AMBROSIA ARTEMISIAEFOLIA- ambrosia artemisiifolia 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.

AMBROSIA ARTEMISIAEFOLIA 200CK MD

Ambrosia artemisiaefolia 200CK

Active ingredient**: See product name on front panel (**contains 0.443 mg of the active ingredient per pellet).

Relieves symptoms of hay fever *

See symptoms on front panel.

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.

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.

lactose, sucrose

BoironUSA.com Info@boiron.com 1-800-BOIRON-1 (1-800-264-7661)

Distributed by Boiron, Inc.

Newtown Square, PA 19073

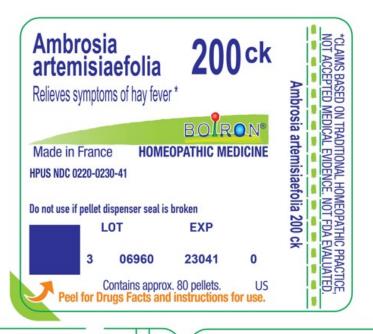
Do not use if pellet dispenser seal is broken.

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.



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 breast-feeding, 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.



AMBROSIA ARTEMISIAEFOLIA

ambrosia artemisiifolia pellet

| Product Information | | | |
|-------------------------|----------------|--------------------|---------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:0220-0230 |
| Route of Administration | ORAL | | |

| Active Ingredient/Active Moiety | | | |
|--|----------------------------|-----------------------------|--|
| Ingredient Name | Basis of Strength | Strength | |
| AMBROSIA ARTEMISIIFOLIA (UNII: 9W34L2CQ9A) (AMBROSIA ARTEMISIIFOLIA - UNII:9W34L2CQ9A) | AMBROSIA ARTEMISIIFOLIA | 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 | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:0220- 0230-41 | 200 [kp_C] in 1 TUBE; Type 0: Not a Combination Product | 01/01/2024 | |

| Marketing Information | | | |
|---------------------------|---|-------------------------|-----------------------|
| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| unapproved homeopathic | | 01/01/2024 | |
| homeopathic | | 01/01/2024 | |

Labeler - Boiron (282560473)

Registrant - Boiron, Inc. (014892269)

| Establishment | | | |
|---------------|---------|-----------|------------------------|
| Name | Address | ID/FEI | Business Operations |
| Boiron | | 282560473 | manufacture(0220-0230) |

Revised: 12/2023 Boiron