

WITCH HAZEL- witch hazel liquid
Freds Inc

Drug Facts

Active ingredient

Witch Hazel 86%

Purpose

Astringent

Uses

- for relief of minor skin irritations due to :
 - insect bites
 - minor cuts
 - minor scrapes

Warnings

For external use only.

- avoid swallowing

When using this product

- avoid contact with the eyes, If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if

condition worsens or symptoms persist for more than 7 days.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

apply to the affected area as often as necessary.

Other information

- keep tightly closed

- store at room temperature

Inactive ingredient

alcohol 14% by volume

Principal Display Panel

Witch Hazel

Alcohol 14% by Volume

ASTRINGENT/HAMAMELIS WATER

For Relief of minor skin irritations due to:

- Insect bites
- Minor cuts
- Minor scrapes

DISTRIBUTED BY: fred's, Inc.

4300 NEW GETWELL RD

MEMPHIS, TN 38118

www.fredsinc.com

Questions or comments:

1-855-331-FRED (3733)

FL OZ (mL)

**TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS
BROKEN OR MISSING**

Package Label

fred's
Pharmacy

NDC 55315-390-16

Witch Hazel

Alcohol 14% by Volume

ASTRINGENT/HAMAMELIS WATER

For relief of minor skin irritations due to:

- ✓ Insect bites
- ✓ Minor cuts
- ✓ Minor scrapes

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4300 NEW GETWELL RD
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100% satisfaction guaranteed

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16 fl oz (1 PT) 473 mL

SKU583294 PLD-C330D LB005493

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PLD-B330C LB002059

FRED'S PHARMACY Witch Hazel

WITCH HAZEL			
witch hazel liquid			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:55315-390
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
	Ingredient Name	Basis of Strength	Strength
	WITCH HAZEL (UNII: 101I4J0U34) (WTCH HAZEL - UNII:101I4J0U34)	WTCH HAZEL	842 mg in 1 mL
Inactive Ingredients			
	Ingredient Name	Strength	
	ALCOHOL (UNII: 3K9958V90M)		

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:55315-390-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/31/2014	06/30/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	12/31/2014	06/30/2024

Labeler - Freds Inc (005866116)

Revised: 11/2023

Freds Inc