ARUNDO MAURITANICA- arundo pliniana 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.

Arundo mauritanica 30C

Arundo mauritanica 30C HPUS

Active ingredient**: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

Relieves itching of the nose and the palate *

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.

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.

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

BoironUSA.com Info@boiron.com

1-800-BOIRON-1 (1-800-264-7661) 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).

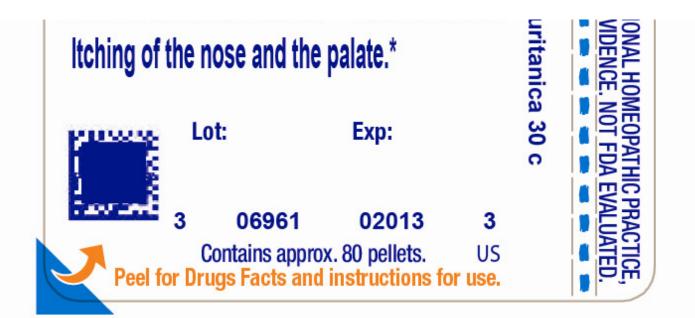
Uses: See symptoms on front panel.

Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breast-feeding, 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.







Drug Facts

Active ingredient***: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

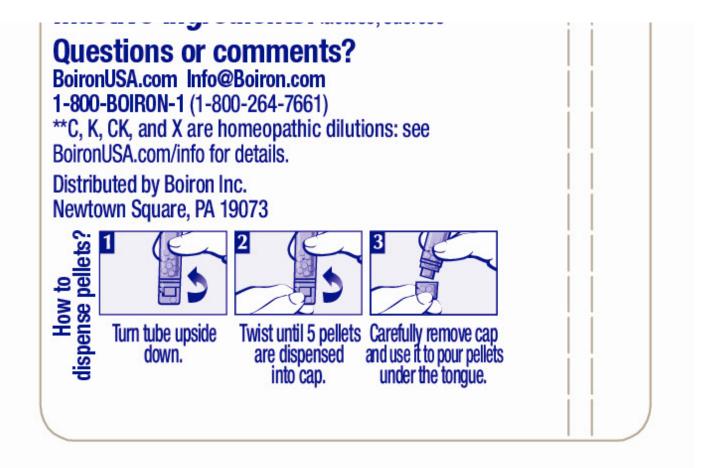
Uses: See symptoms on front panel.

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 incredients: lactose. sucrose



ARUNDO MAURITAN					
arundo pliniana root pellet					
Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:0220-0569	
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
-	Strongth				
Ingre	Strength				
ARUNDO PLINIANA ROOT (UNII: ZXE7LB03WC) (ARUNDO PLINIANA ROOT - ARUNDO PLINIANA ROOT - JNII:ZXE7LB03WC) ARUNDO PLINIANA ROOT			30 [hp_C] in 30 [hp_C]		
Inactive Ingredients					
Ingredient Name					Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)					
SUCROSE (UNII: C151H8M554)					
Product Characteristics					
Color	vhite	Score	3		

Shape ROUND		ROUND	Size	4mm			
Flavor Imj		Imprint Cod	mprint Code				
Contains							
Packaging							
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date		
NDC:0220-0569- 41	30 [hp_C] in Product			03/03/1983			
Marketing Information							
Marketing Category	Applic	ation Number or N Citation	Aonograph	Marketing Start Date	Marketing End Date		
	avor ontains ackaging Item Code NDC:0220-0569- 41 Iarketing Marketing	avor ontains ackaging item Code P NDC:0220-0569- 30 [hp_C] in 41 Product Iarketing Informa Marketing Applic	avor ontains ackaging Item Code Package Descripti NDC:0220-0569- 30 [hp_C] in 1 TUBE; Type 0: Not a 41 Product Product	avor attem code NDC:0220-0569- 41 Application Number or Monograph	avor Imprint Code ontains Imprint Code ackaging NDC:0220-0569- 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product Marketing Start Date NDC:0220-0569- 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0569)				

Revised: 5/2024

Boiron