

DISCOUNT DRUG MART- benzalkonium chloride liquid
Apollo Health and Beauty Care

Gold Antibacterial Liquid Hand Soap 63148-140

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

Helps decrease 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

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 pump bottle, apply onto wet hands. Lather and rinse thoroughly.

Inactive ingredients

Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Sodium Sulfate, Fragrance (Parfum), Tetrasodium EDTA, Citric Acid, Methylchlorisothiazolinone, Methylisothiazolinone, Yellow 5 (CI 19140), Red 4 (CI 14700)

NDC 63148-146-11



AN EMPLOYEE OWNERSHIP COMPANY

Gold Antibacterial liquid hand soap

Benzalkonium Chloride 0.13%
Antibacterial Hand Soap

COMPARE TO
Dial® Gold Antibacterial
Liquid Hand Soap*

06-27031



**KILLS 99.9%
OF BACTERIA**
PARABEN FREE
PHTHALATE FREE

11 FL OZ (325 mL)

Drug Facts

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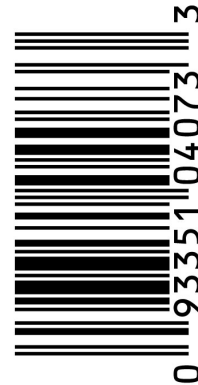
Inactive ingredients Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Sodium Sulfate, Fragrance (Parfum), Tetrasodium EDTA, Citric Acid, Methylchlorisothiazolinone, Methylisothiazolinone, Yellow 5 (CI 19140), Red 4 (CI 17400).

DISTRIBUTED BY: DISCOUNT DRUG
MART INC. MEDINA, OH 44256
WWW.DISCOUNT-DRUGMART.COM
QUESTIONS OR COMMENTS?
330-725-2340
MADE IN CANADA

*This product is not manufactured or distributed by the Henkel Canada Corporation, owner of the registered trademark Dial® Gold Antibacterial Liquid Hand Soap.

Created responsibly at a
LEED-certified facility.

06-27032





Drug Facts	
Active ingredient	Purpose
Benzalkonium Chloride 0.13%	Antibacterial
Uses helps decrease 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 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 ■ Use only to refill a Liquid Hand Soap pump bottle. ■ From pump bottle, apply onto wet hands. ■ Lather and rinse thoroughly.	
Inactive ingredients Water (Aqua), Lauramidopropylamine Oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Sodium Sulfate, Fragrance (Parfum), Tetrasodium EDTA, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, Yellow 5 (CI 19140), Red 4 (CI 17400).	

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SATISFACTION GUARANTEED
 IF DISSATISFIED, RETURN UNUSED PORTION AND PACKAGE TO STORE WHERE PURCHASED. IF UNABLE TO RETURN TO STORE, SEND REASON FOR DISSATISFACTION, NAME, ADDRESS AND EMPTY PACKAGE TO: DISCOUNT DRUG MART, 211 COMMERCE DRIVE, MEDINA, OHIO 44256

Created responsibly at a LEED-certified facility.



06-27034

DISCOUNT DRUG MART

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:63148-146
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
COCAMIDE MEA (UNII: C80684146D)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
WATER (UNII: 059QF0KO0R)	
YELLOW 5 (UNII: I753WB2F1M)	
CITRIC ACID (UNII: 2968PHW8QP)	
GLYCERIN (UNII: PDC6A3C0OX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
TETRASODIUM EDTA (UNII: MP1J8420LU)	

RED 4 (UNII: X3W0AM1JLX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:63148-146-52	1540 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/28/2026	
2	NDC:63148-146-11	325 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/28/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	04/28/2026	

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(63148-146)

Revised: 4/2026

Apollo Health and Beauty Care