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 5C

Bryonia 5C

(**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).

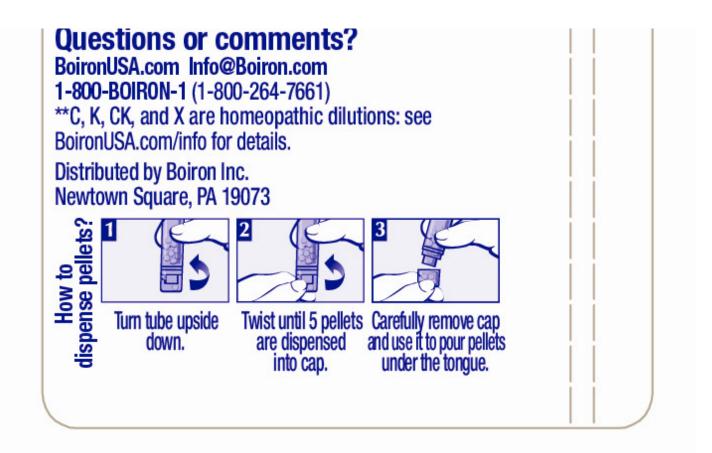
Uses: See symptoms on front panel.

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									
bryonia alba root pellet									
Product Information									
Product Type	HUMAN OTC DRUG Item Code (Source)		NDC:0220-0939						
Route of Administration	ORAL								
Active Ingredient/Active									
Ingred	Strength								
BRYONIA ALBA ROOT (UNII: T7J04 UNII:T7J046YI2B)	T 5 [hp_C] in 5 [hp_C]								
Inactive Ingredients									
	Strength								
LACTOSE, UNSPECIFIED FORM (
SUCROSE (UNII: C151H8M554)									
Product Characteristics									
Color	vhite	Score							
Shape R	OUND	Size		4mm					

Flavor		Imprint C	Imprint Code						
C	ontains								
Packaging									
#	ltem Code	Ρ	ackage Description	Marketing Start Date	Marketing End Date				
1	NDC:0220-0939- 41	5 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983					
Marketing Information									
	Marketing Category	Applic	ation Number or Monograp Citation	h Marketing Start Date	Marketing End Date				
	approved			03/03/1983					

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0939)				

Revised: 11/2023

Boiron