ZEP INSTANT HAND SANITIZER GEL- alcohol liquid
Zep Inc.

66949-117 / 3558 Zep Instant Hand Sanitizer Gel 70%

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 INSTANT HAND SANITIZER GEL

alcohol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-117
Route of Administration	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**
ACRYLATES/VINYL ISODECANOATE CROSSPOLYMER (10000 MPA.S NEUTRALIZED AT 0.5%) (UNII: 2N8MDB79NA)	
WATER (UNII: 059QF0KO0R)	
POLYETHYLENE GLYCOL 300 (UNII: 5655G9Y8AQ)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:66949-117-01	6000 mL in 1 CASE; Type 0: Not a Combination Product	09/28/2020	
2	NDC:66949-117-24	15140 mL in 1 CASE; Type 0: Not a Combination Product	09/28/2020	
3	NDC:66949-117-32	11400 mL in 1 CASE; Type 0: Not a Combination Product	09/28/2020	
4	NDC:66949-117-10	3785 mL in 1 CASE; Type 0: Not a Combination Product	09/28/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/28/2020	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-117)

Revised: 3/2025 Zep Inc.