HAND SANITIZER- ethyl alcohol spray UpLift Brands, LLC

Germ-X 899.000-899AA 99% Natural Germ-X Hand Sanitizer Spray Citrus Scent

Active ingredient

Ethyl alcohol 62%

Purpose

Antiseptic

Uses

- to decrease bacteria on the skin that could cause disease
- recommended for repeated use

Warnings

For external use only: hands

Flammable, keep away from fire or flame

When using this product

- keep out of eyes. In case of contact with eyes, flush thoroughly with water.
- avoid contact with broken skin
- do not inhale or ingest

Stop use and ask a doctor if

- irritation or redness develop
- condition persists for more than 72 hours

Keep out of reach of children.

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

Directions

- spray hands thoroughly with product and allow to dry without wiping
- for children under 6, use only under adult supervision
- not recommended for infants

Other information

- do not store above 105°F
- may discolor some fabrics
- harmful to wood finishes and plastics

Inactive ingredients

water, glycerin, Aloe barbadensis leaf juice, fragrance, limonene

Questions?

1-888-593-0593

Distributed By:

Vi-Jon, LLC,

St. Louis, MO 63114

Disclaimer

*Effective at eliminating 99.99% of many common harmful germs & bacteria in as little as 15 seconds.

FORMULA MADE IN USA

EMPLOYEE OWNED

principal display panel

germ-X_®

SINCE 1997

KILLS 99.99% OF GERMS*

HAND SANITIZER SPRAY

CITRUS SCENT

*KILLS GERMS IN 15 SECONDS

FORMULA MADE IN USA

EMPLOYEE OWNED

2 FL OZ (59 mL)





HAND SANITIZER

ethyl alcohol spray

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:83986-899

Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name

Basis of Strength

ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII: 3K9958V90M)

ALCOHOL

ALCOHOL

62 mL in 100 mL

Inactive Ingredients Ingredient Name Strength WATER (UNII: 059QF0KO0R) GLYCERIN (UNII: PDC6A3C0OX) ALOE VERA LEAF (UNII: ZY81Z83H0X) LIMONENE OXIDE, (+)- (UNII: 278IM94GXB)

Packaging						
#	Item Code	Package Description	Marketing Start Date	Marketing End Date		
1	NDC:83986- 899-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	06/06/2016			

Marketing Information						
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC Monograph Drug	505G(a)(3)	06/06/2016				

Labeler - UpLift Brands, LLC (119091527)

Revised: 4/2024 UpLift Brands, LLC