AQUA MARINA- sodium chloride 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.

Aqua marina 200CK HPUS

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Active ingredient**: See product name on front panel (**contains 0.443 mg of the active ingredient per pellet).

Uses: See symptoms on front panel.

Relieves runny nose due to allergies, worse in morning *

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.

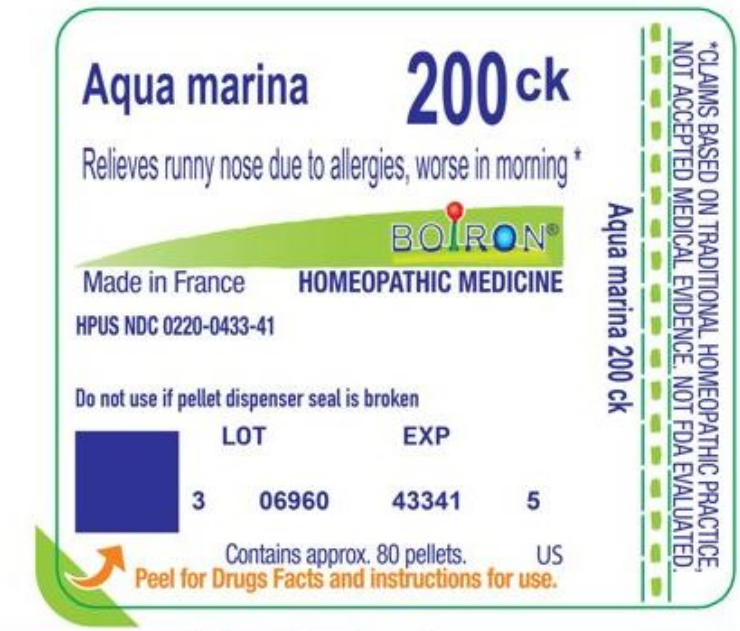
Keep out of reach of children.

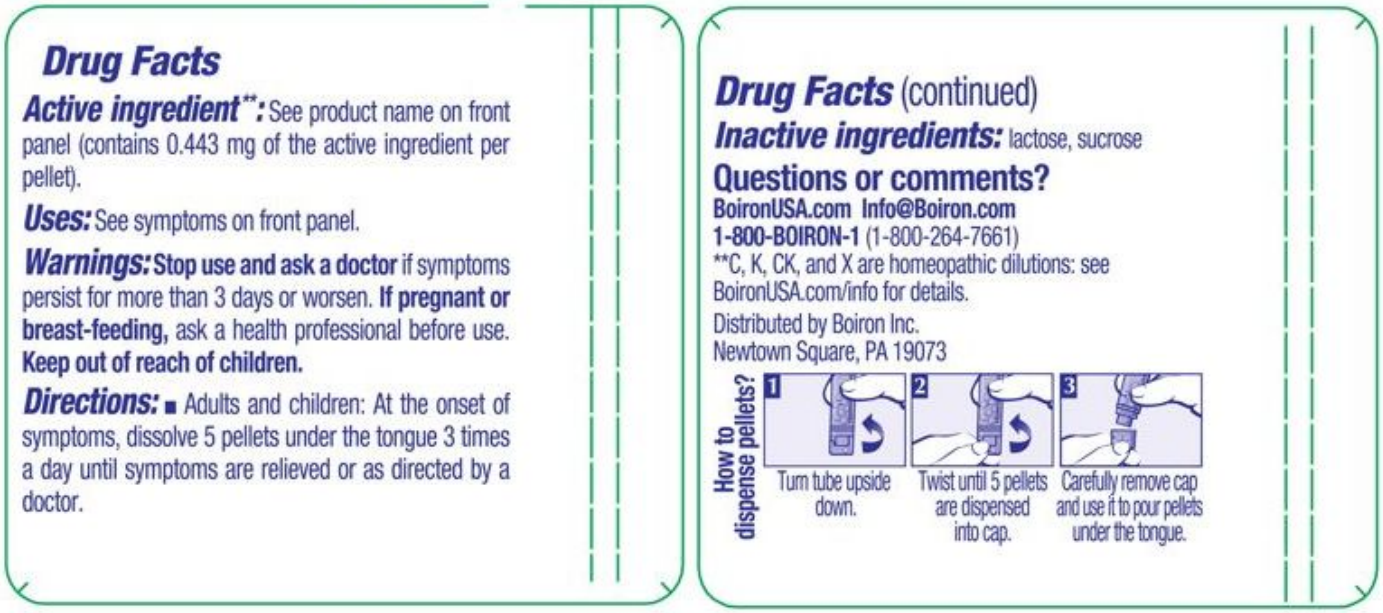
Adults and children: At the onset of symptoms, dissolve 5 pellets under the tongue 3 times a day until symptoms are relieved or as directed by a doctor.

lactose, sucrose

BoironUSA.com Info@boiron.com
1-800-BOIRON-1 (1-800-264-7661)
Distributed by Boiron, Inc.
Newtown Square, PA 19073

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.





AQUA MARINA

sodium chloride pellet

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0220-0433
Route of Administration	ORAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	200 [kp_C] in 200 [kp_C]

Inactive Ingredients

Ingredient Name	Strength
SUCROSE (UNII: C151H8M554)	
LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G)	

Product Characteristics

Color	white	**Score**	
Shape	ROUND	**Size**	4mm
Flavor		**Imprint Code**	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0220-0433-41	200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product	01/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		01/01/2024	

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment

Name	Address	ID/FEI	Business Operations
Boiron		282560473	manufacture(0220-0433)

Revised: 3/2024 Boiron