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.

Sulphur 6X

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(**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.

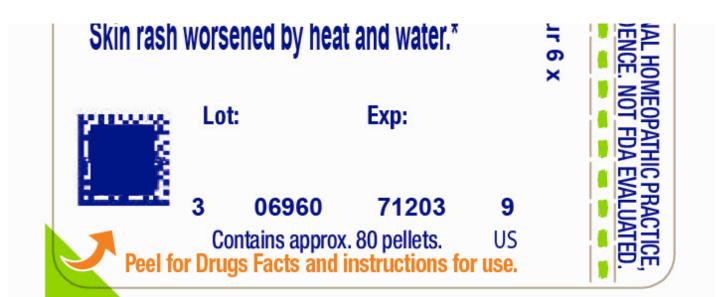
*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

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

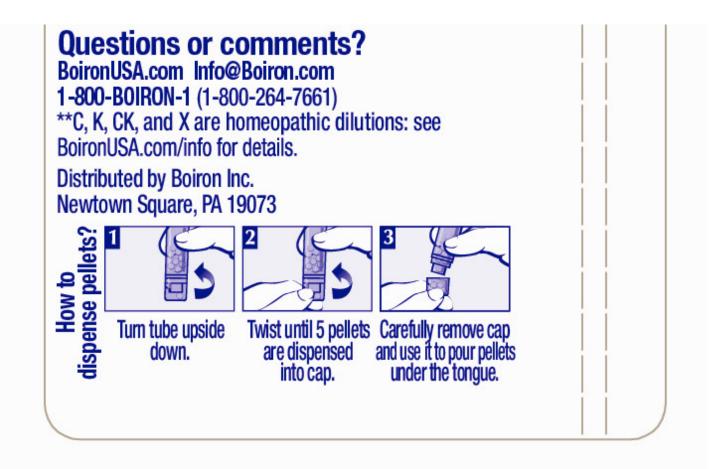
Uses: See symptoms on front panel.

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							
Product Information							
Product Type	HUMAN OTC DRUG	Item Co	de (Source)	NDC:02	220-4823	
Route of Administration	ORAL						
Active Ingredient/Active	Moiety						
Ingred	Basis o	sis of Strength		Strength			
SULFUR (UNII: 70FD1KFU70) (SULFUR - UNII:70FD1KFU70)			SULFUR			6 [hp_X]	
Inactive Ingredients							
						rength	
SUCROSE (UNII: C151H8M554)							
LACTOSE (UNII: J2B2A4N98G)							
Product Characteristics							
00101	white	Score					
Shape	ROUND	Size			4	1mm	
Flavor		Imprint Code					

Сс	ontains					
Packaging						
#	ltem Code	Package Description	Marketing Start Date		Marketing End Date	
1	NDC:0220-4823- 41	80 in 1 TUBE; Type 0: Not a Combination Product	03/03/1983			
Μ	larketing l	nformation				
Marketing Information						
	Marketing Category	Application Number or Monograpl Citation		Marketing Start Date	Marketing End Date	
	approved meopathic			03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron Inc. (014892269)

Establishment						
Name	Address	ID/FEI	Business Operations			
Boiron		282560473	manufacture(0220-4823)			

Revised: 11/2023

Boiron