ULTRA PROFESSIONAL ANTIBACTERIAL FOAMING HAND- 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

Use

reduces amount of bacteria on hands

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

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

Directions

- Apply a palmful to hands and forearms.
- Scrub thoroughly for at least 15 seconds.
- Rinse completely and dry.

Inactive Ingredients

Water, Lauramine Oxide, Glycerin, PEG-120 Methyl Glucose Dioleate, Fragrance, Yellow 5, Blue 1





To reorder, contact your Ultra Professional Distributor 816-876-2266 www.UltraProLine.com

Contains 8 Bags

LBL1657-1.0

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

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LOT#

EXP:

ULTRA PROFESSIONAL ANTIBACTERIAL FOAMING HAND

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:65585-584
Route of Administration	TOPICAL		

l	Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength	
	BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6 Y)	BENZALKONIUM CHLORIDE	1 mg in 1 mL	

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
LAURAMINE O XIDE (UNII: 4F6 FC4MI8 W)			
GLYCERIN (UNII: PDC6A3C0OX)			
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)			
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			

Product Characteristics			
Color	green (apple green (dispensed as a white foam))	Score	
Shape		Size	
Flavor		Imprint Code	
Contains			

l	Packaging				
	# Item Code	Package Description	Marketing Start Date	Marketing End Date	
	1 NDC:65585-584-01	8 in 1 BOX	04/17/2019		
ı	1	1000 mL in 1 BAG; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/17/2019	

Labeler - Whisk Products, Inc. (834270639)

Establishment				
Name	Address	ID/FEI	Business Operations	

Whisk Products, Inc. 834270639 manufacture(65585-584)

Revised: 4/2019 Whisk Products, Inc.