STERIL-WIZE ANTI-BACTERIAL FOAMING 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 Foaming 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.

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, Glycerine, Caprylyl Glucoside, Tetrasodium EDTA, Benzisothiazol, Methylisothiazol, Fragrance, Citric Acid, Aloe Vera, Yellow #5, Red #40.

Package Labeling:946.35 mL





Package Labeling:3.785L



STERIL-WIZE ANTI-BACTERIAL FOAMING HAND

benzalkonium chloride soap

Produ	duct Information			
Produc	t T yp e	HUMAN OTC DRUG	Item Code (Source)	NDC:77051-001
Route o	f Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
COCAMIDO PRO PYL BETAINE (UNII: 50CF3011KX)		

GLYCERIN (UNII: PDC6A3C0OX)	
CAPRYLYL GLUCO SIDE (UNII: V109 WUT6 RL)	
EDETATE SO DIUM (UNII: MP1J8420LU)	
CITRIC ACID MONO HYDRATE (UNII: 2968 PHW8 QP)	
ALOE VERA LEAF (UNII: ZY8 1Z8 3H0 X)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

F	Packaging			
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77051-001- 32	946.35 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/16/2020	
2	NDC:77051-001- 37	3785 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