

WITCH HAZEL- witch hazel liquid
Hydrox Laboratories

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

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.

KEEP OUT OF REACH OF CHILDREN

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

Directions

Apply to the affected area as often as necessary. 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.

Inactive ingredient

Alcohol 14% by volume

DIST. BY:

HYDROX LABORATORIES

825-B Tollgate Rd. • Elgin, IL 60123

PRINCIPAL DISPLAY PANEL - 473 mL Bottle Label

Personal Care

FreshMoment

Witch Hazel

Natural Astringent

Cleans and Refreshes Skin

No Artificial Fragrance

or Dyes

MADE IN USA

16 FL. OZ. (473mL)

Personal Care



Witch Hazel

*Natural Astringent
Cleans and Refreshes Skin
No Artificial Fragrance
or Dyes*

MADE IN USA

16 FL. OZ. (473mL)

NDC 10565-040-16 REF D0082

Fresh Moment Witch Hazel is a natural formulation that gently cleans and refreshes your skin without stripping essential moisture.

Witch Hazel can also be used to soothe minor cuts, scrapes and insect bites.

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DIST. BY:
HYDROX LABS/ANTHONY'S
825-S Toigate Rd. • Bpph, IL 60123



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WITCH HAZEL				
witch hazel liquid				
Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:10565-040	
Route of Administration	TOPICAL			
Active Ingredient/Active Moiety				
Ingredient Name		Basis of Strength	Strength	
Witch Hazel (UNII: 101I4J0U34) (Witch Hazel - UNII:101I4J0U34)		Witch Hazel	86 g in 100 mL	
Inactive Ingredients				
Ingredient Name			Strength	
Alcohol (UNII: 3K9958V90M)				
Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date

1	NDC:10565-040-16	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	11/13/20 17	
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Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC MONOGRAPH FINAL	part347	12/21/20 11	

Labeler - Hydrox Laboratories (025164302)

Registrant - Hydrox Laboratories (025164302)

Establishment

Name	Address	ID/FEI	Business Operations
Hydrox Laboratories		025164302	MANUFACTURE(10565-040) , label(10565-040) , pack(10565-040)

Revised: 3/2020

Hydrox Laboratories