ALPHA FOAMING E-2- benzalkonium chloride soap Whisk Products, 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.

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Skin Antimicrobial

Uses

- Reduces amount of bacteria on hands
- Recommended for repeated use

Warnings

For external use only.

When using this product, avoid contact with eyes. In case of eye contact, flush eyes with water.

Stop use and ask a doctor if irritation or redness develops, or if 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

- Pump a small amount of foam into palm of hand.
- Rub thoroughly over all surfaces of both hands for at least fifteen seconds.
- Rinse with potable water.

Inactive Ingredients

Water, Lauramine Oxide, Glycerin, PEG-120 Methyl Glucose Dioleate







Hand Sanitizing Wash

Drug Facts

Active Ingredient Purpose Benzalkonium Chloride 0.1% Skin Antimicrobial

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Whisk Products, Inc. Wentzville, MO 63385

Contains: Eight Bags

Net Contents Each: 33.8 fl. oz. (1 qt. 1.8 fl. oz.) 1000 mL Total Net Volume: 270.4 fl. oz. (2 gal. 14.4 fl. oz.) 8 L

ALPHA FOAMING E-2

benzalkonium chloride soap

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:65585-514

Route of Administration TOPICAL

Active Ingredient/Active Moiety

8		
Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM -	BENZALKONIUM	1 mg
UNII:7N6JUD5X6Y)	CHLORIDE	in 1 mL

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)				
LAURAMINE O XIDE (UNII: 4F6 FC4MI8 W)				
GLYCERIN (UNII: PDC6A3C0OX)				

Product Characteristics				
Color	white (colorless, water-white (dispensed as a white foam))	Score		
Shape		Size		
Flavor		Imprint Code		
Contains				

1	ackaging			
#	t Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:65585-514-01	4 in 1 BOX	04/23/2020	
1		1000 mL in 1 BAG; Type 0: Not a Combination Product		

2 NDC:65585-514-02	8 in 1 BOX	04/23/2020	
2	1000 mL in 1 BAG; Type 0: Not a Combination Product		
Marketing Information			
Wat Keing Intelligence			
Marketing Categor	ry Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not fin	nal part333E	04/23/2020	

Labeler - Whisk Products, Inc. (834270639)

Establishment				
Name	Address	ID/FEI	Business Operations	
Whisk Products, Inc.		834270639	manufacture(65585-514)	

Revised: 4/2020 Whisk Products, Inc.