

EQUATE ANTIBACTERIAL CITRUS HAND- benzalkonium chloride liquid
Wal-Mart Store 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.

Drug Facts

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

- irritation or redness develops and lasts.

Keep out of reach of children.

In case of accidental ingestion, get medical help or contact a Poison Control Center immediately.

Directions

- apply onto wet hands.
- lather and rinse thoroughly.

Other information

store at room temperature.

Inactive ingredients

Water (Aqua), Cetrimonium Chloride, Glycerin, Lauramidopropylamine Oxide, Sodium Chloride, Cocamide MEA, PEG-120 Methyl Glucose Dioleate, Fragrance (Parfum), Citric Acid, Tetrasodium EDTA, Sodium Sulfate, Methylchloroisothiazolinone, Methylisothiazolinone, Citrus Nobilis (Mandarin Orange) Peel Extract, Camellia Sinensis Leaf Extract, Zingiber Officinale (Ginger) Root Extract, Yellow 5 (CI 19140), Blue 1 (CI 42090).

Questions or comments?

1-888-287-1915

Label Copy



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DISTRIBUTED BY: Wal-Mart Stores, Inc., Bentonville, AR 72716
 PRODUCT OF CANADA
 *This product is not manufactured or distributed by Soft Soap Enterprises Inc., owner of registered trademark SOFT SOAP®.
 †vs. Equate 7.5 oz Liquid Hand Soap.

EQUATE ANTIBACTERIAL CITRUS HAND

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:49035-701
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0K00R)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	

EDETATE SODIUM (UNII: MP1J8420LU)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
TANGERINE PEEL (UNII: JU3D414057)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
GINGER (UNII: C5529G5JPQ)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:49035-701-11	333 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/05/2017	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	01/05/2017	

Labeler - Wal-Mart Store Inc (051957769)

Registrant - Apollo Health and Beauty Care (201901209)

Establishment			
Name	Address	ID/FEI	Business Operations
Apollo Health and Beauty Care		201901209	manufacture(49035-701)