

THE SPATHECARY FOAMING LAVENDER- benzalkonium chloride 0.13% soap
Larry (Xiamen) Hi Tech 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.

The Spathecary Foaming Hand Soap Lavender

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Uses

to help reduce bacteria that potentially can cause disease.

Warnings

For external use only.

When using this product

When using this product. Avoid contact with eyes. In case of eye contact, rinse eyes thoroughly with water.

Stop use and ask a doctor

if irritation or rash occurs. These may be signs of a serious condition.

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 20 seconds. Wash skin, rinse thoroughly, and dry.

Other Information

Store between 15-30C (59-86F)

Inactive Ingredients

aqua/water/EAU, Sodium Laureth Sulfate, cocamidopropylamine oxide, cocamidopropyl betaine, glycerin, parfum, methylchloroisothiazolinone, Methylisothiazolinone, Magnesium chloride, Magnesium nitrate, Aloe Barbadensis Leaf Juice, Disodium EDTA, Citric Acid, Tocopheryl Acetate, FD &C yellow No.5 (CI 19140), FD&C Red No. 33 (CI 17200), FD&C Blue No. 1 (CI 42090)

Questions or Comments?

P.O. Box 230258, Brooklyn, NY 11223

The Spathecary Foaming Hand Soap Lavender

benzalkonium chloride 0.13% soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73166-111
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.65 g in 500 mL

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	
MAGNESIUM CHLORIDE ANHYDROUS (UNII: 59XN63C8VM)	
ANHYDROUS CITRIC ACID (UNII: XF417D3PSL)	
GLYCERIN (UNII: PDC6A3C0OX)	
WATER (UNII: 059QF0K00R)	
.ALPHA.-TOCOPHEROL ACETATE (UNII: 9E8X80D2L0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
DISODIUM EDTA-COPPER (UNII: 6V475AX06U)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C BLUE NO. 1 (UNII: HBR47K3TBD)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73166-111-16	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/01/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/01/2020	

Labeler - Larry (Xiamen) Hi Tech Co., Ltd (529759328)

Registrant - G2 Beauty Inc (124608169)

