

POLLEN GUARD RAGWEED RAGWEED HAY FEVER SYMPTOM RELIEF - ambrosia artemisiaefolia liquid

Western Allergy Services Ltd.

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.

Short Ragweed (Ambrosia Artemisiaefolia) 1X

Hay Fever Symptom Relief

temporarily relieves symptoms of hay fever resulting from exposure to ragweed such as:

- runny nose
- nasal congestion/pressure
- sneezing
- itching
- water, itchy eyes
- hives

Do not use

- if you have asthma, severe immunodeficiencies, malignancies, or autoimmune disease
- if you take beta blockers

When using this product

- avoid other drugs and medications
- avoid food, drink, or teeth brushing for at least 5 minutes after each dose

Stop use and ask a doctor if

- condition persists or worsens
- mouth, throat, chest or abdominal discomfort occurs
- you experience hives, itching, or shortness of breath

If pregnant or breast feeding, ask a health professional before use.

Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.

Directions

- drop of extract may be placed under tongue, retained under the tongue for 2 minutes, then swallowed
- adults: take 1 drop (0.05 mL) 2 or 3 times per day before meals
- children over 3 years of age: only use treatment regimen if child has experienced more than 6 months of symptoms and is documented for allergies to ragweed
- children under 3 years of age: ask a doctor

- Store between 2-8 degrees C
- Do not use if safety seal is broken

Glycerin, Water

Questions or Comments?

<p>Drug Facts (continued)</p> <p>Stop use and ask a doctor if •condition persists or worsens •mouth, throat, chest or abdominal discomfort occurs •you experience hives, itching, or shortness of breath</p> <p>If pregnant or breast feeding, ask a health professional before use. Keep out of reach of children. In case of overdose, get medical help or contact a Poison Control Center right away.</p> <p>Directions •drop of extract may be placed under the tongue, retained under the tongue for 2 minutes, then swallowed •adults: take 1 drop (0.05 mL) 2 or 3 times per day before meals •children over 3 years of age: only use treatment regimen if child</p>	<p>Drug Facts (continued)</p> <p>has experienced more than 6 months of symptoms and is documented for allergies to ragweed •children under 3 years of age: ask a doctor</p> <p>Other Information •Store between 2-8° C •Do not use if safety seal is broken.</p> <p>Inactive Ingredients Glycerin, Water</p> <p>Questions or Comments? 1-866-335-5294 Monday through Friday 9am-5pm</p> <p>Western Allergy Services Mississauga, ON L4Z 2R6 Product of Canada www.pollenguard.com (WV)</p>		<p>Drug Facts</p> <p>Active Ingredients Short Ragweed (<i>Ambrosia Artemisiaefolia</i>) 1x...Hay Fever Symptom Relief</p> <p>Purpose Uses temporarily relieves symptoms of hay fever resulting from exposure to ragweed such as: •runny nose •nasal congestion/pressure •sneezing •itching •watery, itchy eyes •hives</p> <p>Warnings Do not use •if you have asthma, severe immunodeficiencies, malignancies, or autoimmune disease •if you take beta blockers When using this product •avoid other drugs and medications •avoid food, drink, or teeth brushing for at least 5 minutes after each dose</p>
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ambrosia artemisiaefolia liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76097-003
Route of Administration	ORAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
AMBROSIA ARTEMISIIFOLIA (UNII: 9W34L2CQ9A) (AMBROSIA ARTEMISIIFOLIA - UNII:9W34L2CQ9A)	AMBROSIA ARTEMISIIFOLIA	1 [hp_X] in 5 mL

Inactive Ingredients

Ingredient Name	Strength
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76097-003-05	5 mL in 1 BOTTLE, UNIT-DOSE		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
unapproved homeopathic		10/03/2011	

Labeler - Western Allergy Services Ltd. (208003467)

Registrant - Western Allergy Services Ltd. (208003467)

Establishment

Name	Address	ID/FEI	Business Operations
Quantum Allergy Canada, Inc.		246829324	manufacture

Revised: 9/2011

Western Allergy Services Ltd.