

PURELOGIC FOAMING HAND SANITIZER EUCALYPTUS MINT- 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 EM 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 irritation 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, Polyaminopropyl biguanide, Citric acid, Disodium EDTA, Benzophenone-3, Methylchloroisothiazolinone, Methylisothiazolinone, Fragrance, Cetylpyridinium chloride, Blue 1 (CI42090)



PURELOGIC FOAMING HAND SANITIZER EUCALYPTUS MINT			
benzalkonium chloride solution			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77731-056
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 LAURO AMPHO ACETATE (UNII: SLK428451L)			
CO CAMIDO PROPYLAMINE O XIDE (UNII: M4SL82J7HK)			
WATER (UNII: 059QF0KO0R)			
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)			
CO CAMIDO PROPYL BETAINE (UNII: 5OCF3O11KX)			
GLYCERIN (UNII: PDC6A3C0OX)			
OLEA EURO PAEA LEAF (UNII: MJ95C3OH47)			
CETYL PYRIDINIUM CHL O RIDE (UNII: D9OM4SK49P)			
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)			
DISODIUM HEDTA (UNII: KME849MC7A)			
BENZOPHENONE (UNII: 701M4TTV9O)			
METHYL CHLORO ISOTHIAZOLINONE (UNII: DEL7T5QRPN)			
METHYLISOTHIAZOLINO NE (UNII: 229D0E1QFA)			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77731-056-01	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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