AMMONIUM CARBONICUM- ammonium carbonate 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.

Ammonium carbonicum 200CK

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

Cold symptoms with stuffy nose and nosebleeds*

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.

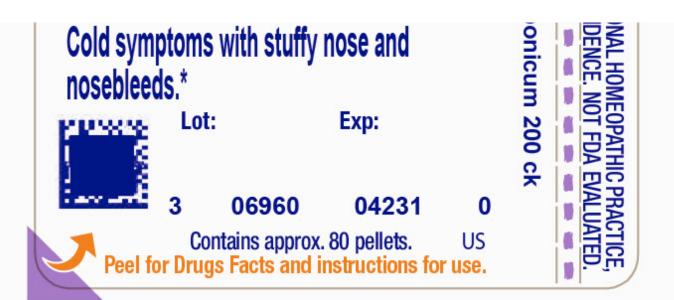
*C,K,CK, and X are homeopathic dilutions: see BoironUSA.com/info for details.

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



AMMONIUM CARBONICUM						
ammonium carbonate pellet						
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Product Information						
Product Type	HUMAN OTC DRUG	Item Cod	e (Source)	NDC:0220-0242		
Route of Administration	ORAL					
Active Ingredient/Active Moiety						
Ingred	Strength					
			200 [kp_C] in 200 [kp_C]			
Inactive Ingredients						
	Strength					
LACTOSE, UNSPECIFIED FORM						
SUCROSE (UNII: C151H8M554)						
Product Characteristics						
Color	white	Score				
Shape	ROUND	Size		4mm		

Flavor		Imprint Code					
Contains							
D	· • • • • • • • • •						
Packaging							
#	ltem Code	Р	Package Description		Marketing Start Date	Marketing End Date	
1	NDC:0220- 0242-41	200 [kp_C] in Product	1 TUBE; Type 0: Not a	a Combination	03/03/1983		
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(0220-0242)			

Revised: 1/2023

Boiron