WITCH HAZEL- witch hazel liquid Humco Holding Group, Inc.

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.

Private Label Witch Hazel

Drug Facts

Active Ingredient

Witch Hazel

Purpose

Astringent

Indications

For relief of minor skin Irritations due to minor cuts minor scraps

insect bites

Warnings

For external use only avoid contact with eyes

If contact occurs rinse thoroughly with water.

When using this product stop using and contact a doctor if

condition persists or gets worse symptoms do not improve within 7 days

Keep out of reach of children.

In case of accidental ingestion, seek professional assistance or contact a Poison Control Center immediately.

In case of eye contact flush eyes with running water for 15 minutes.

Directions

Apply liberally to the afflicted area as often as necessary

Inactive ingredients

Good Neighbor Label



Sunmark Label



Health Mart Label





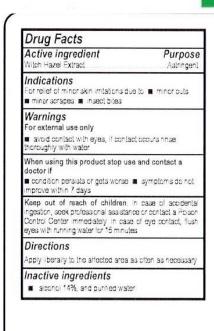
*Warning: Do not use if tamper evident seal is broken or missing. This product is sealed with a breakaway cap ring.

> A46037 02 2023

MSKESSON

Another Quality Product Distributed by McKesson 6555 State Highway 161, Les Celinas, TX 75039 Money Back Guarantee healthmart.com

B012020





broken or missing. This product is sealed with a breakaway cap ring.

MSKESSON

Another Quality Product Distributed by McKesson One Post Street, San Francisco, CA 94104 Money Back Guarantee

R100317 Made in U.S.A.

WITCH HAZEL

witch hazel liquid

Product Information

HUMAN OTC DRUG NDC:0395-9125 Product Type Item Code (Source)

Route of Administration	TOPICAL
Route of Administration	IOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength

WITCH HAZEL (UNII: 10 114J0 U34) (WITCH HAZEL - UNII: 10 114J0 U34)

WITCH HAZEL 855 mg in 1 mL

Inactive Ingredients

macuve ingreatents		
Ingredient Name	Strength	
ALCOHOL (UNII: 3K9958V90M)		
WATER (UNII: 059QF0KO0R)		

Packaging

l	#	Item Code	Package Description	Marketing Start Date	Marketing End Date
ı	1	NDC:0395-9125-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	0 1/0 1/20 16	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	0 1/0 1/20 16	

Labeler - Humco Holding Group, Inc. (825672884)

Registrant - Humco Holding Group, Inc. (825672884)

Establishment			
Name	Address	ID/FEI	Business Operations
Humco Holding Group, Inc.		825672884	analysis(0395-9125), manufacture(0395-9125), pack(0395-9125), label(0395-9125)

Revised: 6/2020 Humco Holding Group, Inc.