

HANDSOAP- benzalkonium chloride soap
CVS Pharmacy

CVS, 51316-703

Purpose

Antibacterial

Benzalkonium Chloride.

For External Use only

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

ask a doctor if Irritation or redness develops and lasts

Apply onto dry hands. Work into a lather, rinse and dry thoroughly.

Store at room temperature

Water (Aqua), Lauramidopropylamine oxide, Glycerin, Cetrimonium Chloride, Sodium Chloride, Cocamide MIPA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Red 40 (CI 16035), Yellow 5 (CI 19140), Red 33 (CI 17200)

Helps eliminate bacteria on skin

NDC 51316-703-50



Antibacterial Liquid Hand Soap Refill

CLEAN SCENT

COMPARE TO
Softsoap® Crisp Clean
Antibacterial Liquid Hand Soap*

50 FL OZ (1.47 L)

06-26101

Drug Facts

Active ingredient	Purpose
Benzalkonium Chloride 0.13%	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 ■ 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 liquid hand soap pump bottle, apply onto wet hands.
■ lather and rinse thoroughly.

Other information store at room temperature.

Inactive ingredients Cetrimonium Chloride, Citric Acid, Cocamide MIPA, Fragrance (Parfum), Glycerin, Lauramidopropylamine Oxide, Methylchloroisoethiazolinone, Methylisothiazolinone, PEG-120 Methyl Glucose Dioleate, Red 33 (CI 17200), Red 40 (CI 16035), Sodium Chloride, Sodium Sulfate, Tetrasodium EDTA, Water (Aqua), Yellow 5 (CI 19140).

*This product is not manufactured or distributed by the Colgate-Palmolive Company, owner of the registered trademark Softsoap®.

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#262246



06-27040



NDC
51316-703-11

Drug Facts

Active ingredient	Purpose
Benzalkonium Chloride 0.13%	Antibacterial

Uses helps eliminate bacteria on hands.

Warnings For external use only.

When using this product avoid contact with eyes. If this occurs, 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 ■ apply onto wet hands.
■ lather and rinse thoroughly.

Other information store at room temperature.

Inactive ingredients Cetrimonium Chloride, Citric Acid, Cocamide MIPA, Fragrance (Parfum), Glycerin, Lauramidopropylamine Oxide, Methylchloroisothiazolinone, Methylisothiazolinone, PEG-120 Methyl Glucose Dioleate, Red 33 (CI 17200), Red 40 (CI 16035), Sodium Chloride, Sodium Sulfate, Tetrasodium EDTA, Water (Aqua), Yellow 5 (CI 19140).

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06-27039

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HANDSOAP

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51316-703
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
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

SODIUM SULFATE (UNII: 0YPR65R21J)
GLYCERIN (UNII: PDC6A3C0OX)
COCAMIDE MIPA (UNII: 21X4Y0VTB1)
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
FD&C RED NO. 40 (UNII: WZB9127XOA)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)
EDETATE SODIUM (UNII: MP1J8420LU)
D&C RED NO. 33 (UNII: 9DBA0SBB0L)
WATER (UNII: 059QF0KO0R)
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)
FRAGRANCE CLEAN ORC0600327 (UNII: 329LCV5BTF)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51316-703-56	1660 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2024	
2	NDC:51316-703-11	333 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2024	
3	NDC:51316-703-50	1470 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	04/01/2024	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	12/01/2021	

Labeler - CVS Pharmacy (062312574)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment

Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care		201901209	manufacture(51316-703)