

ZEP INSTANT HAND SANITIZER- ethanol liquid
Zep Inc.

66949-246 / 0878 Instant Hand Sanitizer

☐ **Active Ingredient**

Alcohol 60%

☐ **Purpose**

Antiseptic

☐ **Uses**

- Hand sanitizing to decrease bacteria on 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 quantity 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 the use of this product.

☐ **Directions**

- Use pump and apply gel to hands.
- 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 20 to 25°C (68 to 77°F).
- Dispose in accordance with all applicable federal, state and local regulations.

☐ **Inactive ingredients**

Deionized Water, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Diisopropanolamine, Fragrance

☐ **Questions or comments?**

Call 1-800-I-BUY-ZEP (1-800-428-9937)





Instant Hand Sanitizer
Gel

ZEP INSTANT HAND SANITIZER

ethanol liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:66949-246
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	60 mL in 100 mL

Inactive Ingredients

Ingredient Name	Strength
CARBOMER INTERPOLYMER TYPE A (55000 CPS) (UNII: 59TL3WG5CO)	
DIISOPROPANOLAMINE (UNII: 0W44HYL8T5)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-246-16	4800 mL in 1 CASE; Type 0: Not a Combination Product	01/29/2004	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/29/2004	12/31/2024

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-246)

Revised: 10/2023

Zep Inc.