ZEP PROFESSIONAL ALCOHOL HAND SANITIZING- ethanol liquid
Zep Inc.

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.

Zep Professional Alcohol Hand Sanitizing Gel

Active ingredient

Alcohol 60%

Purpose

Antiseptice

Uses

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

Warnings

For external use only.

Flammable. Keep away from fire, flame or spark.

When using this product

- Avoid eye contact.
- If in eyes, rinse promptly and thoroughly with water.

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

If swallowed, get medical help or contact a Poison Control Center immediately.

Keep out of reach of children and pets.

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 allowed 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.

Inactiv ingredients

deionized water, acrylates/C10-30 alkyl acrylate crosspolymer, diisopropanolamine, fragrance

Questions or comments?

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

ZEP PROFESSIONAL ALCOHOL HAND SANITIZING

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-109
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	6 mL in 10 mL

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

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66949-109-00	500 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/07/2009	
2	NDC:66949-109-01	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/07/2009	

3	NDC:66949-109-24	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/07/2009	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/07/2009	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		030471374	manufacture(66949-109)

Revised: 11/2016 Zep Inc.