

UNIVERSAL WITCH HAZEL- witch hazel solution
Universal Distribution Center LLC

WITCH HAZEL SOLUTION

Drug Facts

Active Ingredients:

All natural distilled witch hazel 100%
(contains natural grain alcohol 14%)

Purpose

Astringent

Usage:

The soft toner is specifically developed to meet ultimate needs of purifying and conditioning for an array of skin types. It is a calm and non-drying formula which removes remaining traces of soap. It removes make-up or excess oils making your skin feeling soft, healthful, and smooth. It also smoothens the skin after shaving. Use as often as necessary. Apply directly from the bottle or dampen a cotton pad and smoothly wipe it off.

Warning:

For external use only. Avoid contact with eyes. If contact occurs, flush thoroughly with water.

Keep out of reach of children.

Direction:

- remove protective inner seal
- moisten cotton ball or cleansing pad and gently wipe skin
- apply as often as necessary

Ingredients:

Water, Hamamelis Virginiana (Witch Hazel) Leaf Water, Alcohol (14% by Volume), Benzyl Alcohol, Methylchloroisothiazolinone, methylisothiazolinone.

SKIN CLEANSER

MAKE-UP REMOVER

AFTER SHAVING

For Face & Body

Gentle Relief For

100% NATURAL ASTRINGENT

Specifically formulated for purifying and conditioning of the skin.

Distributed by:

Universal Distribution Center LLC

Edison, NJ 08817

www.universaldc.com

This Product is Not Tested on Animals

Made in Jordan

Packaging



SKIN
CLEANSER



MAKE-UP
REMOVER



AFTER
SHAVING

For Face & Body

Gentle Relief For



6 fl. oz. (177 mL)



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Item#55605



Made in Jordan



UNIVERSAL WITCH HAZEL

witch hazel solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-414
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 101I4J0U34) (WTCH HAZEL - UNII:101I4J0U34)	WTCH HAZEL	3 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HAMAMELIS VIRGINIANA LEAF WATER (UNII: 8FP93ED6H2)	
ALCOHOL (UNII: 3K9958V90M)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-414-06	177 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	07/01/2023	

Labeler - Universal Distribution Center LLC (019180459)