

**ZEP FUZION INSTANT HAND SANITIZER- alcohol liquid**  
**Zep Inc.**

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**66949-119 / 3558 Zep Fuzion Instant Hand Sanitizer**

***Active Ingredient***

Ethanol 70% v/v

***Purpose***

Antiseptic

***Uses***

- To help reduce germs and bacteria on the skin.
- Recommended for repeated use.
- No rinsing required.

***Warnings***

**Flammable.** Keep away from fire, flame, or spark. **For external use only.**

**Do not use**

in the eye; if in eyes, rinse thoroughly with water.

**When using this product**

- Do not swallow.
- If swallowed, do not induce vomiting and if individual is conscious, give large quantities of water to drink and consult a physician immediately.

**Stop use and ask doctor**

**Stop use and ask doctor if** skin irritation or redness persists for more than 72 hours.

**Keep out of reach of children and pets**

**Keep out of reach of children and pets. Children must be supervised in use of this product.**

***Directions***

- Apply gel to hands.
- Rub into hands for at least 20 seconds or until dry.

### Other information

- Keep container closed and stored in a dry area at temperatures between 68°F and 77°F (20°C and 25°C).
- Do not reuse empty container.
- Dispose in accordance with all applicable federal, state and local regulations

### Inactive ingredients

Deionized Water, PEG-6 (and) Acrylates/Vinyl Crosspolymer, Fragrance.

### Questions or comments?

**Call 1-877-I-BUY-ZEP (1-877-428-9937)**

### Package Label - Principal Display Panel



### ZEP FUZION INSTANT HAND SANITIZER

alcohol liquid

#### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66949-119
<b>Route of Administration</b>	TOPICAL		

#### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

#### Inactive Ingredients

Ingredient Name	Strength
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	
WATER (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-119-16	4800 mL in 1 CASE; Type 0: Not a Combination Product	11/02/2020	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	11/02/2020	

**Labeler -** Zep Inc. (030471374)

## Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-119)

Revised: 10/2023

Zep Inc.