STERIL WIZE ANTI BACTERIAL LIQUID HAND- benzalkonium chloride soap
Nostalgia Products LLC

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.

Steril-Wize Anti-Bacterial Liquid Hand Soap

Drug Facts

Active ingredient

Benzalkonium Chloride .13%

Purpose

Antiseptic

Uses

- For handwashing to decrease bacteria on the skin.

Warnings

For external use only.

When using this product

avoid contact with eyes. In case of eye contact, flush eyes with water. Do not ingest.

Stop use and ask a doctor

If irritation or redness develops and conditions persist for more than 72 hours.

Keep out of reach of children.

If swallowed, get medical help or contact the Poison Control Center right away.

Directions

- Apply soap into hands, wet as needed
- Lather vigorously for at least 15 seconds · Wash skin, rinse thoroughly and dry.

Inactive Ingredients

Water,Cocamidopropyl Betaine, PEG 150 Distearate, Caprylyl Glucoside, Glycerine, Cacomide PAMMI, Tetrasodium EDTA, Fragrance, Citric Acid, Aloe Vera, Benzisothiazol, Methylisothiazol, Yellow #5, Red #40.

Package Labeling:

STERIL WIZE ANTI BACTERIAL LIQUID HAND

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77051-002
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)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CAPRYLYL GLUCOSIDE (UNII: V109WUT6RL)	
GLYCERIN (UNII: PDC6A3C0OX)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77051-002-32	946.35 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/16/2020	

Labeler - Nostalgia Products LLC (080132557)

Revised: 7/2020 Nostalgia Products LLC