ANAGALLIS ARVENSIS- anagallis arvensis 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.

Anagallis arvensis 30C

Anagallis arvensis 30C HPUS

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

Uses: See symptoms on front panel.

Relieves skin rash with intense itching *

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.

lactose, sucrose

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.

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).

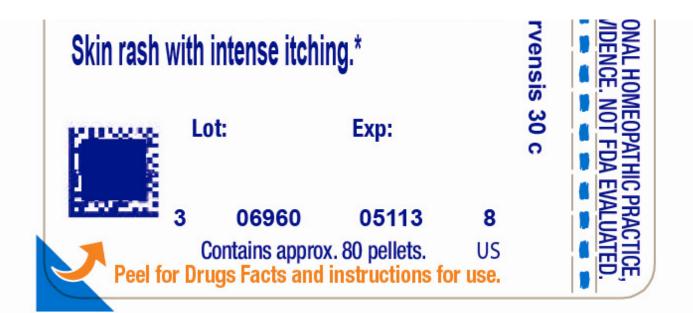
Uses: See symptoms on front panel.

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).

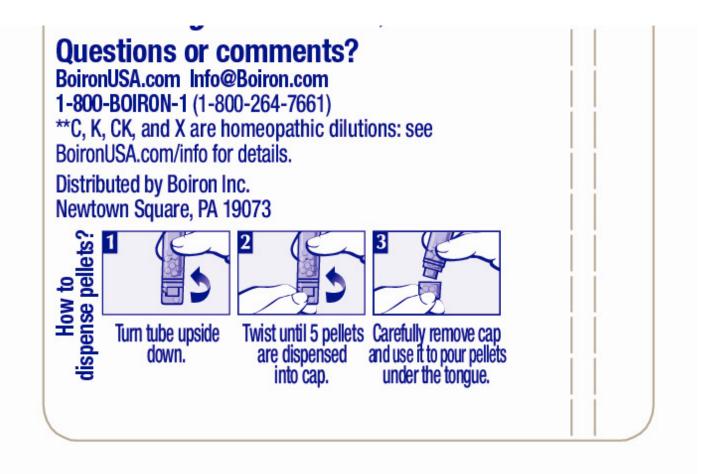
Uses: See symptoms on front panel.

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 ingredients: lactose, sucrose



ANAGALLIS ARVEN	SIS					
anagallis arvensis pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:0220-0312		
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingre	Strength					
ANAGALLIS ARVENSIS (UNII: 46883LR90E) (ANAGALLIS ARVENSIS - UNII:46883LR90E)			ANAGALLIS ARVENSIS	30 [hp_C] in 30 [hp_C]		
Inactive Ingredients						
Inactive Ingredients	Ingredient Na	me		Strength		
-		me		Strength		
LACTOSE, UNSPECIFIED FORM		me		Strength		
Inactive Ingredients LACTOSE, UNSPECIFIED FORM SUCROSE (UNII: C151H8M554)		me		Strength		
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)	me		Strength		

			Size		4mm	
avor	Imprint (Imprint Cod	de		
ntains						
Packaging						
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
NDC:0220-0312- 41	30 [hp_C] in Product			03/03/1983		
	£	••				
Marketing Information						
Marketing Category	Applic	ation Number or I Citation	Monograph	Marketing Start Date	Marketing End Date	
approved meopathic				03/03/1983		
	Ackaging Item Code NDC:0220-0312- 41 Arketing Category Approved	Ackaging Item Code NDC:0220-0312- 41 Broduct Antering Marketing Category Applic	Ackaging Package Descripti Item Code Package Descripti NDC:0220-0312- 30 [hp_C] in 1 TUBE; Type 0: Not a 41 Product arketing Application Number or I Category Application Number or I approved Image: Construct of the second	Ackaging Package Description Item Code Package Description NDC:0220-0312- 41 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product arketing Category Application Number or Monograph Citation	Ackaging Marketing Start Date Item Code Package Description Marketing Start Date NDC:0220-0312- 41 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 arketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date Approved 03/03/1983 03/03/1983	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0312)				

Revised: 5/2024

Boiron