BERYLLIUM METALLICUM- beryllium 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.

Beryllium metallicum 30C

Beryllium metallicum 30C

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

Painful dry cough*

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



BERYLLIUM METALLICUM							
beryllium pellet							
Product Information							
Product Type	HUMAN OTC DRUG	ltem	Code (Source)	NDC:0220-0839			
Route of Administration	ORAL						
Active Ingredient/Active Moiety							
Ingredient Name Basis of Strength				Strength			
ingream							
BERYLLIUM (UNII: OW5102UV6N)		5102UV6N)	BERYLLIUM	30 [hp_C] in 30 [hp_C]			
-		5102UV6N)		30 [hp_C] in 30 [hp_C]			
BERYLLIUM (UNII: OW5102UV6N)		5102UV6N)		30 [hp_C] in 30 [hp_C]			
-		5102UV6N)		30 [hp_C] in 30 [hp_C]			
BERYLLIUM (UNII: OW5102UV6N)				30 [hp_C] in 30 [hp_C] Strength			
BERYLLIUM (UNII: OW5102UV6N)	(BERYLLIUM - UNII:OW						
BERYLLIUM (UNII: OW5102UV6N)	(BERYLLIUM - UNII:OW						
BERYLLIUM (UNII: OW5102UV6N) Inactive Ingredients SUCROSE (UNII: C151H8M554)	(BERYLLIUM - UNII:OW						
BERYLLIUM (UNII: OW5102UV6N) Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FORM	(BERYLLIUM - UNII:OW Ingredient Nat						
BERYLLIUM (UNII: OW5102UV6N) Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FORM	(BERYLLIUM - UNII:OW Ingredient Nat I (UNII: J2B2A4N98G)	me					
BERYLLIUM (UNII: OW5102UV6N) Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FORM Product Characteristics Color	(BERYLLIUM - UNII:OW Ingredient Nat I (UNII: J2B2A4N98G)	me Score					
BERYLLIUM (UNII: OW5102UV6N) Inactive Ingredients SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FORM	(BERYLLIUM - UNII:OW Ingredient Nat I (UNII: J2B2A4N98G)	me					

С	ontains								
P	Packaging								
#	Item Code	Package Description	Marketing Start Date	Marketing End Date					
1	NDC:0220-0839- 41	30 [hp_C] 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(0220-0839)				

Revised: 11/2023

Boiron