

HAND SANITIZER- alcohol gel
UpLift Brands, LLC

Germ-X 456.001/456AC
70% Alcohol Foaming Hand Sanitizer

Active ingredient

Ethyl alcohol 70%

Purpose

Antiseptic

Use

- 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 develops
- condition persists for more than 72 hours

Keep out of reach 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, PEG-8 Dimethicone, meadowfoamamidopropyl betaine, glyceryl caprylate/caprate, glycerin, tocopheryl acetate, isopropyl myristate

Adverse reaction

DSP-TN-21091 DSP-MO-20087

Pat. 9,161,982

DISTRIBUTED BY: UPLIFT BRANDS, LLC

ST. LOUIS, MO 63114

1-866 MY GERM X

Principal display panel

germ-x[®]

PRO

ADVANCED

FOAMING HAND SANITIZER

1000 mL (1.05 QT) 33.8 FL OZ



Drug Facts

Active Ingredient	Purpose
Ethyl alcohol 70%	Antiseptic

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

Warnings
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Flammable, keep away from fire or flame.
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Stop use and ask a doctor if ■ irritation and 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.

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ST. LOUIS, MO 63114 • 1-876-MY GERMOX
L001813188 PaL 9.181.962

HAND SANITIZER

alcohol gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83986-456
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
WATER (UNII: 059QF0K00R)	
PEG-8 DIMETHICONE (UNII: GIA7T7640D)	
MEADOWFOAMAMIDOPROPYL BETAINE (UNII: HNV0L650LG)	
GLYCERYL CAPRYLATE/CAPRATE (UNII: G75155W10N)	
GLYCERIN (UNII: PDC6A3C00X)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging

Marketing Start Marketing End

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-456-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2024	
2	NDC:83986-456-04	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2024	
3	NDC:83986-456-44	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2024	12/31/2025
4	NDC:83986-456-86	1000 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/15/2024	
5	NDC:83986-456-77	500 mL in 1 POUCH; Type 0: Not a Combination Product	04/15/2024	12/20/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/15/2024	

Labeler - UpLift Brands, LLC (119091527)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-456)

Revised: 1/2026

UpLift Brands, LLC