AURUM METALLICUM- gold 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 metallicum 10M

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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.

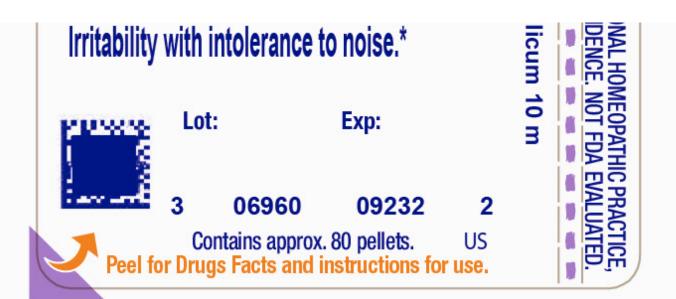
*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

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.

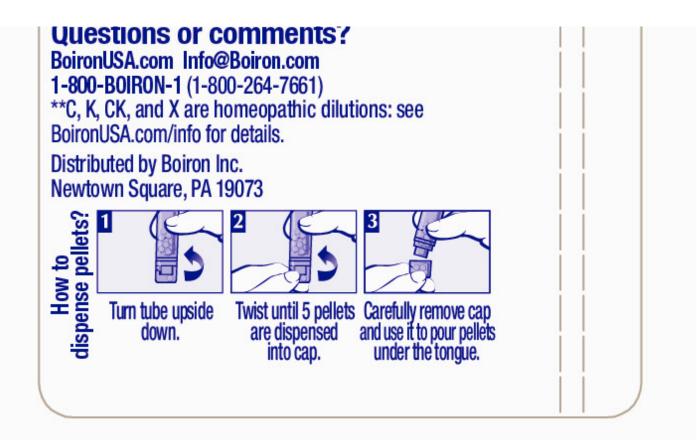
1-800-BOIRON-1 (1-800-264-7661), BoironUSA.com Info@boiron.com Distributed by Boiron, Inc. Newtown Square, PA 19073





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



Μ						
HUMAN OTC DRUG		ltem Code (Source)	1	NDC:0220-0671		
ORAL						
Moiety						
Name		Basis of Strength		Strength		
- UNII:79Y1949PYO)		GOLD	10 [hp	_M] in 10 [hp_M]		
Inactive Ingredients Ingredient Name Strength						
-						
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G) SUCROSE (UNII: C151H8M554)						
white	Score					
ROUND	Size			4mm		
	ORAL Moiety Name - UNII: 79Y1949PYO) Ingredient Na (UNII: J2B2A4N98G) white	HUMAN OTC DRUG ORAL • Moiety Name - UNII:79Y1949PYO) • Ingredient Name (UNII: J2B2A4N98G) • UNII: Socre	HUMAN OTC DRUG Item Code (Source) ORAL Item Code (Source) ORAL Basis of Strength Name Basis of Strength OUNII:79Y1949PYO) GOLD	HUMAN OTC DRUG Item Code (Source) ORAL Moiety Name Basis of Strength OUNII: 79Y1949PYO) GOLD 10 [hp (UNII: j2B2A4N98G)		

Contains							
Packaging							
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:0220-0671- 41	10 [hp_M] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983				
Marketing Information							
	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	approved meopathic		03/03/1983				

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0671)				

Revised: 2/2023

Boiron