PHOSPHORUS- phosphorus 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.

Phosphorus 12X

Phosphorus 12X

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

Dizziness With Sleeplessness*

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).

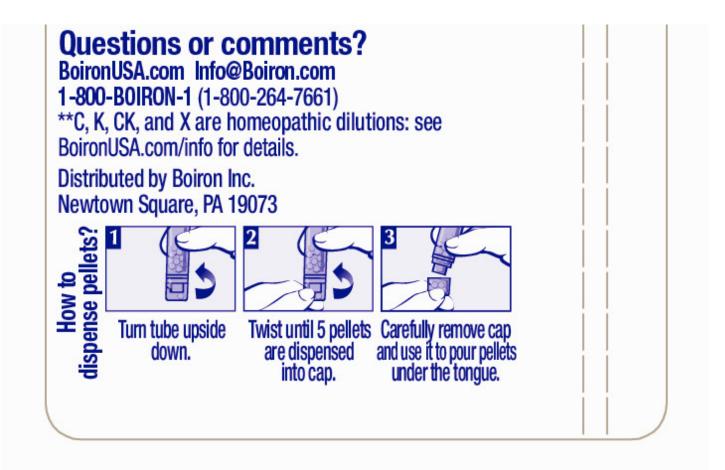
Uses: See symptoms on front panel.

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



PHOSPHORUS								
phosphorus pellet								
Product Information								
Product Type	HUMAN OTC DRUG		Item Code (Source)			NDC:0220-4004		
Route of Administration	ORAL							
Active Ingredient/Active	e Moiety							
Ingr	edient Name			Bas	is of Strength		Strength	
PHOSPHORUS (UNII: 27YLU75U4W) (PHOSPHORUS - UNII:27YLU75U4W) PHOS				PHOS	SPHORUS		12 [hp_X]	
Inactive Ingredients								
						Stren	Strength	
SUCROSE (UNII: C151H8M554)								
LACTOSE (UNII: J2B2A4N98G)								
Product Characteristics								
Color	white	Score						
Shape	ROUND	Size				4	mm	
Flavor		Impri	nt Code					

Contains								
Packaging								
#	ltem Code	Package Description	Marketing Start Date		Marketing End Date			
1	NDC:0220-4004- 41	80 in 1 TUBE; Type 0: Not a Combination Product	03/03/1983					
M	larketing I	nformation						
	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	Business Operations				
Boiron		282560473	manufacture(0220-4004)				

Revised: 7/2023

Boiron