HIMARK- benzalkonium chloride liquid
Apollo Health and Beauty Care

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.

63148-401, HiMark Antibacterial Hand soap Refill.

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

Helps eliminate bacteria on hands

Warnings

For External Use only

When using this product

- Avoid contact with eyes. In case of contact, rinse thoroughly with water.

Stop use and ask a doctor if

Stop use and ask a doctor if

- irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, seek medical attention or contact a poison control center immediately.

Directions

- From Liquid Hand Soap pump bottle, apply onto wet hands
- Lather and rinse thoroughly.

Other Information

Store at room Temperature.

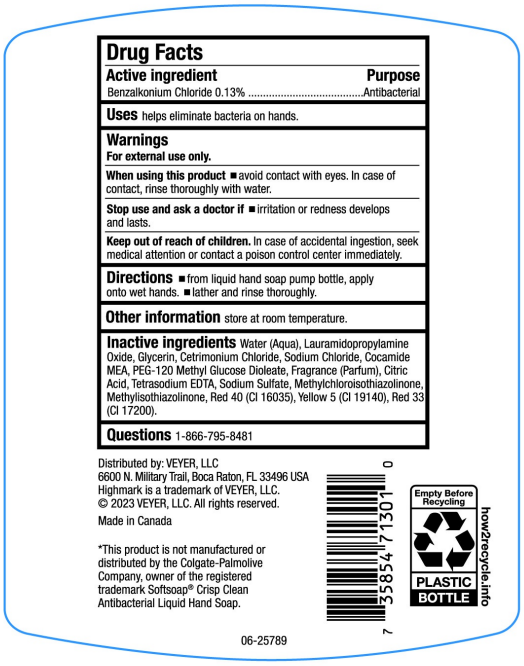
Inactive Ingredients

Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5(CI19140), Red 33 (CI17200).

Questions

1-866-795-8481

Principal Display Panel- 50 Oz



Principal Display Panel-7.5 Oz

HIMARK

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:63148-401
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	130 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	
BASIC YELLOW 5 (UNII: 07BP340B4T)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYL GLUCOSE DIOLEATE (UNII: FA9KFJ4Z6P)	

METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-401-07	221 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2023	
2	NDC:63148-401-50	1478 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	06/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	06/01/2023	

Labeler - Apollo Health and Beauty Care (201901209)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care		201901209	manufacture(63148-401)

Revised: 6/2023 Apollo Health and Beauty Care