

**SPATHERAPY - benzalkonium chloride lotion**  
**Xiamen Olivee Daily Use Chemical Co., Ltd.**

*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.*

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**Drug Facts**

Inactive Ingredients - Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Glycerin, DMDM Hydantoin, Benzalkonium Chloride, Fragrance, Xanthan Gum, Disodium EDTA, Citric Acid. May contain: CI 14700, CI 17200, CI 19140, CI 42090

Active Ingredient

Benzalkonium Chloride 0,2 %

Purpose

Antibacterial

Use - handwashing to decrease bacteria on skin

Directions - wet hands, work into rich lather - rinse well

Warnings

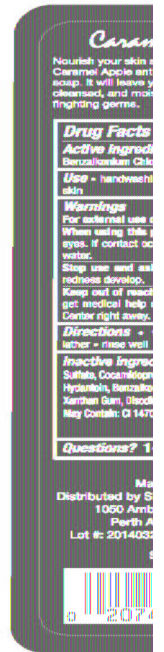
For external use only

When using this product, avoid contact with eyes. If contact occurs, rinse thoroughly with water.

Stop use and ask a doctor if irritation and redness develop

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

Questions? 1-800-FDA-1088



## SPATHERAPY

benzalkonium chloride lotion

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:56136-009
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	.56 g in 293 g

### Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH-3 SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	

GLYCERIN (UNII: PDC6A3C0OX)	
DMDM HYDANTOIN (UNII: BYR0546TOW)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:56136-009-01	293 g in 1 BOTTLE		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	03/27/2014	

**Labeler** - Xiamen Olivee Daily Use Chemical Co., Ltd. (526021375)

### Establishment

Name	Address	ID/FEI	Business Operations
Xiamen Olivee Daily Use Chemical Co., Ltd.		526021375	manufacture(56136-009)

Revised: 4/2014

Xiamen Olivee Daily Use Chemical Co., Ltd.