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 200CK

Glonoinum 200CK

(**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

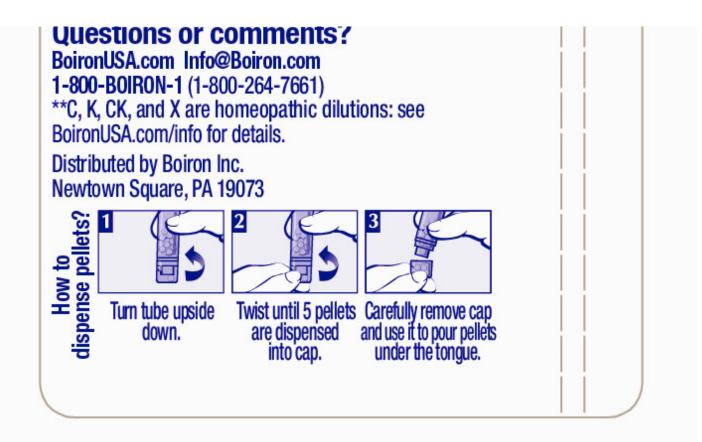




Drug Facts Active ingredient** See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before 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 Co	de (Source)	NDC:0220-2279
Route of Administration	ORAL				
Active Ingredient/Active	Moiety				
Ingred	ient Name		Basis of Strength	Strength	
NITROGLYCERIN (UNII: G59M7S0WS3) (NITROGLYCERIN - UNII:G59M7S0WS3)				NITROGLYCERIN	200 [kp_C] in 200 [kp_C]
Inactive Ingredients					
	Strength				
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)					
SUCROSE (UNII: C151H8M554)					
Product Characteristics					
Color	white	Score			
Shape	ROUND	Size			4mm

Flavor		Imprint Code					
Contains							
D	I 						
Packaging							
#	ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
1	NDC:0220- 2279-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product		a Combination	03/03/1983		
M	larkotina	Informa	tion				
Marketing Information							
	Marketing Category	Applic	ation Number or N Citation	Monograph	Marketing Start Date	Marketing End Date	
	approved meopathic				03/03/1983		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-2279)				

Revised: 10/2023

Boiron