GLONOINUM- nitroglycerin 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.

Glonoinum 5C

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

Sudden hot flash and headache*

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).

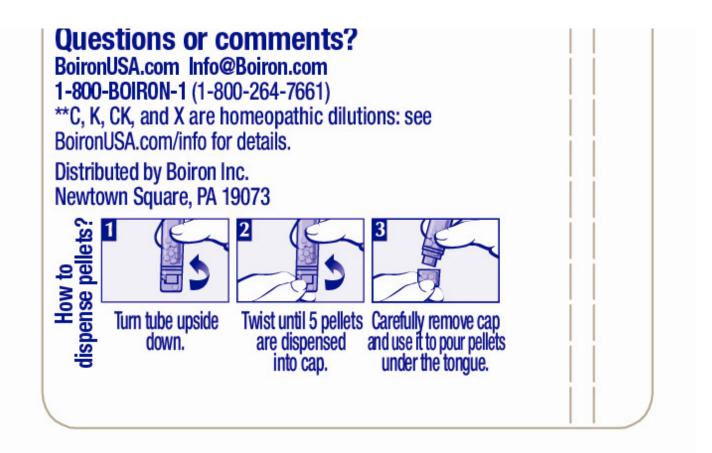
Uses: See symptoms on front panel.

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



GLONOINUM					
nitroglycerin pellet					
Product Information					
Product Type	HUMAN OTC DRUG	ltem	Code (So	urce)	NDC:0220-2290
Route of Administration	ORAL				
Active Ingredient/Active	e Moiety				
Ingre	dient Name			Basis of Strength	Strength
NITROGLYCERIN (UNII: G59M7S0 UNII:G59M7S0WS3))WS3) (NITROGLYCERII	N -	NITP	ROGLYCERIN	5 [hp_C] in 5 [hp_C]
Inactive Ingredients					
	Strength				
SUCROSE (UNII: C151H8M554)					
LACTOSE, UNSPECIFIED FORM	(UNII: J2B2A4N98G)				
Product Characteristics					
Color	white	Score			
Shape	ROUND	Size			4mm

Flavor		Imprint C	Imprint Code						
Contains									
Ρ	ackaging								
#	ltem Code	Ρ	ackage Description	Marketing Start Date	Marketing End Date				
1	NDC:0220-2290- 41	5 [hp_C] in 1 Product	TUBE; Type 0: Not a Combination	03/03/1983					
Marketing Information									
	Marketing Category	Application Number or Monograph Citation		h Marketing Start Date	Marketing End Date				

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment								
Name	Address	ID/FEI	Business Operations					
Boiron		282560473	manufacture(0220-2290)					

Revised: 11/2023

Boiron