ORNITHOGALUM UMBELLATUM- ornithogalum umbellatum 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.

Ornithogalum umbellatum 30C

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

Heartburn with pain while swallowing*

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).

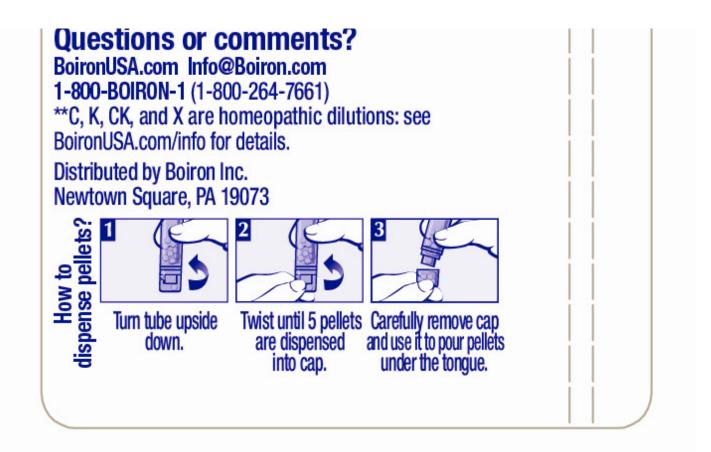
Uses: See symptoms on front panel.

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



ORNITHOGALUM UN	1BELLATUM						
ornithogalum umbellatum pellet							
Product Information							
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Product Type	HUMAN OTC DRUG	Item Code (Source)			NDC:0220-3820		
Route of Administration	ORAL						
Active Ingredient/Active	Moiety						
Basis of						<u>.</u>	
Ingredient Name Strength						Strength	
ORNITHOGALUM UMBELLATUM UMBELLATUM - UNII:9NS3M2Y78S)	TUM (UNII: 9NS3M2Y78S) (ORNITHOGALUM 78S) ORNITHOGALUM UMBELLATUM				30 [hp_C] in 30 [hp_C]		
Inactive Ingredients							
Ingredient Name						Strength	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)							
SUCROSE (UNII: C151H8M554)							
Product Characteristics							
Color	white	Score	e				
Shape	ROUND	Size				4mm	

Flavor		Imprint Code					
Contains							
Packaging							
#	Item Code	P	Package Description		Marketing Start Date	Marketing End Date	
	NDC:0220-3820- 41	30 [hp_C] in 3 Product	[hp_C] in 1 TUBE; Type 0: Not a Combination duct		03/03/1983		
М	arkotina	Informa	tion				
Marketing Information							
	Marketing Category	Applic	ication Number or Monograph Citation		Marketing Start Date	Marketing End Date	
	approved meopathic				03/03/1983		
hor	neopathic				02/02/1802		

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment							
Name	Address	ID/FEI	Business Operations				
Boiron		282560473	manufacture(0220-3820)				

Revised: 11/2023

Boiron