PURELOGIC FOAMING HAND SANITIZER FRENCH LAVENDER- benzalkonium chloride solution
Argento sc by sicura 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.

AB0028 FL foaming

Active Ingredient(s)

Benzalkonium Chloride 0.13%

Purpose:

Antibacterial

Use

Hand sanitizer to help reduce bacteria on skin that could cause disease

Recommended for repeated use

Warnings

For external use on hands only.

When using this product

When using this product, keep out of eyes. In case of contact with eyes, flush thoroughly with water

Do not ingest or inhale

Avoid contact with broken skin

Stop use and ask a doctor

if irrigation and redness develop or if condition persists for more than 72 hours

Keep out of reach of children

if swallowed, get medical help or contact a Poison Control Center immediately

Directions

- Place enough product in your palm to thoroughly cover your hands. Rub hands together briskly until dry
- Children under 6 years of age should be supervised when using this product

Other information

To prevent discoloration, avoid contact with clothing

Inactive ingredients

Water, Sodium lauroamphoacetate, Cocamidopropylamine oxide, Cocamidopropyl betaine, Glycerin, Olea europaea (olive) leaf extract, Polyminopropyl biguanide, Citric acid, Disodium EDTA, Benzophenone-3, Methylchloroisothiazolinone, Methylisothiazolinone, Fragrance, Cetylpyridinium chloride, Red 33 (CI17200), Blue 1 (CI42090)

PURELOGIC FOAMING HAND SANITIZER FRENCH LAVENDER

benzalkonium chloride solution

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:77731-055
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
SODIUM LAUROAMPHOACETATE (UNII: SLK428451L)	
COCAMIDOPROPYLAMINE OXIDE (UNII: M4SL82J7HK)	
WATER (UNII: 059QF0KO0R)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
OLEA EUROPAEA LEAF (UNII: MJ95C3OH47)	
CETYLPYRIDINIUM CHLORIDE (UNII: D9OM4SK49P)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
DISODIUM HEDTA (UNII: KME849MC7A)	
BENZOPHENONE (UNII: 701M4TTV9O)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77731-055-01	500 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/31/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/31/2020	

Labeler - Argento sc by sicura inc. (168718778)

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