## EQUISETUM ARVENSE- equisetum arvense top 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.

-----

## Equisetum arvense 30C

Equisetum arvense 30C

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

Bedwetting\*

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>EQUISETUM ARVEN</b>	SE			
equisetum arvense top pelle	et			
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (S	Source)	NDC:0220-1893
Route of Administration	ORAL			
<b>Active Ingredient/Active</b>	e Moiety			
Ingr	Strength			
EQUISETUM ARVENSE TOP (UN - UNII:1DP6Y6B65Z)	II: 1DP6Y6B65Z) (EQU	30 [hp_C] in 30 [hp_C]		
Inactive Ingredients				
	Strength			
LACTOSE, UNSPECIFIED FORM				
SUCROSE (UNII: C151H8M554)				
<b>Product Characteristics</b>	;			
Color	white	Score		
Shape	ROUND	Size		4mm

Flavor		Imprint Code					
ntains							
Packaging							
Item Code	P	Package Description		Marketing Start Date	Marketing End Date		
	30 [hp_C] in 3 Product			03/03/1983			
Marketing Information							
<b>U</b>							
Marketing Category	Applic	ation Number or I Citation	Monograph	Marketing Start Date	Marketing End Date		
approved meopathic				03/03/1983			
	Item Code NDC:0220-1893- 41 Iarketing Marketing Category approved	ackaging         Item Code       Pathods         NDC:0220-1893-       30 [hp_C] in 1         41       Product         Iarketing       Information         Marketing       Applic         Category       Applic         approved       Information	Ackaging       Package Descripti         Item Code       Package Descripti         NDC:0220-1893- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Product         Iarketing Information         Marketing Category approved       Application Number or Point	Ackaging       Package Description         Item Code       Package Description         NDC:0220-1893- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product         Iarketing       Application Number or Monograph Citation         Marketing Category       Application Number or Monograph Citation	Ackaging       Marketing Start Date         Item Code       Package Description       Marketing Start Date         NDC:0220-1893- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product       03/03/1983         Iarketing Comparison       Application Number or Monograph Citation       Marketing Start Date         Marketing category       Application Number or Monograph Citation       Marketing Start Date		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-1893)			

Revised: 11/2023

Boiron