AURUM MURIATICUM- gold trichloride 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 9C

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

Irritability with intolerance to noise*

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.

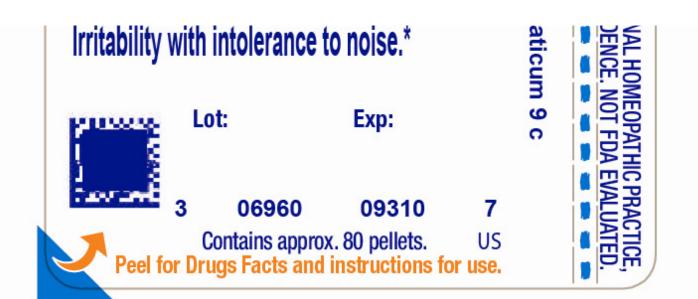
*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

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	ЈМ						
gold trichloride pellet							
Product Information							
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:0220-0674			
Route of Administration	ORAL						
Active Ingradient/Active	Maiaty						
Active Ingredient/Active	emolety		Basis of				
Ingre	Strength						
GOLD TRICHLORIDE (UNII: 1544 UNII:7XM25QYI14)	9 [hp_C] in 9 [hp_C]						
Inactive Ingredients							
	Strength						
LACTOSE, UNSPECIFIED FORM							
SUCROSE (UNII: C151H8M554)							
Product Characteristics	Product Characteristics						
Color	white	Score					
Shape	ROUND	Size		4mm			

Flavor		Imprint C	Imprint Code						
С	ontains								
Packaging									
#	ltem Code	Ρ	ackage Description	Marketing Start Date	Marketing End Date				
1	NDC:0220-0674- 41	9 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983					
Marketing Information									
	Marketing		ation Number or Monograp Citation	h Marketing Start Date	Marketing End Date				
	Category								

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0674)				

Revised: 11/2023

Boiron