AURUM MURIATICUM NATRONUM- sodium tetrachloroaurate 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.

Aurum muriaticum natronatum 200CK

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

Warts, hemorrhoids.*

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



atronatum 200 cl

Lot: Exp:

3 06960

09431

9

7

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).

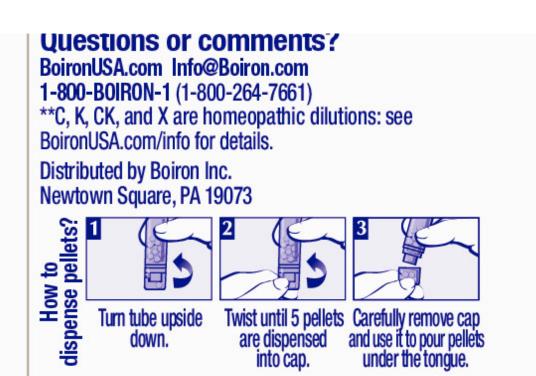
Uses: See symptoms on front panel.

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



AURUM MURIATICUM NATRONUM

sodium tetrachloroaurate pellet

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0220-0646 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | | |
|--|-----------------------------|-----------------------------|--|--|
| Ingredient Name | Basis of Strength | Strength | | |
| SODIUM TETRACHLOROAURATE (UNII: 7FT6QUT299) (TETRACHLOROAURATE ION - UNII:ZNL6IP5PJX) | SODIUM TETRACHLOROAURATE | 200 [kp_C] in 200 [kp_C] | | |

| Inactive Ingredients | | |
|--|----------|--|
| Ingredient Name | Strength | |
| LACTOSE, UNSPECIFIED FORM (UNII: J2B2A4N98G) | | |
| SUCROSE (UNII: C151H8M554) | | |

| Product Characteristics | | | |
|-------------------------|-------|--------------|-----|
| Color | white | Score | |
| Shape | ROUND | Size | 4mm |
| Flavor | | Imprint Code | |

Contains

| Packaging | | | | | |
|-----------|---|----------------------|---|------------|-----------------------|
| | # | Item Code | Item Code Package Description | | Marketing End Date |
| | 1 | NDC:0220- 0646-41 | 200 [kp_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 | | |
| | Application Number or Monograph | Application Number or Monograph Marketing Start Citation Date | |

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

| Establishment | | | |
|---------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| Boiron | | 282560473 | manufacture(0220-0646) |

Revised: 11/2023 Boiron