ANTIBACTERIAL SPRING RAIN- benzalkonium chloride liquid
Brands International Corporation

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.

Studio selection Antibacterial liquid hand soap- Orange Clementine

Active Ingredient - Benzalkonium Chloride - 0.13%

Purpose - Antibacterial

Use for handwashing to decrease bacteria on skin

for external use only - hands 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.

keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center immediately

Direction

* wet hands

* apply to hands

* scrub thoroughly

* rinse thoroughly

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Glycerin, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Vitamin E, Blue# 1, Yellow 5, Fragrance (Parfum)

orange clementine
ANTIBACTERIAL HAND SOAP

Drug Facts

Active ingredients — **Purpose**

Benzalkonium chloride 0.13% w/w ..... Antibacterial

Uses ■ for handwashing to decrease bacteria on the skin.

Warnings

For external use only; hands only

When using this product ■ do not get into eyes, if contact occurs, rinse eye thoroughly with water.

Stop use and ask a doctor if ■ irritation or redness develops.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ Wet hands ■ apply palmful to hands ■ scrub thoroughly ■ rinse

Inactive ingredients Water, Sodium laureth sulfate, Cocamidopropyl betaine, Sodium chloride, Citric acid, Fragrance, Polyquaternium-7, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, YELLOW 5, RED 4.

GO121

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MADE IN CANADA

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ANTIBACTERIAL SPRING RAIN

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:50157-605
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 1000 mL

Inactive Ingredients

Ingredient Name	Strength
EDETATE SODIUM (UNII: MP1J8420LU)	
ACID RED 337 (UNII: 9H247YW69C)	
POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600 KD) (UNII: 0L414VCS5Y)	
SOLVENT RED 4 (UNII: EVE9WNU99R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:50157-605-14	1480 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/14/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	11/14/2022	

Labeler - Brands International Corporation (243748238)

Establishment

Name	Address	ID/FEI	Business Operations
Brands International Corporation		243748238	manufacture(50157-605)

Revised: 11/2022 Brands International Corporation