

HAND SANITIZER- ethyl alcohol spray
UpLift Brands, LLC

Germ-X 721.001/721AB
99% Natural Germ-X Hand Sanitizer Spray Lavender 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

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

Disclaimer

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

Distributed By: Vi-Jon, LLC

8515 Page Ave., St. Louis, MO 63114

www.germx.com

FORMULA MADE IN THE USA

EMPLOYEE-OWNED

Principal display panel

germ-X®

SINCE 1997

KILLS 99.99% OF GERMS

HAND SANITIZER SPRAY

LAVENDER 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:83986-721
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	0.62 mL in 1 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:83986-721-20	59 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	04/26/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
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OTC Monograph Drug	505G(a)(3)	04/26/2024	
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Labeler - UpLift Brands, LLC (119091527)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-721)

Revised: 4/2024

UpLift Brands, LLC