

**NATRUM MURIATICUM- 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.*

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**Natrum muriaticum 30C**

Natrum muriaticum 30C

(\*\*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



Runny nose due to allergies, worse in morning.\*



Lot:

Exp:

3

06960

51113

7

Contains approx. 80 pellets.

US

Peel for Drugs Facts and instructions for use.

aticum 30 c

VAL HOMEOPATHIC PRACTICE,  
JENCE. NOT FDA EVALUATED.

## 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

# Questions or comments?

BoironUSA.com Info@Boiron.com

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How to dispense pellets?



Turn tube upside down.



Twist until 5 pellets are dispensed into cap.



Carefully remove cap and use it to pour pellets under the tongue.

## NATRUM MURIATICUM

sodium chloride pellet

### Product Information

|                         |                |                    |               |
|-------------------------|----------------|--------------------|---------------|
| Product Type            | HUMAN OTC DRUG | Item Code (Source) | NDC:0220-3599 |
| Route of Administration | ORAL           |                    |               |

### Active Ingredient/Active Moiety

| Ingredient Name  | Basis of Strength | Strength                  |
|--|-------------------|---------------------------|
| <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X) (CHLORIDE ION - UNII:Q32ZN48698) | SODIUM CHLORIDE   | 30 [hp_C]<br>in 30 [hp_C] |

### Inactive Ingredients

| Ingredient Name                                     | Strength |
|---|----------|
| <b>LACTOSE, UNSPECIFIED FORM</b> (UNII: J2B2A4N98G) |          |
| <b>SUCROSE</b> (UNII: C151H8M554)                   |          |

### Product Characteristics

|       |       |       |     |
|-------|-------|-------|-----|
| Color | white | Score |     |
| Shape | ROUND | Size  | 4mm |

| <b>Flavor</b>                |  | <b>Imprint Code</b>                                    |                      |                    |
|------------------------------|--|--|----------------------|--------------------|
| <b>Contains</b>              |  |  |                      |                    |
| <b>Packaging</b>             |  |  |                      |                    |
| #                            | Item Code                                | Package Description                                    | Marketing Start Date | Marketing End Date |
| 1                            | NDC:0220-3599-41                         | 30 [hp_C] in 1 TUBE; Type 0: Not a Combination Product | 03/03/1983           |                    |
| <b>Marketing Information</b> |  |  |                      |                    |
| Marketing Category           | Application Number or Monograph Citation | Marketing Start Date                                   | Marketing End Date   |                    |
| unapproved homeopathic       |  | 03/03/1983   |                      |                    |

**Labeler** - Boiron (282560473)

**Registrant** - Boiron, Inc. (014892269)

| <b>Establishment</b> |         |           |                        |
|----------------------|---------|-----------|------------------------|
| Name                 | Address | ID/FEI    | Business Operations    |
| Boiron               |         | 282560473 | manufacture(0220-3599) |

Revised: 1/2023

Boiron