# BROOKSTONE LEMON ZEST ADVANCED FOAMING HAND SANITIZER- 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.

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#### 77731-087/AB0054LZ-YEL foaming

#### **Active Ingredient(s)**

Benzalkonium Chloride 0.13%

Purpose:

Antibacterial

#### Use

Decreases bacteria on skin that could cause diseases

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**

- Spread product on hands thoroughly and allow to dry, do not wipe off
- For children under 6, use only under adult supervision
- Not recommended for use with infants

#### Other information

Do not store above 105F (40.6C)

May discolor some fabrics

Harmful to plastics and wood finished

#### **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, Yellow 5 (CI19140)



### **BROOKSTONE LEMON ZEST ADVANCED FOAMING HAND SANITIZER**

benzalkonium chloride solution

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77731-087		
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
SO DIUM LAURO AMPHO ACETATE (UNII: SLK428451L)	
CO CAMIDO PRO PYLAMINE O XIDE (UNII: M4SL82J7HK)	
WATER (UNII: 059QF0KO0R)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
COCAMIDO PRO PYL BETAINE (UNII: 50 CF3O 11KX)	
GLYCERIN (UNII: PDC6A3C0OX)	
OLEA EUROPAEA LEAF (UNII: MJ95C3OH47)	
CETYLPYRIDINIUM CHLO RIDE (UNII: D9 O M4S K49 P)	
CITRIC ACID MO NO HYDRATE (UNII: 2968 PHW8 QP)	
<b>DISO DIUM HEDTA</b> (UNII: KME849 MC7A)	
BENZOPHENO NE (UNII: 70 1M4TTV9 O)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229 D0 E1QFA)	

Packaging			
# Item Code	Package Description	<b>Marketing Start Date</b>	<b>Marketing End Date</b>

1 NDC:77731-087-01 5	00 mL in 1 BOTTLE; Type 0: Not a Combination Product	09/17/2020		
Marketing Information				
Marketing Category	y Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph not fina	al part333A	09/17/2020		

## Labeler - Argento sc by sicura inc. (168718778)

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