

ANTIBACTERIAL HAND WASH SPRING RAIN- benzalkonium chloride soap
UpLift Brands, LLC

Germ-X D12.000/D12AB
Antibacterial Hand Soap Spring Rain

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 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, lauramine oxide, cocamidopropyl betaine, lauramidopropylamine oxide, glycerin, myristamidopropylamine oxide, fragrance, hexyl cinnamal, linalool, limonene, disteareth-75 IPDI, PEG-150 distearate, citric acid, sodium chloride, tetrasodium EDTA, benzophenone-4, sodium benzoate, ext. violet 2, green 3

Adverse reaction

Distributed by: Vi-Jon, LLC, St. Louis, MO 63114

FORMULA MADE IN USA

EMPLOYEE-OWNED

Principal panel display

germ-x

ANTIBACTERIAL

LIQUID HAND SOAP

SPRING RAIN

Dermatologist Tested

12FL OZ (355 ML)

germ-x



ANTIBACTERIAL HAND WASH SPRING RAIN

benzalkonium chloride soap

Product Information

Product Type

HUMAN OTC DRUG

Item Code (Source)

NDC:83986-012

Route of Administration	TOPICAL
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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
EDETATE SODIUM (UNII: MP1J8420LU)	
SULISOBENZONE (UNII: 1W6L629B4K)	
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
GLYCERIN (UNII: PDC6A3C0OX)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
.ALPHA.-HEXYLCINNAMALDEHYDE (UNII: 7X6O37OK2I)	
LINALOOL, (+/-)- (UNII: D81QY6I88E)	
LIMONENE, (+)- (UNII: GFD7C86Q1W)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
EXT. D&C VIOLET NO. 2 (UNII: G5UX3K0728)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83986-012-32	355 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	10/12/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	10/12/2022	

Labeler - UpLift Brands, LLC (119091527)

Registrant - Consumer Product Partners, LLC (119091520)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091520	manufacture(83986-012)

Establishment

Name	Address	ID/FEI	Business Operations
Consumer Product Partners, LLC		119091514	manufacture(83986-012)

Revised: 3/2024

UpLift Brands, LLC