# WITCH HAZEL- witch hazel liquid Geiss, Destin & Dunn

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.

-----

#### GoodSense Witch Hazel

## Active ingredient

Witch hazel 86%

## **Purpose**

Astringent

#### Use

for relief of minor skin irritations due to:

- insect 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** ingredient

alcohol 14% by volume

## adverse reactions

This product is not manufactured or distributed by Dickinson Brands, Inc., distributor of T.N Dickinson Witch Hazel\*

Distributed By: Geiss, Destin & Dun, Inc

Peachtree City, GA 30269 www.valuelabels.com 822.001/822AA

## principal display panel

**GOOD SENSE** 

Witch Hazel USP

Hammamelis Water

Astringent

For relief of minor skin irritations due to:

- Insect bites
- Minor cuts
- Minor scrapes

Compare to active ingredients of T.N. Dickinson's Witch Hazel\* 100% SATISFACTION GUARANTEED

Square bottles ues 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

<b>.</b> .		T (		. •	
Drod	uct	Into	rm:	atinn	

Product Type HUMAN OTC DRUG Item Code (Source) NDC:50804-822

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 10 114J0 U34) (WITCH HAZEL - UNII:10 114J0 U34)	WITCH HAZEL	979 mg in 1 mL

Inactive I	ng re die nts
------------	---------------

Ingredient Name	Strength
-----------------	----------

ALCOHOL (UNII: 3K9958V90M)

Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date
	1 NDC:50804-822-	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	0 4/22/20 11	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	04/22/2011	

## Labeler - Geiss, Destin & Dunn (076059836)

## **Registrant -** Vi-Jon (790752542)

Establishment				
Name	Address	ID/FEI	Business Operations	
Vi-Jon		790752542	manufacture(50804-822)	

Revised: 11/2019 Geiss, Destin & Dunn