ALLIUM CEPA- onion 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.

Allium cepa 200CK

Allium cepa 200CK

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

Runny Nose With Clear Discharge*

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

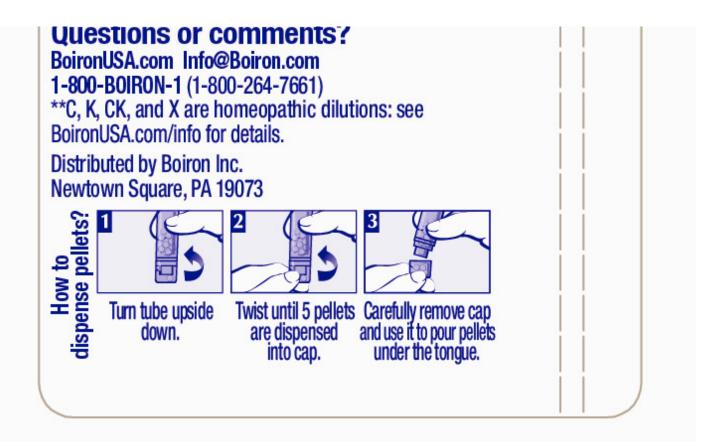




Drug Facts Active ingredient*: See product name on front panel (contains 0.443 mg of the active ingredient per pellet). Uses: See symptoms on front panel. Warnings: Stop use and ask a doctor if symptoms persist for more than 3 days or worsen. If pregnant or breastfeeding, ask a health professional before 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



ALLIUM CEPA					
onion pellet					
Product Information					
Product Type	HUMAN OTC DRUG		ltem Code (Source)		NDC:0220-0178
Route of Administration	ORAL				
Active Ingredient/Active	e Moietv				
Ingredient Name			Basis of Strength		Strength
		ONION	200 [<p_c] 200="" [kp_c]<="" in="" td=""></p_c]>	
Inactive Ingredients					
	Strength				
SUCROSE (UNII: C151H8M554)					
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)					
Product Characteristics	;				
Color	white	Score			
Shape	ROUND	Size			4mm
Flavor		Imprir	nt Code		

Co	ontains							
Packaging								
#	ltem Code	Package Description	Marketing Start Date	Marketing End Date				
1	NDC:0220- 0178-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combinat Product	on 03/03/1983					
Marketing Information								
	Marketing Category	Application Number or Monograp Citation	h 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(0220-0178)					

Revised: 10/2023

Boiron