

HAND WASH- benzalkonium chloride soap
UpLift Brands, LLC

Germ-X 460.001/460AB
Antibacterial Foaming Hand Wash

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

for handwashing to decrease bacteria on the skin

Warnings

For external use only: hands only

When using this product

- avoid contact with eyes. If contact occurs, rinse eyes thoroughly with water.

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
- apply palmful to hands
- scrub thoroughly
- rinse thoroughly

Inactive ingredients

water, cocamidopropyl betaine, lauramidopropylamine oxide, lauramine oxide, myristamidopropylamine oxide, glycerin, fragrance, citric acid, tetrasodium EDTA,

benzophenone-4, sodium benzoate, red 4, yellow 5

Adverse reaction

DISTRIBUTED BY: UPLIFT BRANDS, LLC
ST. LOUIS, MO 63114 1-866 MY GERM X

Principal display panel

germ-x[®]

PRO

ANTIBACTERIAL

FOAMING

HAND SOAP

TRICLOSAN FREE

Helps kill harmful germs

Quick foaming

Gentle and mild formula

7.5 FL OZ (221 mL)



HAND WASH

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:83986-460
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
GLYCERIN (UNII: PDC6A3C0OX)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SULISOBENZONE (UNII: 1W6L629B4K)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-460-96	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	
2	NDC:83986-460-44	532 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	02/15/2025
3	NDC:83986-460-04	750 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	
4	NDC:83986-460-45	1150 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	02/05/2024	
5	NDC:83986-460-08	3780 mL in 1 BOTTLE, PLASTIC; Type 1: Convenience Kit of Co-Package	02/05/2024	02/15/2025

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)	02/05/2024	
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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(83986-460)

Establishment			
Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83986-460)

Revised: 5/2025

UpLift Brands, LLC