## HELLEBORUS NIGER- helleborus niger 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.

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## Helleborus niger 200CK

Helleborus niger 200CK

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

Nausea, vomiting, diarrhea\*

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

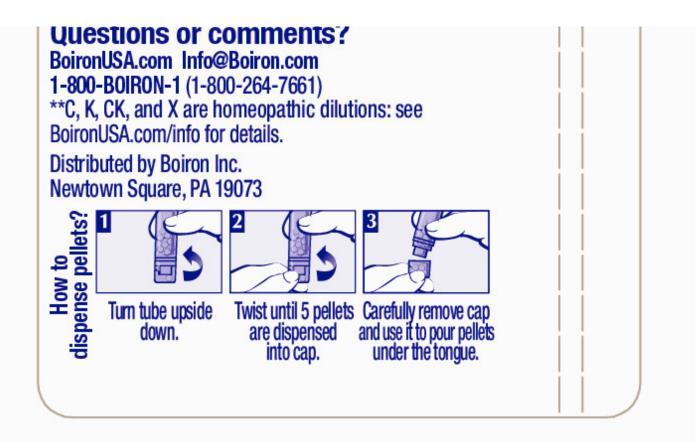




## Drug Facts Active ingredient\*\*: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional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



<b>HELLEBORUS NIGER</b>					
helleborus niger root pellet					
Product Information					
Product Type	HUMAN PRESCRIPTION DRUG		ltem Code (Source)		NDC:0220-2450
Route of Administration	ORAL				
<b>.</b>					
Active Ingredient/Active	Molety				
Ingred	Basis of Strength	Strength			
HELLEBORUS NIGER ROOT (UNII - UNII:608DGJ6815)	608DGJ6815) (HELLE	200 [kp_C] in 200 [kp_C]			
Inactive Ingredients					
	Strength				
SUCROSE (UNII: C151H8M554)					
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)					
Product Characteristics					
Color	vhite	Score			
Shape F	OUND	Size			4mm

Fla	avor			Imprint Cod	e	
Contains						
D	<b>I </b>					
Packaging						
#	ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date
1	NDC:0220- 2450-41	200 [kp_C] in Product	1 TUBE; Type 0: Not a	a Combination	03/03/1983	
М	larketing	Informa	tion			
	<b>U</b>					
	Marketing Category	Applic	ation Number or I Citation	Monograph	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(0220-2450)			

Revised: 7/2023

Boiron