ZEP HANDSTAND ANTIMICROBIAL- benzalkonium chloride liquid
Zep Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

66949-429 Handstand Antimicrobial

Benzalkonium Chloride 0.13%

Antibacterial handwashing

Uses

Handwash to help decrease bacteris on the skin.

For external use only.

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

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

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.

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 33

ZEP HANDSTAND ANTIMICROBIAL

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:66949-429
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
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
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)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:66949-429-01	800 mL in 1 BOX; 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 not final	part333A	08/23/2017	

Labeler - Zep Inc. (030471374)

Establishment

Name	Address	ID/FEI	Business Operations
Kutol Products Company		004236139	manufacture(66949-429)

Revised: 12/2019 Zep Inc.