CALCAREA SULPHURICA- calcium sulfate anhydrous 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.

Calcarea sulphurica 30C

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

Pustular Acne*

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).

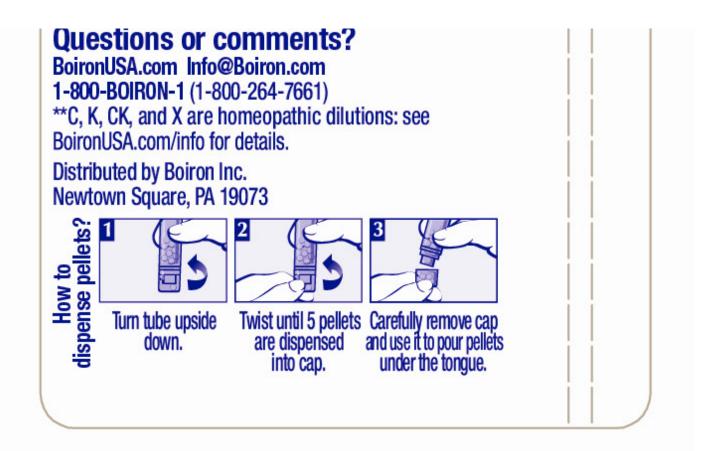
Uses: See symptoms on front panel.

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



CALCAREA SULPHURICA								
calcium sulfate anhydrous pellet								
Product Information								
Product Type	HUMAN OTC DRUG	Item Code (Source)		NDC:	NDC:0220-0999			
Route of Administration	ORAL							
Active Ingredient/Active Moiety								
Ingred	ient Name		Basis of Streng	th	Strength			
CALCIUM SULFATE ANHYDROUS - UNII:2M83C4R6ZB)	CIUM SULFATE ANHYDROUS (UNII: E934B3V59H) (CALCIUM CATION CALCIUM SULFATE ANHYDROUS (UNII: E934B3V59H) (CALCIUM CATION CALCIUM SULFATE ANHYDROUS				0 [hp_C] n 30 [hp_C]			
Inactive Ingredients								
		Strength						
LACTOSE, UNSPECIFIED FORM								
SUCROSE (UNII: C151H8M554)								
Product Characteristics								
Color	white	Score						
Shape F	ROUND	Size			4mm			
Flavor		Imprint Code						

ontains			
ackaging			
ltem Code	Package Description	Marketing Start Date	Marketing End Date
NDC:0220-0999- 41	30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	
arketing	Information		
Marketing	Application Number or Monograph	Marketing Start Date	Marketing End Date
Category	Citation	Date	Date
	ackaging Item Code NDC:0220-0999- 41	Ackaging Package Description Item Code Package Description NDC:0220-0999- 41 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product arketing Information	Ackaging Marketing Start Item Code Package Description Marketing Start NDC:0220-0999- 41 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-0999)				

Revised: 2/2023

Boiron