MOISTURIZING HAND SANITIZER- ethyl alcohol gel
UpLift Brands, LLC

Germ-X 447.001/447AC
Moisturizing Foaming Hand Sanitizer

Claims

germ-x

PRO

MOISTURIZING

FOAMING HAND SANITIZER

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 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.

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, glycerin, tocopheryl acetate, isopropyl myristate

Adverse Reactions

DISTRIBUTED BY: UPLIFT BRANDS, LLC

ST. LOUIS, MO 63114

1-866 MY GERMX

DSP-TN-21091 DSP-MO-20087

Principal display panel

germ-X®

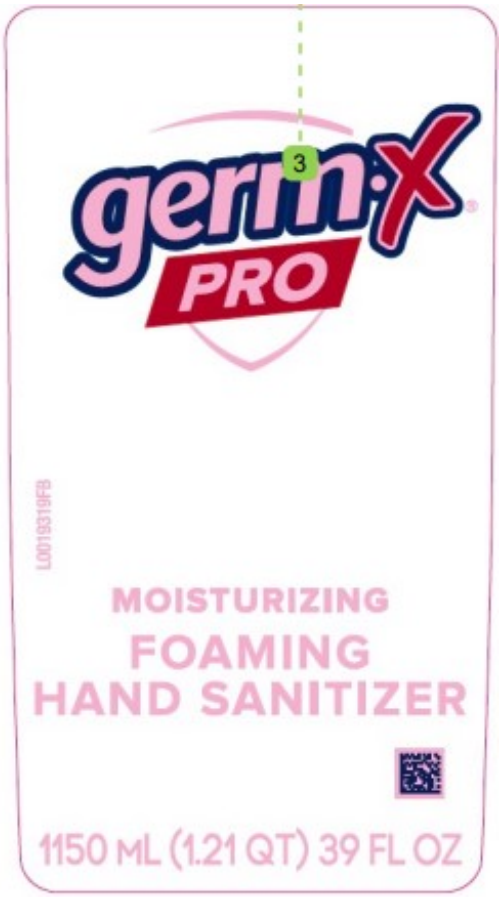
PRO

MOISTURIZING

FOAMING

HAND SANITIZER

1150 mL (1.21 QT) 39 FL OZ



MOISTURIZING HAND SANITIZER

ethyl alcohol gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:83986-447
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
ALCOHOL (UNII: 3K9958V90M) (ALCOHOL - UNII:3K9958V90M)	ALCOHOL	62 mL in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
PEG-8 DIMETHICONE (UNII: GIA7T764OD)	
MEADOWFOAMAMIDOPROPYL BETAINE (UNII: HNV0L650LG)	
GLYCERIN (UNII: PDC6A3C0OX)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-447-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/12/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505(a)(3)	04/12/2024	

Labeler - UpLift Brands, LLC (119091527)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-447)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83986-447)

Revised: 11/2024 UpLift Brands, LLC