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 1M

Berberis vulgaris 1M

(**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).

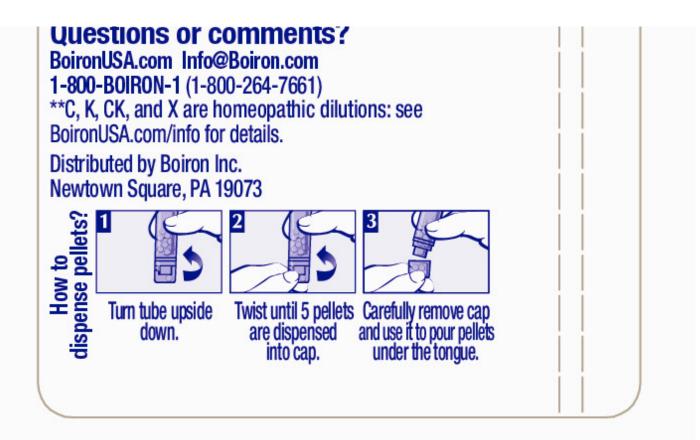
Uses: See symptoms on front panel.

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 VULGAR	IS						
berberis vulgaris root bark p	ellet						
Product Information							
Product Type	HUMAN OTC DRUG	Item Code (Item Code (Source)		NDC:0220-0841		
Route of Administration	ORAL						
Active Ingredient/Active Moiety							
Ingredient Name Basis of Stren					Strength		
			BERBERIS VULGARIS ROOT BARK		1 [hp_M] in 1 [hp_M]		
Inactive Ingredients							
Ingredient Name					Strength		
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)							
SUCROSE (UNII: C151H8M554)							
Product Characteristics	i						
Color	white	Score					
Shape	ROUND	Size		4	4mm		
Flavor		Imprint Code					

ntains			
ckaging			
ltem Code	Package Description	Marketing Start Date	Marketing End Date
	1 [hp_M] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
arketing	Information		
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing Enc Date
pproved neopathic		03/03/1983	
	ckaging Item Code NDC:0220-0841- 41 arketing Category pproved	ckaging Item Code Package Description NDC:0220-0841- 1 [hp_M] in 1 TUBE; Type 0: Not a Combination Product arketing Application Number or Monograph Citation pproved Image: Comparison of the section of t	ckaging Marketing Start Date Item Code Package Description Marketing Start Date NDC:0220-0841- 1 [hp_M] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 arketing Information 03/03/1983 Marketing Category Application Number or Monograph Citation Marketing Start Date pproved 03/03/1983

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-0841)			

Revised: 2/2023

Boiron