HAND SANITIZER GEL- ethanol liquid
Zep Inc.

66949-137 / 4508 HDX Hand Sanitizer Gel

Active ingredient

Ethanol 70% v/v

Purpose

Antiseptic

Uses

■ To help reduce bacteria that can potentially cause disease

■ For use when soap and water are not available

Warnings

For external use only.

Flammable. Keep away from source of heat or fire

Do Not Use

■ on children less than 2 months of age

■ on open skin wounds

■ on broken or damaged skin

When using this product

When using this product avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water. Do not inhale.

Stop use and ask a doctor

Stop use and ask a doctor if irritation or redness develops and lasts.

Keep out of reach of children and pets

Keep out of reach of children and pets. In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

■ For occasional and personal use.

■ Rub thoroughly into hands for at least 30 seconds. Allow to dry.

■ Children under 6 years should be supervised when using this product.

Other information

■ store at a temperature below 110 °F (43 °C).

■ may discolor certain fabrics or surfaces.

Inactive ingredients

Water, Triisopropanolamine, Acrylates/C10-30 Alkyl Acrylate Crosspolymer, Tocopheryl Acetate.

Questions?

1-800-514-6729

HAND SANITIZER GEL

ethanol liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-137
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 CROSSPOLYMER-6 (UNII: 4GXD0Q3OS3)	
ALPHA-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
WATER (UNII: 059QF0KO0R)	
TRIISOPROPANOLAMINE (UNII: W9EN9DLM98)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-137-28	15140 mL in 1 CASE; Type 0: Not a Combination Product	07/05/2021	
2	NDC:66949-137-08	2840 mL in 1 CASE; Type 0: Not a Combination Product	07/05/2021	12/31/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	07/05/2021	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Zep Inc.		112125310	manufacture(66949-137)

Revised: 10/2023 Zep Inc.