

CLOROX ANTIMICROBIAL- benzalkonium chloride soap
Brand Buzz LLC

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.

Clorox Antibacterial Hand Soap (Orange Fusion)

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

- For handwashing to decrease bacteria on skin

Warnings

For external use only.

In case of accidental ingestion, drink a glass of water to dilute. If eye contact occurs, rinse thoroughly with water. If irritation persists, contact a Poison Control Center right away.

Keep out of reach of children.

Directions

- Apply onto wet hands, lather for 30 seconds and rinse thoroughly.

Other Information

- Avoid freezing and excessive heat above 40C (104F).

Inactive ingredients

Water, Sodium Laureth Sulfate, Sodium Chloride, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Laureth-3, Sodium Bicarbonate, Disodium EDTA, Fragrance, Methylchlorisothiazolinone, Methylisothiazoline, Yellow 6(Ci 15985), Red 7(Ci 16255).

Do not add bleach.

Not for use in dishwashers.

PHOSPATE-FREE

BLEACH-FREE

Distributed by: Brand Buzz LLC., 1407 Broadway, New York, NY 10018

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Questions or Comments?

Visit us at www.brandbuzzcp.com or call us at: 1-888-508-4750

For more product ingredient information, visit www.brandbuzzcp.com/ingredients

Clorox

Antimicrobial

HandSoap

Ultra Concentrated Dishwashing Liquid

Orange Fusion

with Oxi

Bleach-Free

8FL OZ

(0.5 PT) 236 mL

Label



Antibacterial
Hand Soap

Orange Fusion



Ultra Concentrated
Dishwashing Liquid

Bleach-Free

with
OXI

8 FL OZ
(0.5 PT) 236 mL



22FL OZ

(1.375 PT) 650 mL

The Clorox logo is a diamond shape with a red border and a yellow inner border. The word "CLOROX" is written in white, bold, sans-serif capital letters across the center of the diamond. A small "TM" trademark symbol is located at the bottom right of the diamond.

CLOROX™

Antibacterial
Hand Soap

Orange Fusion



Ultra Concentrated



CLOROX ANTIMICROBIAL

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69540-0030
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
C12-14 PARETH-12 (UNII: M0LJS773XW)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM BICARBONATE (UNII: 8MDF5V39QO)	

METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
WATER (UNII: 059QF0KO0R)	
PONCEAU 4R (UNII: Z525CBK9PG)	
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)	

Product Characteristics

Color		Score	
Shape		Size	
Flavor	ORANGE	Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69540-0030-1	236 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/21/2021	
2	NDC:69540-0030-2	650 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/21/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	04/07/2014	

Labeler - Brand Buzz LLC (079266204)