HAND SPRITZER- ethyl alcohol gel
Hand Spritzer Llc

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.

Hand Spritzer

Drug Facts

Active Ingredient(s)

Ethyl alcohol 80% v/v

Purpose

Antiseptic

Use(s)

Hand Sanitizer to help reduce bacteria that potentially can cause disease.
For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

• in children less than 2 months of age

• on open skin wounds

When using this product keep out of eyes, ears, and mouth. In case of contact with eyes,rinse eyes thoroughly with water.

Stop use and ask a doctor if irritation or rash occurs. These may be signs of a serious condition.

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

Directions

• Place enough product on hands to cover all surfaces. Rub hands together until dry.
• Supervise children under 6 years of age when using this product to avoid swallowing.

Other information

• Store between 15-30C (59-86F)
• Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients RODI water, hydrogen peroxide, glycerin, methocellulose, carbopol, aloe

Ethyl Alcohol Antiseptic

80% Topical Solution

Hand Sanitizer

Non-sterile Solution

Packaging

Drug Facts
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Ethyl Alcohol Antiseptic

80% Topical Solution

Hand Sanitizer

Non-sterile Solution

[235 mL]

NDC 77276-428-08

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Ethyl Alcohol Antiseptic

80% Topical Solution

Hand Sanitizer

Non-sterile Solution

[500 mL]

NDC 77276-428-16

HAND SPRITZER

ethyl alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77276-428
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	80 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
HYDROGEN PEROXIDE (UNII: BBX060AN9V)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLCELLULOSE, UNSPECIFIED (UNII: Z944H5SN0H)	
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)	
ALOE (UNII: V5VD430YW9)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77276-428-08	235 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/23/2020	
2	NDC:77276-428-16	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/23/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/23/2020	

Labeler - Hand Spritzer Llc (130846471)

Revised: 7/2020 Hand Spritzer Llc