

LUCKY SUPER SOFT FOAMING- benzalkonium chloride soap
Delta Brands, Inc

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.

LckyHandSoapWldFlwrsBZK013

Active ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

for hand washing to decrease bacteria on the skin

Warnings

For external use only

When using this product

avoid contact with the eyes. In case of eye contact, flush with water.

Stop use and ask a doctor if

irritation and redness develops

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

■ wet hands and apply soap ■ lather all surfaces of hands and fingers by rubbing vigorously for at least 20 seconds ■ rinse hands well and dry

Inactive ingredients

water, glycerin, lauramine oxide, cetrimonium chloride, cocamidopropyl betaine, decyl

glucoside, hydroxypropyl methylcellulose, fragrance, zinc sulfate, citric acid, sodium chloride, tetrasodium EDTA, alcohol, methylchloroisothiazolinone, methylisothiazolinone

Package Label



LUCKY SUPER SOFT FOAMING

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:20276-138
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)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	

GLYCERIN (UNII: PDC6A3C00X)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
HYPROMELLOSE, UNSPECIFIED (UNII: 3NXW29V3WO)	
ZINC SULFATE (UNII: 89DS0H96TB)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
EDETATE SODIUM (UNII: MP1J8420LU)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
ALCOHOL (UNII: 3K9958V90M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:20276-138-75	222 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	01/29/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	01/29/2021	

Labeler - Delta Brands, Inc (102672008)

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