ZEP ALCOHOL SANITIZER- 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 Alcohol Sanitizer

Active ingredient

Alcohol 62%

Purpose

Antiseptic

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.

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

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

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

Inactive ingredients

Water, Glycerine

Questions or comments?

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

Net Contents: 33.8 fl. oz. (1 Liter)

ZEP ALCOHOL SANITIZER

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-900
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	6.2 mL in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66949-900-00	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2000	
2	NDC:66949-900-21	3785 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2000	
3	NDC:66949-900-85	208198 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2000	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/07/2000	

ZEP ALCOHOL SANITIZER

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-357
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	6.2 mL in 10 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66949-357-15	1200 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	08/07/2000	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	08/07/2000	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		030471374	manufacture(66949-900, 66949-357)

Revised: 11/2016 Zep Inc.