

**BIRCH JUNIPER- birch juniper 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.*

**Birch Juniper**

Directions: FOR ORAL USE ONLY.

Take 2 times daily. Ages 12 and older: 1-2 teaspoons. Ages 2-11: 1/2 teaspoon. Under age 2: Consult a doctor.

Active Ingredients: 1gm contains: 400mg Betula (Birch Leaf) 1X, 300mg Juniperus (Juniper berry) 1X

Inactive Ingredients: Distilled water, Sucanat, Honey, Citric acid

"prepared using rhythmical processes"

Use: Supports normal kidney function.

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.

SHAKE WELL BEFORE USE. REFRIGERATE AFTER OPENING. BEST WHEN USED WITHIN 90 DAYS OF OPENING.

Questions? Call 866.642.2858

Made with care by Uriel, East Troy, WI 53120

shopuriel.com Lot:



**BIRCH JUNIPER**

birch juniper liquid

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:48951-2149
<b>Route of Administration</b>	ORAL		

**Active Ingredient/Active Moiety**

<b>Ingredient Name</b>	<b>Basis of Strength</b>	<b>Strength</b>
<b>JUNIPER BERRY</b> (UNII: O84B5194RL) (JUNIPER BERRY - UNII:O84B5194RL)	JUNIPER BERRY	1 [hp_X] in 1 mL
<b>BETULA PENDULA BARK</b> (UNII: 40S83Y133C) (BETULA PENDULA BARK - UNII:40S83Y133C)	BETULA PENDULA BARK	1 [hp_X] in 1 mL

**Inactive Ingredients**

<b>Ingredient Name</b>	<b>Strength</b>
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>HONEY</b> (UNII: Y9H1V576FH)	
<b>SUCROSE</b> (UNII: C151H8M554)	
<b>CITRIC ACID MONOHYDRATE</b> (UNII: 2968PHW8QP)	

**Packaging**

<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:48951-2149-9	240 mL in 1 BOTTLE, GLASS; Type 0: Not a Combination Product	11/03/2021	

**Marketing Information**

<b>Marketing Category</b>	<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
unapproved homeopathic		11/03/2021	

**Labeler** - Uriel Pharmacy Inc (043471163)**Establishment**

<b>Name</b>	<b>Address</b>	<b>ID/FEI</b>	<b>Business Operations</b>
Uriel Pharmacy Inc		043471163	manufacture(48951-2149)