

**HAND SANITIZER- ethyl alcohol gel**  
**UpLift Brands LLC**

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**Germ-X Advanced Hand Sanitizer Original Scent**  
**529.000/529AA rev 1**

**Active Ingredient**

Ethyl alcohol 70%

**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 heat and 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 and redness develops
- 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**

- wet 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, glyceryl caprylate/caprinate, glycerin, isopropyl myristate, tocopheryl acetate, carbomer or acrylates/C10-30 alkyl acrylate crosspolymer, fragrance, benzophenone-4

**Disclaimer**

DSP-TN-21091 DSP-MO-20087 DSP-MO-28 DSD-MO-20068

\*Effective at eliminating more than 99.99% of many common harmful germs and bacteria in as little as 15 seconds.

FORMULA MADE IN USA

EMPLOYEE OWNED

**Adverse reaction**

Dist. by: Vi-Jon, LLC 8515 Page Ave

St. Louis, MO 63114

**Principal display panel**

germ-x®

since 1997

ADVANCED

70% alcohol hand sanitizer

Kills more than 99.99% of Germs\*

with added moisturizers

33.8 FL OZ (1L) 1.05QT



L0019747FC



33.8 FL OZ (1L) 1.05 QT

## HAND SANITIZER

ethyl alcohol gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83986-529
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>ALCOHOL</b> (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	70 mL in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>GLYCERYL CAPRYLATE/CAPRATE</b> (UNII: G7515SW10N)	
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	

<b>ISOPROPYL MYRISTATE</b> (UNII: 0RE8K4LNJS)	
<b>.ALPHA.-TOCOPHEROL ACETATE</b> (UNII: 9E8X80D2L0)	
<b>CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE</b> (UNII: 0A5MM307FC)	
<b>CARBOMER COPOLYMER TYPE B (ALLYL PENTAERYTHRITOL CROSSLINKED)</b> (UNII: 809Y72KV36)	
<b>SULISOBENZONE</b> (UNII: 1W6L629B4K)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-529-16	59 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/22/2022	
2	NDC:83986-529-34	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/22/2022	
3	NDC:83986-529-32	355 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/22/2022	
4	NDC:83986-529-50	710 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/22/2022	
5	NDC:83986-529-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	12/22/2022	

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M505G(a)(3)	12/22/2022	

**Labeler** - UpLift Brands LLC (119091527)

**Registrant** - Consumer Product Partners, LLC (119091520)

### Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-529)

### Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83986-529)

Revised: 2/2024

UpLift Brands LLC