WITCH HAZEL- witch hazel liquid
Pharmacy Vlaue Alliance LLC

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Witch Hazel
822

Active ingredient

Witch hazel 86%

Purpose

Astringent

use

for relief of minor skin irritations due to:

- insects bites
- minor cuts
- minor scrapes

Warnings

For external use only

When using this product

- avoid contact with the eyes

Stop use and ask a doctor if

- condition worsens or symptoms persist for more than 7 days

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

apply as often as needed

inactive ingredients

alcohol 14% by volume

Questions

Call 1-888-593-0593

*This product is not manufactued or distributed by Dickinson Brand, Inc., distributorof T.N. Dickinson

Witch Hazel.

If for any reason you are not satisfied with this product, please return it to the store where purchased for a full refund.

Distributed by: Phamacy Value Alliance LLC

407 East Lancaster Avenue, Wayne, PA 19087

www.emersongroup.com

Principal Display Panel

COMPARE TO T.N.DICKINSON'S*

Premier Value Witch Hazel u.s.p.

astringent

hamamelis water

for relief of minor skin irritations due to:

- insects bites
- minor cuts
- minor scrapes

Square bottle uses less plastic than a similarly sized round bottle

Recyclable (if available in your area)

16 FL OZ (1 PT) 473 mL

WITCH HAZEL

witch hazel liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:68016-822
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH 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:68016-822-43	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/15/2003	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	08/15/2003	

Labeler - Pharmacy Vlaue Alliance LLC (101668460)

Registrant - Vi-Jon (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		790752542	manufacture(68016-822)

Revised: 5/2020 Pharmacy Vlaue Alliance LLC