

SUNMARK WITCH HAZEL- witch hazel liquid
Strategic Sourcing Services LLC

Sunmark 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

Alcohol 14% and purified water.

Principal Display Panel

Drug Facts	
Active ingredient	Purpose
Witch Hazel Extract.....	Astringent
Indications	
For relief of minor skin irritations due to ■ minor cuts ■ minor scrapes ■ insect bites	
Warnings	
For external use only ■ avoid contact with eyes, if contact occurs rinse thoroughly with water.	
When using this product stop use 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 affected area as often as necessary	
Inactive ingredients	
■ alcohol 14%, and purified water	

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

MEKESSON

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Money Back Guarantee
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NDC 49348-181-38

witch

hazel

A lcohol 14%

16 FLOZ (1 PT) 473 mL

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SUNMARK WITCH HAZEL

witch hazel liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49348-181
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
WITCH HAZEL (UNII: 101I4J0U34) (WTCH HAZEL - UNII:101I4J0U34)	WTCH HAZEL	855 mg in 1 mL

Inactive Ingredients

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

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49348-181-38	473 mL in 1 BOTTLE; Type 0: Not a Combination Product	01/01/2016	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	01/01/2016	

Labeler - Strategic Sourcing Services LLC (116956644)

Registrant - Pharma Nobis, LLC (118564114)

Establishment

Name	Address	ID/FEI	Business Operations
Pharma Nobis, LLC		118564114	analysis(49348-181) , manufacture(49348-181) , pack(49348-181) , label(49348-181)

Revised: 12/2023

Strategic Sourcing Services LLC