

**AFCO 5511 SANIFECT E3- ethanol liquid**  
**Zep Inc.**

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**66949-140 / 0900 AFCO Sanifect E3**

***Active ingredient***

Ethanol 70% v/v

***Purpose***

Antiseptic

***Uses***

- Hand sanitizing to decrease bacteria on skin.
- No rinsing required.

***Warnings***

**For external use only.**

**Do not use** in the eyes; if in eyes, rinse promptly and thoroughly with water.

**Flammable.** Keep away from fire, flame or spark.

**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 a doctor**

**Stop use and ask a 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***

- Wet hands thoroughly with spray mist.
- Rub hands together allowing liquid to contact all areas, especially around the nails and cuticles.
- Continue rubbing vigorously until hands are dry.

- No rinsing or toweling is required.

**Other information**

Store at 4 to 27°C (40 to 80°F).

Dispose in accordance with all applicable federal, state and local regulations.

**Inactive ingredients**

Deionized water; Glycerine

**Questions or comments?**

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



<b>AFCO 5511 SANIFECT E3</b>			
ethanol liquid			
<b>Product Information</b>			
<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:66949-140
<b>Route of Administration</b>	TOPICAL		
<b>Active Ingredient/Active Moiety</b>			
<b>Ingredient Name</b>		<b>Basis of Strength</b>	<b>Strength</b>
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)		ALCOHOL	70 mL in 100 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>WATER</b> (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-140-01	208198 mL in 1 DRUM; Type 0: Not a Combination Product	05/05/2025	
2	NDC:66949-140-02	6000 mL in 1 CASE; Type 0: Not a Combination Product	05/05/2025	
3	NDC:66949-140-03	3785 mL in 1 CASE; Type 0: Not a Combination Product	05/05/2025	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	05/05/2025	

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

## Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-140)

Revised: 12/2025

Zep Inc.