ANTIBACTERIAL HANDWASH LAVENDER BOUQUET- benzalkonium chloride liquid
Universal Distribution Center LLC

ANTIBACTERIAL HANDWASH LAVENDER BOUQUET

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

USES

for hand washing to decrease bacteria on the skin.

Warning:

Avoid contact with eyes. In case of contact flush with water. Keep out of direct sunlight. Do not drink, not edible. External use only.

Keep out of reach of children.

Directions:

Pump a small amount of soap into wet hands, rub hands together to create rich lather, then rinse.

Ingredients:

Water, Sodium Laureth Sulfate, Sodium Chloride, Cocamidopropyl Betaine, Cocamide MIPA, Laureth-4, Glycerin*, Polyquaternium-7*, Fragrance, Methylchloroisothiazolinone, Methylisothiazolinone, Tetrasodium EDTA, Citric Acid, Benzophenono-4, Tocopheryl Acetate, FD&C Blue No. 1, D&C Red No. 33

*Contains one or more of these ingredients.

Kill Germs and Odors

Distributed By:
Universal Distribution Center
96 Distribution Boulevard,
Edison, NJ 08817

Made in Turkey

Packaging

ANTIBACTERIAL HANDWASH LAVENDER BOUQUET

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:52000-042
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)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
LAURETH-4 (UNII: 6HQ855798J)	
GLYCERIN (UNII: PDC6A3C0OX)	

POLYQUATERNIUM-7 (70/30 ACRYLAMIDE/DADMAC; 1600000 MW) (UNII: 0L414VCS5Y)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SULISOBENZONE (UNII: 1W6L629B4K)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:52000-042-01	400 mL in 1 BOTTLE; Type 0: Not a Combination Product	08/01/2018	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	08/01/2018	

Labeler - Universal Distribution Center LLC (019180459)

Registrant - Universal Distribution Center LLC (019180459)

Revised: 12/2024 Universal Distribution Center LLC