WITCH HAZEL- witch hazel solution
CHAIN DRUG MARKETING ASSOCIATION 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.

Witch Hazel
100% Natural Astringent
For Face and Body

Active Ingredient **Purpose**

Witch hazel .....Astringent

Gentle Relief for:
Oily, Irritated, Red, Damaged, Blemished, or Inflamed Skin

Use

for relief of minor skin irritations due to:
- insect bites
- minor cuts
- minor scrapes

Warnings

For external use only

When using this product
- avoid contact with eyes

Stop use and ask a doctor if
- condition worsens or symptoms persist for more than 7 days

Keep out of reach of children. If swallowed, get medical help, or contact a Poison Control Center right away.

Directions

apply as often as needed

Inactive ingredients

alcohol and purified water

Questions or comments?
1-800-935-2362 (Mon-Fri 9am-5pm EST)

Distributed by C.D.M.A. lnc.
43157 W 9 Mile Rd
Novi, MI 48375
www.qualitychoice.com

PRINCIPAL DISPLAY PANEL

QC QUALITY CHOICE

NDC 63868-592-16

Witch Hazel

100% Natural Astringent

For Face and Body

Gentle Relief for:

Oily, Irritated, Red, Damaged, Blemished, or Inflamed Skin

16 FL OZ (473 mL)

TAMPER EVIDENT: DO NOT USE IF INNER BOTTLE SEAL IS BROKEN OR MISSING.

Drug Facts

Active Ingredient	***Purpose***
Witch hazel	Astringent

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■ insect bites ■ minor cuts ■ minor scrapes

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SATISFACTION 100% QC GUARANTEED

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43157 W 9 Mile Rd
Novi, MI 48375
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Question: 800-935-2362

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Lot:

Exp:

WITCH HAZEL

witch hazel solution

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63868-592
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
WITCH HAZEL (UNII: 101I4J0U34) (WITCH HAZEL - UNII:101I4J0U34)	WITCH HAZEL	85 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:63868-592-16	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	02/01/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part347	02/01/2021	

Labeler - CHAIN DRUG MARKETING ASSOCIATION INC (011920774)

Registrant - Seaway Pharma Inc. (117218785)

Establishment

Name	Address	ID/FEI	Business Operations
Seaway Pharma Inc.		117218785	manufacture(63868-592)

Revised: 6/2021 CHAIN DRUG MARKETING ASSOCIATION INC