## SULPHURICUM ACIDUM- sulfuric acid 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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## Sulphuricum acidum 30C

Sulphuricum acidum 30C (\*\*contains 0.443 mg of the active ingredient per pellet)

Acid indigestion and cold sores\*

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



SULPHURICUM AC	IDUM					
sulfuric acid pellet						
Product Information						
Product Type	HUMAN OTC DRUG		Item Coo	de (Source)	NDC:0220-4	860
Route of Administration	ORAL					
Active Ingradient/Activ	o Meioty					
Active Ingredient/Activ	e molety					
Ingredient Name Basis of Strength						gth
SULFURIC ACID (UNII: O40UQP6WCF) (SULFURIC ACID - UNII:O40UQP6WCF)				SULFURIC ACID	30 [hp_C] in 30 [hp_C]	]
Inactive Ingredients						
Ingredient Name						ngth
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)						
SUCROSE (UNII: C151H8M554)						
<b>Product Characteristic</b>	S					
Color	white	Score	3			
Shape	ROUND	Size			4mm	

		-	mprint Cod	C				
Contains								
Packaging								
ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date			
NDC:0220-4860- 41	30 [hp_C] in Product			03/03/1983				
arkotina	Informa	tion						
Marketing Category	Applic	ation Number or Mo Citation	onograph	Marketing Start Date	Marketing End Date			
ipproved neopathic				03/03/1983				
	Item Code NDC:0220-4860- A1  arketing Marketing Category	Item Code Para Strain Para Strain Para Strain Para Strain Product Prod	Item Code     Package Descriptio       NDC:0220-4860- 41     30 [hp_C] in 1 TUBE; Type 0: Not a C Product       arketing Information       Marketing Category       Application Number or Marketion       Application Number or Marketion	Item Code       Package Description         NDC:0220-4860- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product         arketing Category       Application Number or Monograph Citation	Item Code       Package Description       Marketing Start Date         NDC:0220-4860- 41       30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product       03/03/1983         arketing Information         Marketing Category       Application Number or Monograph Citation       Marketing Start Date         og/03/1983       03/03/1983			

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment						
Name	Address	ID/FEI	<b>Business Operations</b>			
Boiron		282560473	manufacture(0220-4860)			

Revised: 11/2023

Boiron