

DIAL ANTIMICROBIAL- benzalkonium chloride liquid
Bocchi Laboratories

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.

Dial Antimicrobial

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

- For handwashing to decrease bacteria on the skin

Warnings

For external use only.

Stop use and ask a doctor if

- 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

- Pump into dry hands.
- Lather vigorously for at least 15 seconds
- Rinse thoroughly and dry

Aqua (Water, Eau), Glycerin, Lauramine Oxide, Cetrimonium Chloride, Cocamidopropyl Betaine, Sodium Benzoate, Hydroxypropyl Methylcellulose, Parfum (Fragrance), Zinc Sulfate, Sodium Chloride, Dimethyl Lauramine, Tetrasodium EDTA, Alcohol, Dimethyl Myristamine, CI 42090 (Blue 1), CI 17200 (Red 33).

When using this product

Avoid contact with eyes, in case of eye contact, flush with water



DIAL ANTIMICROBIAL

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:57702-501
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	13 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
DIMETHYL LAURAMINE (UNII: 6V2OM301LZ)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	

GLYCERIN (UNII: PDC6A3C0OX)
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
WATER (UNII: 059QF0KO0R)
DIMETHYL MYRISTAMINE (UNII: 5E4O85D8T2)
SODIUM BENZOATE (UNII: OJ245FE5EU)
ALCOHOL (UNII: 3K9958V90M)
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)
D&C RED NO. 33 (UNII: 9DBA0SBB0L)
ZINC SULFATE (UNII: 89DS0H96TB)
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)
EDETATE SODIUM (UNII: MP1J8420LU)

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:57702-501-08	221 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	05/07/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	05/07/2021	

Labeler - Bocchi Laboratories (013579387)

Registrant - Henkel Corporation (081264055)

Establishment

Name	Address	ID/FEI	Business Operations
Bocchi Laboratories		013579387	manufacture(57702-501)

Revised: 5/2021

Bocchi Laboratories