WITCH HAZEL- witch hazel liquid Medical Products Laboratories, Inc.

Witch Hazel

Warnings

For external use only

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 last for more than 7 days

Keep out of reach of children.

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

MANUFACTURED BY:

MEDICAL PRODUCTS LABORATORIES, INC.

9990 GLOBAL ROAD, PHILADELPHIA 19115

mplusa.com

Questions or comments? Call 1-800-523-0191

Witch Hazel For Face & Body

ASTRINGENT

Relief of minor skin irritations due to:

- Insect bites
- Minor cuts & scrapes

Active ingredient

Witch hazel 86%

Purpose

Astringent

Inactive ingredient:

alcohol 14% by volume

Use

for relief of minor skin irritations due to:

• insect bites • minor cuts • minor scrapes

Directions

apply as often as needed

10733-801-08



10733-801-16



MEDICAL PRODUCTS LABORATORIES, INC.

NDC 10733-801-16

Witch Hazel FOR FACE & BODY

ASTRINGENT

Relief of minor skin irritations due to:

- Insect bites
- · Minor cuts & scrapes

16 FL OZ (1 PT) 473 mL

Active ingredient Witch hazel 96%	Purpose Astringent
USE for relief at minor s ■ insect bites ■ minor co	kin imitations due to uts II minor scrape:
Warnings For external use only	
When using this product ■ avoid contact with the occurs, rinse thoroughly to	yes. If contact
Stop use and ask a doot condition worsens or sy more than 7 days	
Keep out of reach of child it swallowed, get medical Poison Control Center righ	help or contact a
Directions apply as o	itten as needed
Inactive ingredient	

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WITCH HAZEL

Route of Administration

witch hazel liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:10733-801

TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

WITCH HAZEL (UNII: 10114J0U34) (WTCH HAZEL - UNII:10114J0U34)

WTCH HAZEL (BE in 100 mL

Inactive Ingredients					
Ingredient Name	Strength				
ALCOHOL (UNII: 3K9958V90M)					

P	Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date			
1	NDC:10733- 801-08	236 mL in 1 BOTTLE, DISPENSING; Type 0: Not a Combination Product	01/29/2021				
2	NDC:10733- 801-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/29/2021	11/30/2022			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	M015	01/29/2021				

Labeler - Medical Products Laboratories, Inc. (002290302)

Revised: 12/2024 Medical Products Laboratories, Inc.