						
Product Characteristics						
Color	white	Score	•			
Shape	ROUND	Size				4mm
Flavor		Impri	nt Code			

	ontains							
Packaging								
#	ltem Code	Pa	Package Description		Marketing Start Date	Marketing End Date		
	NDC:0220-0649- 41	9 [hp_C] in 1 1 Product	[hp_C] in 1 TUBE; Type 0: Not a Combination oduct		03/03/1983			
Μ	arketing	Informat	ion					
Μ	arketing Marketing Category		ion tion Number or Mo Citation	onograph	Marketing Start Date	Marketing End Date		
una	Marketing		tion Number or Mo	onograph	-	Marketing End Date		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0649)				

Revised: 11/2023

Boiron