BERBERIS VULGARIS- berberis vulgaris root bark 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.

Berberis vulgaris 200CK

Berberis vulgaris 200CK

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

Skin rash with round, red patches and thin flakes*

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



BERBERIS VULGARI	S						
berberis vulgaris root bark pellet							
Product Information							
Product Type	HUMAN OTC DRUG	1	Item Code (Source)		NDC:0220-0837		
Route of Administration	ORAL						
Active Ingredient/Active Moiety							
Ingredient Name Basis of Streng						Strength	
BERBERIS VULGARIS ROOT BARK (UNII: 1TH8Q20J0U) (BERBERISBERBERIS VULGARISVULGARIS ROOT BARK - UNII:1TH8Q20J0U)ROOT BARK			S	200 [kp_C] in 200 [kp_C]			
Inactive Ingredients							
Ingredient Name						Strength	
SUCROSE (UNII: C151H8M554)							
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)						
Product Characteristics							
Color	white	Score					
Shape	ROUND	Size				4mm	
Flavor		Imprin	t Code				

	ontains			
Pa	ackaging			
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220- 0837-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
	arkating			
M	arketing	Information		
M	Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-0837)					

Revised: 1/2023

Boiron