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 30C

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

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

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



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Lot: Exp:

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Contains approx. 80 pellets. US

Peel for Drugs Facts and instructions for use.

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



AQUA MARINA

sodium chloride pellet

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0220-0432
Route of Administration	ORAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SODIUM CHLORIDE (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698)	SODIUM CHLORIDE	30 [hp_C] in 30 [hp_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		

ı	Packaging			
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:0220-0432-	30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product	03/03/1983	

		Marketing Information			
Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
	03/03/1983				
•		Citation Date			

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

Establishment			
Name	Address	ID/FEI	Business Operations
Boiron		282560473	manufacture(0220-0432)

Revised: 10/2023 Boiron