## 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.

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## **Glonoinum 6X**

Glonoinum 6X

(\*\*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.

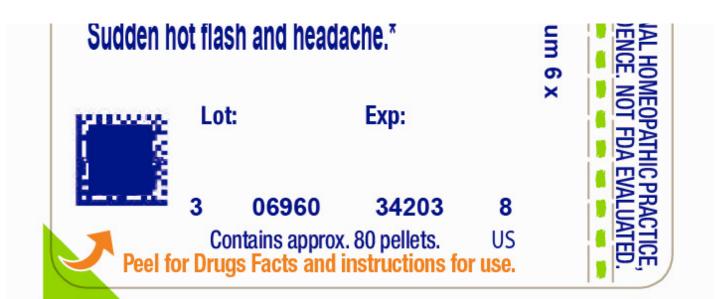
\*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

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**<sup>\*\*</sup>: See product name on front panel (contains 0.443 mg of the active ingredient per pellet).

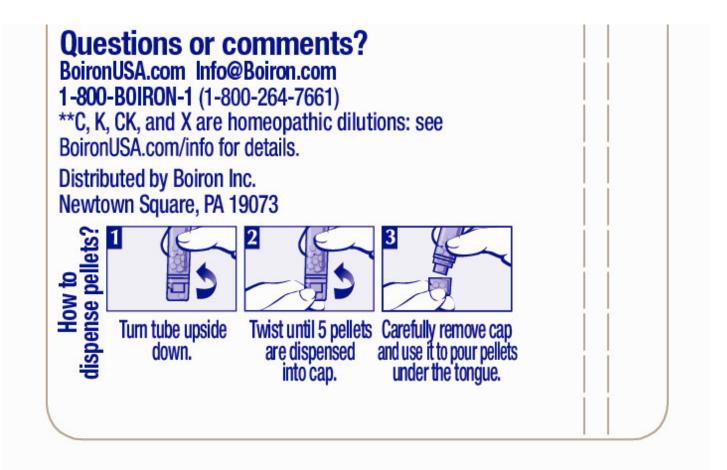
**Uses:** See symptoms on front panel.

**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	Item Code (Sou	urce)	NDC:022	0-2293			
Route of Administration	ORAL							
Active Ingredient/Active Moiety								
Ing	<b>Basis of Strength</b>		Strength					
NITROGLYCERIN (UNII: G59M7S0WS3) (NITROGLYCERIN - UNII:G59M7S0WS3)			NITROGLYCERIN		6 [hp_X]			
Inactive Ingredients								
lr		Strength						
SUCROSE (UNII: C151H8M554)								
LACTOSE (UNII: J2B2A4N98G)								
Product Characteristics								
Product Characteristics	white	Score						
Product Characteristics		Score Size Imprint Code		4m	m			

Сс	ontains						
Packaging							
#	ltem Code	Package Description	N	larketing Start Date	Marketing End Date		
1	NDC:0220-2293- 41	80 in 1 TUBE; Type 0: Not a Combination Product	03/03/1983				
N	larketing l	nformation					
	arketing Information						
	Marketing Category	Application Number or Monograph Citation		Marketing Start Date	Marketing End Date		
	approved meopathic			03/03/1983			

Labeler - Boiron (282560473)

Registrant - Boiron Inc. (014892269)

Establishment							
Name	Address	ID/FEI	<b>Business Operations</b>				
Boiron		282560473	manufacture(0220-2293)				

Revised: 11/2023

Boiron