RUMEX CRISPUS- rumex crispus root 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.

Rumex crispus 9C

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

Dry cough worsened by breathing cold air*

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.

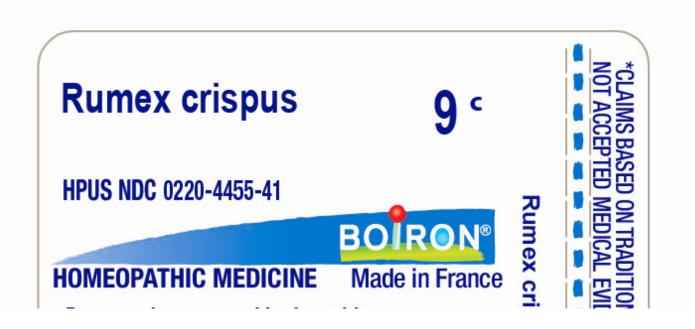
*CLAIMS BASED ON TRADITIONAL HOMEOPATHIC PRACTICE NOT ACCEPTED MEDICAL EVIDENCE. NOT FDA EVALUATED.

*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



pus 9 c

VAL HOMEOPATHIC PR DENCE, NOT FDA EVAL

Lot: Exp:

3 06960 64110 0

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).

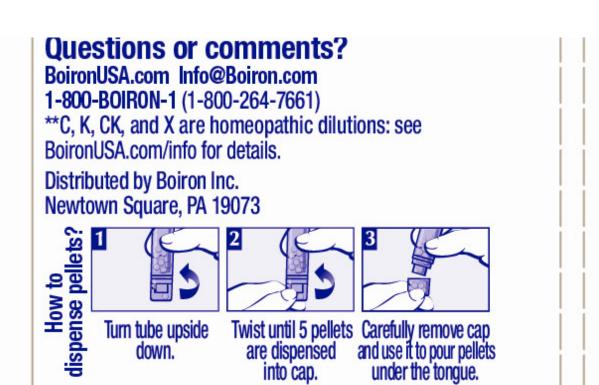
Uses: See symptoms on front panel.

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



RUMEX CRISPUS

rumex crispus root pellet

| Product Information | | | | |
|-------------------------|----------------|--------------------|---------------|--|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0220-4455 | |
| Route of Administration | ORAL | | | |

| Active Ingredient/Active Moiety | | | |
|--|-----------------------|-------------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| RUMEX CRISPUS ROOT (UNII: 9N1RM2S62C) (RUMEX CRISPUS ROOT - UNII:9N1RM2S62C) | RUMEX CRISPUS ROOT | 9 [hp_C] in 9 [hp_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 | Package Description | Marketing Start Date | Marketing End Date |
| | | NDC:0220-4455- 41 | 9 [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 | | |
| | 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(0220-4455) | |

Revised: 11/2023 Boiron