

ANTIBACTERIAL- anti bacterial solution
Collective Science Integration

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 ingredients	Purpose
Benzalkonium Chloride 0.25%	Antibacterial
Uses • for hand washing to decrease bacteria on the skin	
Warnings	
For external use only.	
When using this product avoid contact with eyes. In case of eye contact, flush with water.	
Stop use and ask a doctor if irritation and redness develops	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.	
Directions	
Lather vigorously and wash skin. Rinse and dry thoroughly.	
Inactive ingredients Water, TEA-Lauryl Sulfate, Cocamidopropyl Hydroxylamine, Glycerin, Sodium Chloride, Lavender, Peg-150 Distearate, Glycol Stearate Cocamide DIPA, Disodium EDTA, Mythyl Chloro Isothiazolinone, Methyl Isothiazolinone, CitricAcid.	

Dist. by P&P International Inc.
 El Monte, CA 91731
 Made in USA

www.csihealth.org



ABSP8BR28WH



**ANTI-BACTERIAL
 HAND SOAP**

KILLS 99% OF GERMS

**Lavander + Rosemary
 Essential Oils**

8 FL. OZ.



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ANTIBACTERIAL anti bacterial solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:90055-210
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Route of Administration TOPICAL

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.25 mg in 0.25 mg

Inactive Ingredients

Ingredient Name	Strength
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
LAVENDER OIL (UNII: ZBP1YXW0H8)	
TRIETHANOLAMINE LAURYL SULFATE (UNII: E8458C1KAA)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
CITRIC ACID ACETATE (UNII: DSO12WL7AU)	
GLYCERIN (UNII: PDC6A3C0OX)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
GLYCOL STEARATE (UNII: 0324G66D0E)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:90055-210-08	240 mg in 1 BOTTLE; Type 0: Not a Combination Product	10/09/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/09/2020	

Labeler - Collective Science Integration (133591508)

Establishment

Name	Address	ID/FEI	Business Operations
P&P International		133591508	manufacture(90055-210)

Revised: 2/2021

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