## FUMARIA OFFICINALIS- fumaria officinalis flowering 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.

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## Fumaria officinalis 6X

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

Dry skin rash with 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

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).

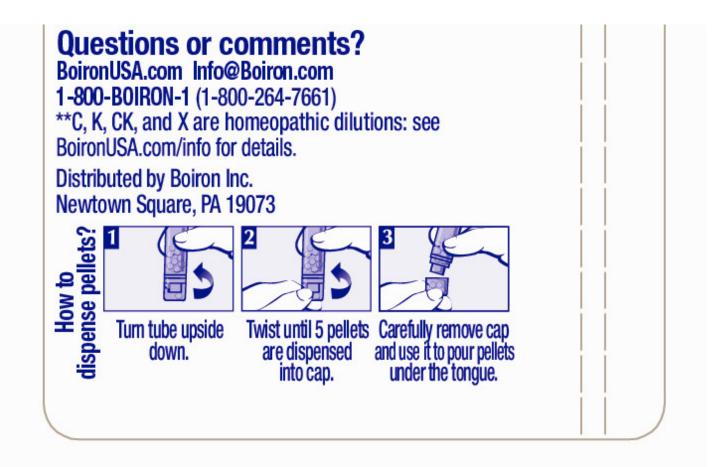
**Uses:** See symptoms on front panel.

**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>FUMARIA OFFICINA</b>	LIS							
fumaria officinalis flowering top pellet								
Product Information								
Product Type	HUMAN OTC DRUG		Item Code (Source)		NDC:0220-2176			
Route of Administration	ORAL							
Active Ingredient/Active Moiety								
Ingredient Name Basis of Stre						Strength		
FUMARIA OFFICINALIS FLOWERING TOP (UNII: VH659J61ZL) (FUMARIAFUMARIA OFFICINOFFICINALIS FLOWERING TOP - UNII: VH659J61ZL)FLOWERING TOP						6 [hp_X] in 6 [hp_X]		
Inactive Ingredients								
Ingredient Name						Strength		
SUCROSE (UNII: C151H8M554)								
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)								
Product Characteristics								
	white	Score						
	ROUND	Size			4n	nm		
Flavor		Imprin	nt Code					

CU	ntains					
Pa	ackaging					
# Item Code Package Description					Marketing Start Date	Marketing End Date
	NDC:0220-2176- 41	6 [hp_X] in 1 Product	TUBE; Type 0: N	ot a Combination	03/03/1983	
		TTOULLE				
			tion			
	arketing Marketing Category	Informa		r or Monograph on	Marketing Start Date	Marketing End Date

Labeler - Boiron (282560473)

## Registrant - Boiron Inc. (014892269)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Boiron		282560473	manufacture(0220-2176)				

Revised: 7/2023

Boiron