## NATRUM SULPHURICUM- sodium sulfate 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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## Natrum sulphuricum 12C

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

Bronchial irritation worsened by humidity\*

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

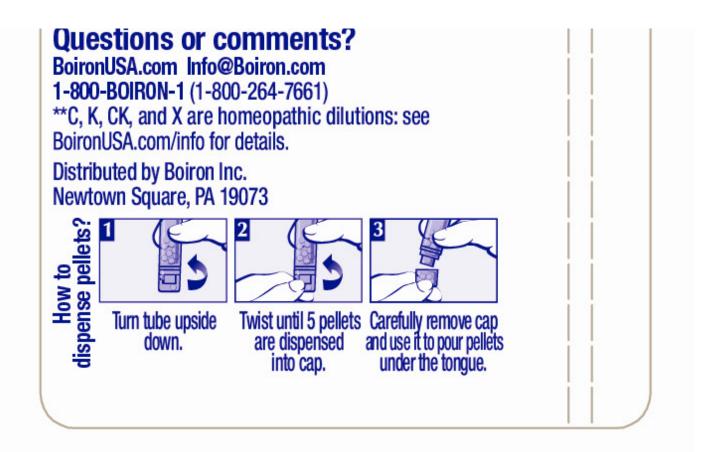




## Drug Facts Active ingredient\*\*: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional 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



NATRUM SULPHUR	ICUM					
sodium sulfate pellet						
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Product Information						
Product Type	HUMAN OTC DRUG		Item Code (S	iource)	NDC:0220-3635	
Route of Administration	ORAL					
<b>Active Ingredient/Active</b>	Moiety					
Ingre	Strength					
SODIUM SULFATE (UNII: 0YPR65) UNII:36KCS0R750)	12 [hp_C] in 12 [hp_C]					
Inactive Ingredients						
	Strength					
LACTOSE, UNSPECIFIED FORM						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						
Color	white	Score	2			
Shape	ROUND	Size			4mm	

Flavor		Im	Imprint Code				
Contains							
Packaging							
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date		
	12 [hp_C] in Product	] in 1 TUBE; Type 0: Not a Combination		03/03/1983			
arkating	Informa	tion					
Marketing mormation							
Marketing Category	Applic	olication Number or Monograph Citation		Marketing Start Date	Marketing End Date		
approved meopathic				03/03/1983			
	ackaging Item Code NDC:0220-3635- 41 Iarketing Category approved	ackaging Item Code NDC:0220-3635- 41 Product Iarketing Informa Marketing Category approved	Item Code Package Description   NDC:0220-3635- 12 [hp_C] in 1 TUBE; Type 0: Not a Co   A1 Product     Iarketing Application Number or Mon   Marketing Application Number or Mon   Category Application Number or Mon	Item Code Package Description   NDC:0220-3635- 12 [hp_C] in 1 TUBE; Type 0: Not a Combination Product   Iarketing Category Application Number or Monograph Citation	Item Code       Package Description       Marketing Start Date         NDC:0220-3635-       12 [hp_C] in 1 TUBE; Type 0: Not a Combination Product       03/03/1983         Iterketing Information         Marketing Category         Application Number or Monograph Citation       Marketing Start Date         03/03/1983       03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-3635)				

Revised: 11/2023

Boiron