## SULPHUR- sulfur 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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## Sulphur 200X

Sulphur 200X (\*\*contains 0.443 mg of the active ingredient per pellet)

Skin Rash Worsened By Heat And Water\*

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**<sup>\*\*</sup>: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

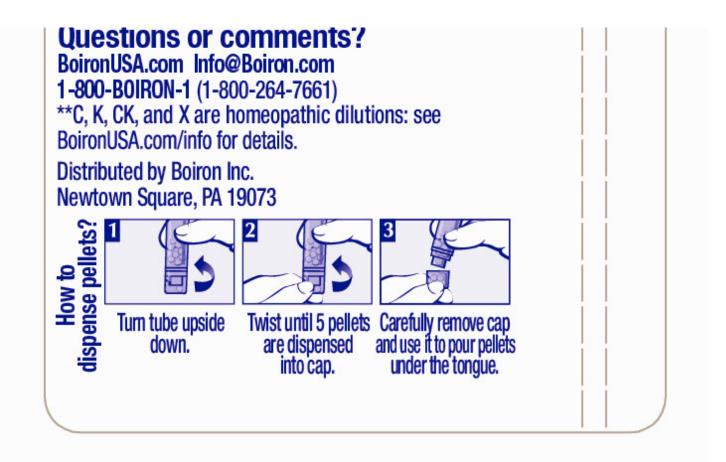
**Uses:** See symptoms on front panel.

**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



SULPHUR sulfur pellet								
<b>Product Information</b>								
Product Type	HUMAN OTC DRUG	ľ	tem Code (Source)		NDC:0220-4813			
Route of Administration	ORAL							
Active Ingredient/Active Moiety								
Ingredien	t Name		<b>Basis of Strength</b>		Strength			
SULFUR (UNII: 70FD1KFU70) (SUI	FUR - UNII:70FD1KFU70	))	SULFUR	200 [h	ıp_X] in 200 [hp_X]			
Inactive Ingredients								
	Ingredient Na	ne			Strength			
SUCROSE (UNII: C151H8M554)	Ingredient Nar	ne			Strength			
SUCROSE (UNII: C151H8M554) LACTOSE, UNSPECIFIED FORM	-	ne			Strength			
	-	ne			Strength			
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)	ne			Strength			
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)	ne Score			Strength			
LACTOSE, UNSPECIFIED FORM Product Characteristics Color	(UNII: J2B2A4N98G)				Strength 4mm			
	(UNII: J2B2A4N98G)	Score Size	t Code					

Packaging					
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date	
	NDC:0220- 4813-41	200 [hp_X] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983		
M	arketing	Information			
M	arketing Marketing Category	Information Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-4813)					

Revised: 10/2023

Boiron