ANTIBACTERIAL- benzalkonium chloride 0.13% liquid
Flex Beauty Labs

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.

Antibacterial- Germout Hand Soap, Triclosan free, Kitchen Lemon

Purpose - Antibacterial

Use help eliminate bacteria on hands

Warnings

For external use only

Stop use and ask a doctor if irritation or redness develops

When using this product

Avoid contact with eyes.In case of contact, flush and rinse with water.

Keep out of reach of children

Except under adult supervision.

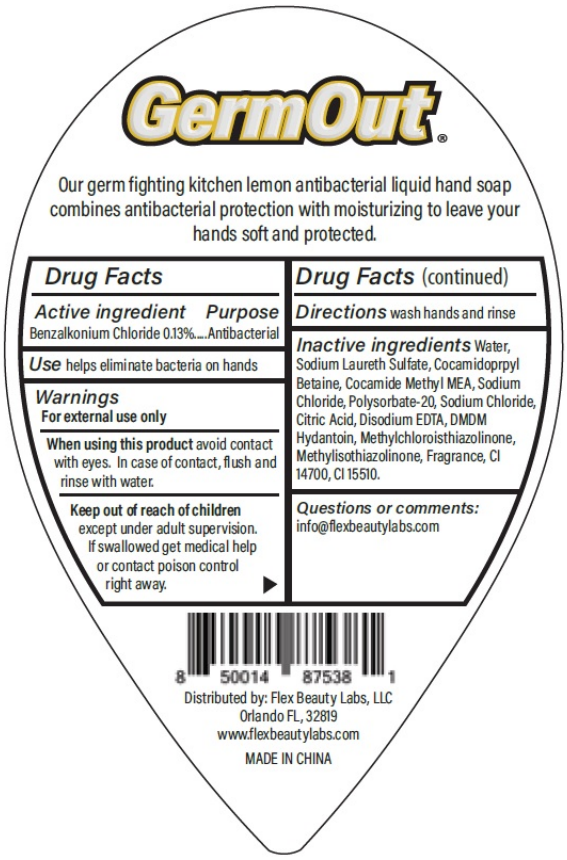
If swallowed get medical help or contact posion control right away.

Directions

Wash hands and rinse

Water, Sodium Laureth Sulfate, Cocamidopropyl Betaine, Sodium Chloride, Citric Acid, Coamide Methyl MEA, Fragrance, Disodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, DMDM Hydantoin ,Cl 14700, Cl 15510.

Benzalkonium Chloride - 0.13%



ANTIBACTERIAL

benzalkonium chloride 0.13% liquid

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:72308-013
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
COCO DIETHANOLAMIDE (UNII: 92005F972D)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

DMDM HYDANTOIN (UNII: BYR0546TOW)	
BENZYL ALCOHOL (UNII: LKG8494WBH)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
FD&C RED NO. 4 (UNII: X3W0AM1JLX)	
D&C ORANGE NO. 4 (UNII: Q1LIY3BO0U)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72308-013-01	325 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	07/10/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/10/2020	

Labeler - Flex Beauty Labs (080858917)

Revised: 7/2020 Flex Beauty Labs