WITCH HAZEL- witch hazel liquid
H-E-B

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
822AA/822.001

Active ingredient

Witch Hazel 86%

Purpose

Astringent

Use

REMOVE STICKER ON PUMP PRIOR TO USE. (FOR PUMP BOTTLES)

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

For Pump

remove sticker prior to use. Pump onto a clean pad and apply as often as needed.

Inactive ingredient

alcohol 14% by volume

adverse reaction

MADE WITH PRIDE & CARE FOR

H-E-B, SAN ANTONIO, TX 78204

principal display panel

PUSH PUMP

H-E-B

Witch Hazel U.S.P.

Astringent

- Cleans & Refreshes
- For Relief of Minor Skin Irritations due to Insects Bites, Minor Cuts and Minor Scrapes

8 FL OZ (236 mL)

WITCH HAZEL

witch hazel liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:37808-822
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	979 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:37808-822-43	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2015	
2	NDC:37808-822-45	946 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/18/2015	
3	NDC:37808-822-99	236 mL in 1 BOTTLE, PUMP; Type 1: Convenience Kit of Co-Package	06/18/2015	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part346	06/18/2015	

Labeler - H-E-B (007924756)

Registrant - Vi-Jon, LLC (790752542)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		790752542	manufacture(37808-822)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon, LLC		088520668	manufacture(37808-822)

Revised: 9/2022 H-E-B