ZEP HANDSTAND ANTIMICROBIAL- benzalkonium chloride liquid
Zep Inc.

66949-127 / U429 Handstand Antimicrobial

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial handwashing

Uses

Handwash to help decrease bacteria on the skin.

Warnings

For external use only.

Do not use

Do not use in the eyes. In case of eye contact, immediately flush with water.

Stop use and ask a doctor

Stop use and ask a doctor if irritation or rash appears and lasts.

Keep out of reach of children

Keep out of reach of children. If swallow, get medical help or contact a Poison Control Center right away.

Directions

- Apply a small amount, covering hands with product for 30 seconds. Add water, lather and rinse.
- Children under 6 years of age should be supervised when using this product.

Inactive Ingredients

Water, Cetrimonium Chloride, Sodium Chloride, Lauramine Oxide, Sorbitol, Disodium EDTA, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Sodium Lauriminodipropionate, Citric Acid, Fragrance, Methylisothiazolinone, Methylchloroisothiazolinone, Yellow 5, Red

1-877-I-BUY-ZEP/1-877-428-9937

ZEP HANDSTAND ANTIMICROBIAL

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-127
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SODIUM LAURIMINODIPROPIONATE (UNII: 7G447D0DH9)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
SORBITOL (UNII: 506T60A25R)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
WATER (UNII: 059QF0KO0R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-127-01	9600 mL in 1 CASE; Type 0: Not a Combination Product	08/23/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/23/2017	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(66949-127)

Revised: 10/2023 Zep Inc.