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

NDC 49348-181-38

witch

hazel

A lcohol 14%

16 FLOZ (1 PT) 473 mL

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) (WITCH HAZEL - UNII:101I4J0U34)	WITCH 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