WITCH HAZEL READYINCASE- witch hazel liquid P & L Development, LLC

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
- avoid swallowing

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 persist for more than 7 days.

If pregnant or breast-feeding,

ask a health professional before use.

Keep out of reach of children.

If swallowed, get medical help or contact a Poison Control Center (1-800-222-1222) right away.

Directions

apply to the affected area as often as necessary.

Other information

- keep tightly closed
- store at room temperature

Inactive ingredient

alcohol 14% by volume

Principal Display Panel

Distilled extract of

Witch Hazel

Astringent

alcohol 14% by volume

hamamelis water

for relief of minor skin irritations due to

- insect bites
- minor cuts
- minor scrapes

floz (PT) mL

Manufactured by:

PL Developments

1874 Hwy 72 W

Clinton, SC 29325

Questions or comments?

Call 1-877-753-3935 Monday-Friday 9AM-5PM EST

TAMPER EVIDENT: DO NOT USE IF PRINTED SAFETY SEAL UNDER CAP IS BROKEN OR MISSING.

Package Label



WITCH HAZEL READYINCASE

witch hazel liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:49580-0390 Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength WITCH HAZEL (UNII: 10114J0U34) (WTCH HAZEL - UNII:10114J0U34) WTCH 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:49580- 0390-6	473 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/31/2014				
Marketing Information							

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M016	08/31/2014			

Labeler - P & L Development, LLC (101896231)

Revised: 10/2023 P & L Development, LLC