

CVS PHARMACY CITRUS FRESH- benzalkonium chloride liquid

CVS PHARMACY

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. If contact occurs, rinse eyes 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

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

Label Copy



CVS PHARMACY CITRUS FRESH

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:59779-724
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
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)
TANGERINE PEEL (UNII: JU3D414057)
COCO MONOETHANOLAMIDE (UNII: C80684146D)
GLYCERIN (UNII: PDC6A3C0OX)
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SODIUM SULFATE (UNII: 0YPR65R21J)
EDETATE SODIUM (UNII: MP1J8420LU)
WATER (UNII: 059QF0KO0R)
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)
GINGER (UNII: C5529G5JPQ)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:59779-724-64	1890 mL in 1 BOTTLE, PLASTIC; 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	02/10/2016	

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(59779-724)