

**GNP WITCH HAZEL- witch hazel liquid**  
**Amerisource Bergen**

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**GNP Witch Hazel**

Drug Facts

**Active Ingredient**

Witch Hazel

**Purpose**

Astringent

**Indications**

For relief of minor skin Irritations due to  
minor cuts  
minor scraps  
insect bites

**Warnings**

For external use only  
avoid contact with eyes  
If contact occurs rinse thoroughly with water.

**When using this product stop using and contact a doctor if**

condition persists or gets worse  
symptoms do not improve within 7 days

**Keep out of reach of children.**

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

In case of eye contact flush eyes with running water for 15 minutes.

**Directions**

Apply liberally to the afflicted area as often as necessary

## Inactive ingredients

Alcohol 14% and purified water.

## Principal Display Panel

**NDC 46122-335-43**

**Witch Hazel**

**(Hamamelis Water)**

**Distilled Extract**

**Alcohol 14% by volume**

**16 fl oz (1 pt) 473 mL**

<p><b>Drug Facts</b></p> <table border="1"> <tr> <td><b>Active ingredient</b></td> <td><b>Purpose</b></td> </tr> <tr> <td>Witch Hazel Extract</td> <td>Astringent</td> </tr> </table> <p><b>Indications</b> For relief of minor skin irritations due to</p> <ul style="list-style-type: none"> <li>■ minor cuts</li> <li>■ minor scrapes</li> <li>■ insect bites</li> </ul> <p><b>Warnings</b> For external use only</p> <ul style="list-style-type: none"> <li>■ avoid contact with eyes. If contact occurs rinse thoroughly with water.</li> </ul> <p>When using this product stop use and contact a doctor if</p> <ul style="list-style-type: none"> <li>■ condition persists or gets worse</li> <li>■ symptoms do not improve within 7 days</li> </ul> <p>Keep out of reach of children. In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately. In case of eye contact, flush eyes with running water for 15 minutes.</p> <p><b>Directions</b> Apply liberally to the affected area as often as necessary</p> <p><b>Inactive ingredients</b></p> <ul style="list-style-type: none"> <li>■ alcohol 14%, and purified water</li> </ul>	<b>Active ingredient</b>	<b>Purpose</b>	Witch Hazel Extract	Astringent	 <p>NDC 46122-335-43 <b>*Safety Sealed</b></p>	<p><b>*Warning:</b> Do not use if tamper evident seal is broken or missing. This product is sealed with a breakaway cap ring.</p>
<b>Active ingredient</b>	<b>Purpose</b>					
Witch Hazel Extract	Astringent					
 <p>Distributed by: AmerisourceBergen 1300 Morris Drive Chesterbrook, PA 19087 Questions or concerns? Visit us at <a href="http://www.mygnp.com">www.mygnp.com</a> Made in U.S.A.</p>	<div style="background-color: #4a4a8a; color: white; padding: 10px; border-radius: 10px;"> <h1 style="margin: 0;">Witch Hazel</h1> <p style="margin: 0;">(Hamamelis Water)</p> <p style="margin: 5px 0 0 20px;"><b>Distilled Extract</b></p> <p style="margin: 5px 0 0 20px;">Alcohol 14% by volume</p> <ul style="list-style-type: none"> <li>• Astringent</li> <li>• Soothes minor skin irritations</li> </ul> <p style="margin: 10px 0 0 20px; font-size: 1.2em;"><b>16 fl oz (1 pt) 473 mL</b></p> </div>	<p>R071015DL      ABC# 470-351 10158335</p>  <p>0 87701 42733 6</p>				

## GNP WITCH HAZEL

witch hazel liquid

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:46122-335
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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<b>WITCH HAZEL</b> (UNII: 101I4J0U34) (WTCH HAZEL - UNII:101I4J0U34)		WTCH HAZEL	855 mg in 1 mL	
<b>Inactive Ingredients</b>				
<b>Ingredient Name</b>			<b>Strength</b>	
ALCOHOL (UNII: 3K9958V90M)				
WATER (UNII: 059QF0KO0R)				
<b>Packaging</b>				
<b>#</b>	<b>Item Code</b>	<b>Package Description</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
1	NDC:46122-335-43	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	
<b>Marketing Information</b>				
<b>Marketing Category</b>		<b>Application Number or Monograph Citation</b>	<b>Marketing Start Date</b>	<b>Marketing End Date</b>
OTC Monograph Drug		M016	01/01/2016	

**Labeler** - Amerisource Bergen (007914906)

**Registrant** - Pharma Nobis, LLC (118564114)

### Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(46122-335) , manufacture(46122-335) , pack(46122-335) , label(46122-335)

Revised: 12/2023

Amerisource Bergen