BRYONIA- bryonia alba root 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.

Bryonia 10M

Bryonia 10M

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

Muscle and joint pain improved by rest*

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).

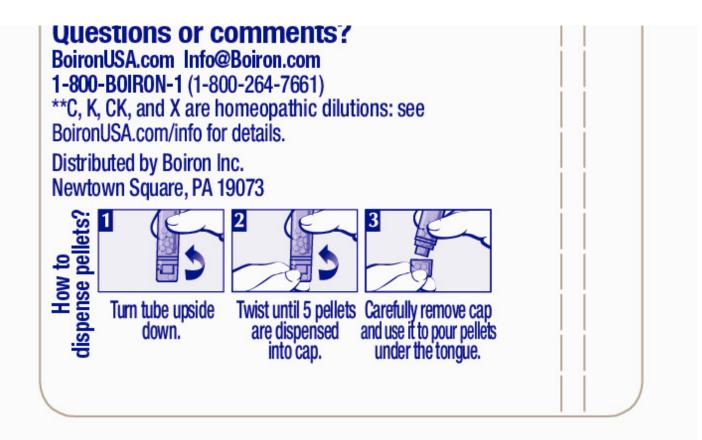
Uses: See symptoms on front panel.

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



BRYONIA				
oryonia alba root pellet				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:0220-0947
Route of Administration	ORAL			
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Active Ingredient/Active	e Molety			
Ingre	Strength			
BRYONIA ALBA ROOT (UNII: T7J046YI2B) (BRYONIA ALBA ROOT - UNII:T7J046YI2B)			BRYONIA ALBA ROOT	10 [hp_M] in 10 [hp_M]
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
Product Characteristics	5			
Color	white	Score		
Shape	ROUND	Size		4mm

	Flavor		Imprint Code				
ntains							
Packaging							
Item Code	Р	Package Description		Marketing Start Date	Marketing End Date		
NDC:0220-0947- 41	10 [hp_M] in Product			03/03/1983			
arkating	Informa	tion					
Marketing information							
Marketing Category	Applic	ation Number or N Citation	Ionograph	Marketing Start Date	Marketing End Date		
approved meopathic				03/03/1983			
	Item Code NDC:0220-0947- 41 arketing Category approved	Item Code P NDC:0220-0947- 10 [hp_M] in Product 41 Product arketing Informa Applic Category Approved Applic Category	Item Code Package Descripti NDC:0220-0947- 41 10 [hp_M] in 1 TUBE; Type 0: Not a Product arketing Category Information Marketing Category Application Number or N Citation	Ackaging Package Description Item Code Package Description NDC:0220-0947- 41 10 [hp_M] in 1 TUBE; Type 0: Not a Combination Product arketing Application Number or Monograph Citation Appproved Application Number or Monograph	Ackaging Marketing Start Date Item Code Package Description Marketing Start Date NDC:0220-0947- 41 10 [hp_M] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 arketing Information Mumber or Monograph Citation Number or Monograph Date Marketing Category Application Number or Monograph Citation 03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-0947)			

Revised: 11/2023

Boiron