HYDRASTIS CANADENSIS- goldenseal 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.

Hydrastis canadensis 200CK

Hydrastis canadensis 200CK

Less than 10 -12 mg hydrastine, berberine alkaloids per pellet

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

Post-Nasal Drip*

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



HOMEOPAIHIC MEDICINE Made in France

HPUS NDC 0220-2556-41

Less than 10⁻¹² mg hydrastine, berberine alkaloids per pellet Do not use if pellet dispenser seal is broken



LOT: EXP:

06960

372318

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 lactore curren

Questions or comments?

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dispense pellets?







Turn tube upside down.

into cap.

Twist until 5 pellets Carefully remove cap are dispensed and use it to pour pellets under the tongue.

Hydrastis canadensis 200 ck

HPUS NDC 0220-2556-41



HOMEOPATHIC MEDICINE

Made in France

Post-nasal drip.*



Exp:

Hydrastis canadensis 200 ck





US

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.

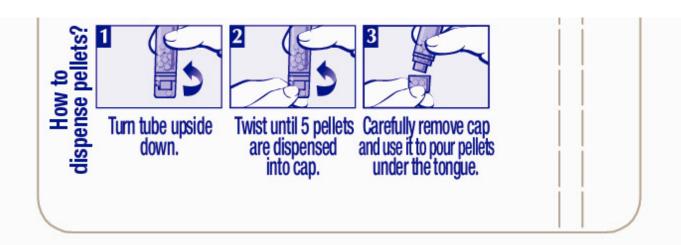
Drug Facts (continued) Inactive ingredients: lactose, sucrose

Questions or comments?

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HYDRASTIS CANADENSIS

goldenseal pellet

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:0220-2556

Route of Administration ORAL

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|-----------------|----------------------|----------|
| | Strength | _ |

GOLDENSEAL (UNII: ZW3Z11D0JV) (GOLDENSEAL - UNII:ZW3Z11D0JV) GOLDENSEAL - GOLDENSEAL 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

| П | - actualing | | | | |
|---|-------------|----------------------|---|-------------------------|-----------------------|
| | # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| | 1 | NDC:0220- 2556-41 | 200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product | 03/03/1983 | |

| Marketing Information | | | | |
|---------------------------|---|-------------------------|-----------------------|--|
| 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)

| Establishment | | | |
|---------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| Boiron | | 282560473 | manufacture(0220-2556) |

Revised: 12/2023 Boiron