CANTHARIS- lytta vesicatoria 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.

Cantharis 15C

Cantharis 15C

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

Blisters with burning pain*

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).

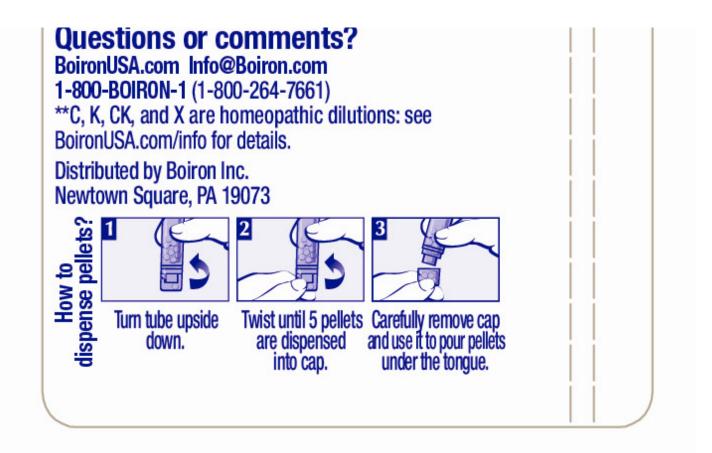
Uses: See symptoms on front panel.

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



CANTHARIS						
lytta vesicatoria pellet						
Product Information						
Product Type	HUMAN OTC DRUG	Item Code	(Source)	NDC:0220-1079		
Route of Administration	ORAL					
Active Ingredient/Active	Moiety					
Ingredient Name Basis of Strength				Strength		
LYTTA VESICATORIA (UNII: 3Q034RO3BT) (LYTTA VESICATORIA - UNII: 3Q034RO3BT) LYTTA VESICATORIA			15 [hp_C] in 15 [hp_C]			
Inactive Ingredients						
	Strength					
LACTOSE, UNSPECIFIED FORM						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						
Color	white	Score				
Shape	ROUND	Size		4mm		

ntains						
Packaging						
ltem Code	P	Package Description		Marketing Start Date	Marketing End Date	
IDC:0220-1079-	15 [hp_C] in 2 Product	1 TUBE; Type 0: Not a	Combination	03/03/1983		
rkoting	Informa	tion				
Marketing information						
Marketing Category	Applic	ation Number or M Citation	lonograph	Marketing Start Date	Marketing End Date	
pproved neopathic				03/03/1983		
	Item Code DC:0220-1079-1 arketing Marketing Category	Item Code DC:0220-1079- 1 To [hp_C] in The second	Item Code Package Description IDC:0220-1079-1 15 [hp_C] in 1 TUBE; Type 0: Not a Product arketing Application Number or M Category opproved 0	Item Code Package Description DC:0220-1079- 1 15 [hp_C] in 1 TUBE; Type 0: Not a Combination Product arketing Category Application Number or Monograph Citation	Item Code Package Description Marketing Start Date IDC:0220-1079- 15 [hp_C] in 1 TUBE; Type 0: Not a Combination Product 03/03/1983 arketing Information Marketing Category Application Number or Monograph Citation Marketing Start Date oproved 03/03/1983 03/03/1983 03/03/1983	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment					
Name	Address	ID/FEI	Business Operations		
Boiron		282560473	manufacture(0220-1079)		

Revised: 11/2023

Boiron