

ARTEMESIA ABSINTHIUM 5 SPECIAL ORDER- artemesia absinthium 5 special order liquid Uriel Pharmacy Inc.

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.

Artemesia absinthium 5 Special Order

Directions: FOR ORAL USE ONLY.

Take 3-4 times daily. Ages 12 and older: 10 drops. Ages 2-11: 5 drops. Under age 2: Consult a doctor.

Active Ingredient: 100 gm contains: 5g Absinthium ex herba 1X

Inactive Ingredients: Distilled water, 20% Organic grain alcohol

Use: Temporary relief of headache.

KEEP OUT OF REACH OF CHILDREN.

Warnings: Claims based on traditional homeopathic practice, not accepted medical evidence. Not FDA evaluated. Do not use if allergic to any ingredient. Consult a doctor before use for serious conditions or if conditions worsen or persist. If pregnant or nursing, consult a doctor before use. Do not use if safety seal is broken or missing.

Questions? Call 866.642.2858 Made with care by Uriel, East Troy, WI 53120 www.urielpharmacy.com

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**Artemesia
absinthium 5%
Special Order**
**Homeopathic Liquid
net vol. 2 fl. oz (60ml)**

ARTEMESIA ABSINTHIUM 5 SPECIAL ORDER

artemesia absinthium 5 special order liquid

Product Information

| | | | |
|-------------------------|----------------|--------------------|----------------|
| Product Type | HUMAN OTC DRUG | Item Code (Source) | NDC:48951-1260 |
| Route of Administration | ORAL | | |

Active Ingredient/Active Moiety

| Ingredient Name | Basis of Strength | Strength |
|--|-------------------|------------------|
| WORMWOOD (UNII: F84709P2XV) (WORMWOOD - UNII:F84709P2XV) | WORMWOOD | 1 [hp_X] in 1 mL |

| Inactive Ingredients | | | | |
|----------------------------|------------------|---|----------------------|--------------------|
| Ingredient Name | | | Strength | |
| WATER (UNII: 059QF0KO0R) | | | | |
| ALCOHOL (UNII: 3K9958V90M) | | | | |
| | | | | |
| Packaging | | | | |
| # | Item Code | Package Description | Marketing Start Date | Marketing End Date |
| 1 | NDC:48951-1260-3 | 60 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product | 09/01/2009 | |
| | | | | |
| Marketing Information | | | | |
| Marketing Category | | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
| unapproved homeopathic | | | 09/01/2009 | |
| | | | | |

Labeler - Uriel Pharmacy Inc. (043471163)

| Establishment | | | |
|---------------------|---------|-----------|-------------------------|
| Name | Address | ID/FEI | Business Operations |
| Uriel Pharmacy Inc. | | 043471163 | manufacture(48951-1260) |

Revised: 4/2018

Uriel Pharmacy Inc.