

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.

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



Bronchial irritation worsened by humidity.*



Lot:

Exp:

3

06960

51311

7

Contains approx. 80 pellets.

US

Peel for Drugs Facts and instructions for use.

uricum 12 c

VAL HOMEOPATHIC PRACTICE,
INC. NOT FDA EVALUATED.

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

Questions or comments?

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Carefully remove cap
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NATRUM SULPHURICUM

sodium sulfate pellet

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0220-3635
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SODIUM SULFATE (UNII: 0YPR65R21J) (SODIUM SULFATE ANHYDROUS - UNII:36KCS0R750)	SODIUM SULFATE	12 [hp_C] in 12 [hp_C]

Inactive Ingredients

Ingredient Name	Strength
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	
SUCROSE (UNII: C151H8M554)	

Product Characteristics

Color	white	Score	
Shape	ROUND	Size	4mm

Flavor		Imprint Code		
Contains				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-3635-41	12 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
unapproved homeopathic		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