CUPRUM METALLICUM- copper 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.

Cuprum metallicum 10M

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

Sudden muscle cramps*

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



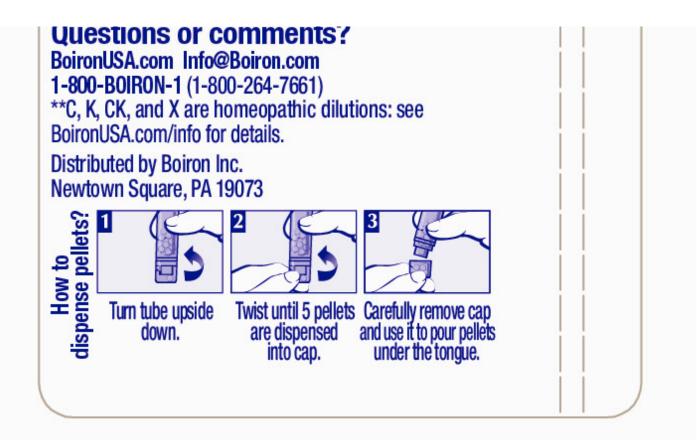


Drug Facts Active ingredient**: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before 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



CUPRUM METALLICUM							
copper pellet							
Product Information							
	HUMAN OT		tem Code (Source)	NDC:0220-1651			
Product Type		C DRUG	tem Code (Source)	NDC:0220-1051	•		
Route of Administration	ORAL						
Active Ingredient/Activ	ve Moiety						
Active migreatend/Activ							
•	-		Basis of Strength	Strength			
-	ent Name	89U1901C5)	Basis of Strength COPPER	Strength	_M]		
Ingredie	ent Name	89U1901C5)		_	_M]		
Ingredie	ent Name	39U1901C5)		_	9_M]		
Ingredie COPPER (UNII: 789U1901C5) (C	opper - UNII:78	89U1901C5) ent Name		_	_		
Ingredie COPPER (UNII: 789U1901C5) (C	opper - UNII:78			10 [hp_M] in 10 [hp	_		
Ingredie COPPER (UNII: 789U1901C5) (C Inactive Ingredients	ent Name OPPER - UNII:78 Ingredie	ent Name		10 [hp_M] in 10 [hp	_		
Ingredie COPPER (UNII: 789U1901C5) (C Inactive Ingredients SUCROSE (UNII: C151H8M554)	ent Name OPPER - UNII:78 Ingredie	ent Name		10 [hp_M] in 10 [hp	_		
Ingredie COPPER (UNII: 789U1901C5) (C Inactive Ingredients SUCROSE (UNII: C151H8M554)	ent Name OPPER - UNII:78 Ingredie M (UNII: J2B2A4	ent Name		10 [hp_M] in 10 [hp	_		
Ingredie COPPER (UNII: 789U1901C5) (C Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FOR Product Characteristic	ent Name OPPER - UNII:78 Ingredie M (UNII: J2B2A4	ent Name		10 [hp_M] in 10 [hp	_		
Ingredie COPPER (UNII: 789U1901C5) (C Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FOR	ent Name OPPER - UNII:78 Ingredia M (UNII: J2B2A4	ent Name		10 [hp_M] in 10 [hp	_		

Co	ontains						
Packaging							
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:0220-1651- 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(0220-1651)					

Revised: 11/2023

Boiron