#### ARSENICUM SULPHURATUM FLAVUM- arsenic trisulfide 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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## Arsenicum sulphuratum flavum 200CK

Arsenicum sulphuratum flavum 200CK HPUS

Less than 10<sup>-12</sup> mg arsenic per pellet

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 dry cough aggravated by cold \*

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

## BoironUSA.com Info@boiron.com

**1-800-BOIRON-1** (1-800-264-7661) Distributed by Boiron, Inc. Newtown Square, PA 19073

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.



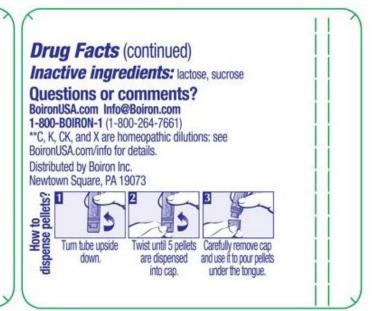
## **Drug Facts**

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**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.



ARSENICUM SULPHURATUM FLAVUM arsenic trisulfide pellet							
HUMAN OTC DRUG	Item Code	e (Source)	NDC:0220-0602				
ORAL							
Active Ingredient/Active Moiety							
Ingredient Name							
ARSENIC TRISULFIDE (UNII: 44SIJ800OX) (ARSENIC CATION (3+) - UNII:C96613F5AV)			200 [kp_C] in 200 [kp_C]				
	HUMAN OTC DRUG ORAL Moiety ient Name	HUMAN OTC DRUG Item Code ORAL Moiety ient Name	HUMAN OTC DRUG Item Code (Source) ORAL Moiety ient Name Basis of Strength				

Inactive Ingredients							
Ingredient Name						Strength	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)							
รบ	CROSE (UNII: C	151H8M554)					
Dr	oduct Char	actoristics	_				
	lor	acteristics	<b>&gt;</b> white	Score			
	ape		ROUND	Size		4mm	
	avor		NOOND	Imprint Co	de		
	ntains						
cu	incams						
Packaging							
#	ltem Code	P	ackage De	escription	Marketing Start Date	Marketing End Date	
	NDC:0220- 0602-41	200 [kp_C] in Product	1 TUBE; Type	e 0: Not a Combination	01/01/2024		
Marketing Information							
	Marketing Category	Applic		ber or Monograph ation	Marketing Start Date	Marketing End Date	
una	approved				01/01/2024		

# Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	<b>Business Operations</b>					
Boiron		282560473	manufacture(0220-0602)					

Revised: 4/2024

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Boiron