

ANTIBACTERIAL LAVENDER BOUQUET- benzalkonium chloride liquid
Universal Distribution Center LLC

ANTIBACTERIAL HAND SOAP LAVENDER BOUQUET

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

for handwashing 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 or redness develops • condition persists for more than 72 hours

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

Directions

• wet hands and apply soap • lather all surfaces of hands and fingers by rubbing vigorously for at least 20 seconds • rinse hands well and dry

Inactive ingredients

Water(Aqua), Sodium Laureth Sulfate, Cocamidopropyl Betaine, Cocamide Methyl MEA, Citric Acid, Methylisothiazolinone, Iodopropynyl Butylcarbamate, Fragrance, Sodium Chloride, D&C Red No. 33, FD&C Blue No. 1.

NEW

kill Germs and Odors

Distributed by:
Universal Distribution Center LLC
 330 Applegarth Road, Monroe Township, NJ08831
www.universaldc.com
 Made in China

Packaging



ANTIBACTERIAL LAVENDER BOUQUET

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52000-435
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
WATER (UNII: 059QF0KO0R)	

SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCOYL METHYL MONOETHANOLAMINE (UNII: 79G1T427CF)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

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Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-435-64	1893 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	01/31/2026	

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Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	01/31/2026	

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Labeler - Universal Distribution Center LLC (019180459)