

EVERGREEN FRENCH LAVENDER SCENTED ANTIBACTERIAL HANDSOAP-

benzalkonium chloride liquid

**GENC GIDA KOZMETİK VE KİMYASAL URUNLER SANAYI VE TICARET LIMITED
SIRKETI**

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.

Evergreen FRENCH LAVENDER ANTIBACTERIAL HAND SOAP

Drug Facts

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

☐ **Uses** • ☐ 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.

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

Directions

- Pump into hands, wet as needed
- Lather vigorously for at least 15 seconds
- Wash skin, rinse thoroughly and dry

Inactive ingredients

Water, Sodium Laureth Sulfate, Cocamide MIPA, Sodium Chloride, Cocamidopropyl Betaine, Glycerin, Fragrance, Tetrasodium EDTA, Citric Acid, Methylchloroisothiazolinone, Methylisothiazolinone, Benzophenone-4, CI 42090 (Blue 1), CI 17200 (Red 33)

BEST WAY TO GET RID OF GERMS

Imported By:

Evergreen USA LLC

Hoboken, NJ 07030

www.egreenusa.com

PRODUCT OF TURKEY

Packaging



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**ANTIBACTERIAL
HAND SOAP**
**FRENCH
LAVENDER**

1 91414 01058 4

**17 FL OZ
(500 ml)**

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EVERGREEN FRENCH LAVENDER SCENTED ANTIBACTERIAL HANDSOAP

benzalkonium chloride liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:74600-504
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)	
COCO MONOISOPROPANOLAMIDE (UNII: 21X4Y0VTB1)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	

EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SULISOBENZONE (UNII: 1W6L629B4K)	
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:74600-504-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	04/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333E	04/01/2020	

Labeler - GENC GIDA KOZMETIK VE KIMYASAL URUNLER SANAYI VE TICARET LIMITED SIRKETI (356827969)

Establishment

Name	Address	ID/FEI	Business Operations
GENC GIDA KOZMETIK VE KIMYASAL URUNLER SANAYI VE TICARET LIMITED SIRKETI		356827969	manufacture(74600-504)

Revised: 8/2020

GENC GIDA KOZMETIK VE KIMYASAL URUNLER SANAYI VE TICARET LIMITED SIRKETI