

PURELOGIC ANTIBACTERIAL WIPES REFRESHING ALOE 2 TRAVEL PACKS-

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.

WI0015AL

Active Ingredient(s)

Benzalkonium Chloride 0.2%

Purpose:

Disinfectant

USE

Decrease bacteria on skin that could cause disease

Recommended for repeated use

warnings

For external use only on hands. Flammable, keep way from flame or fire

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 and 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

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

other information

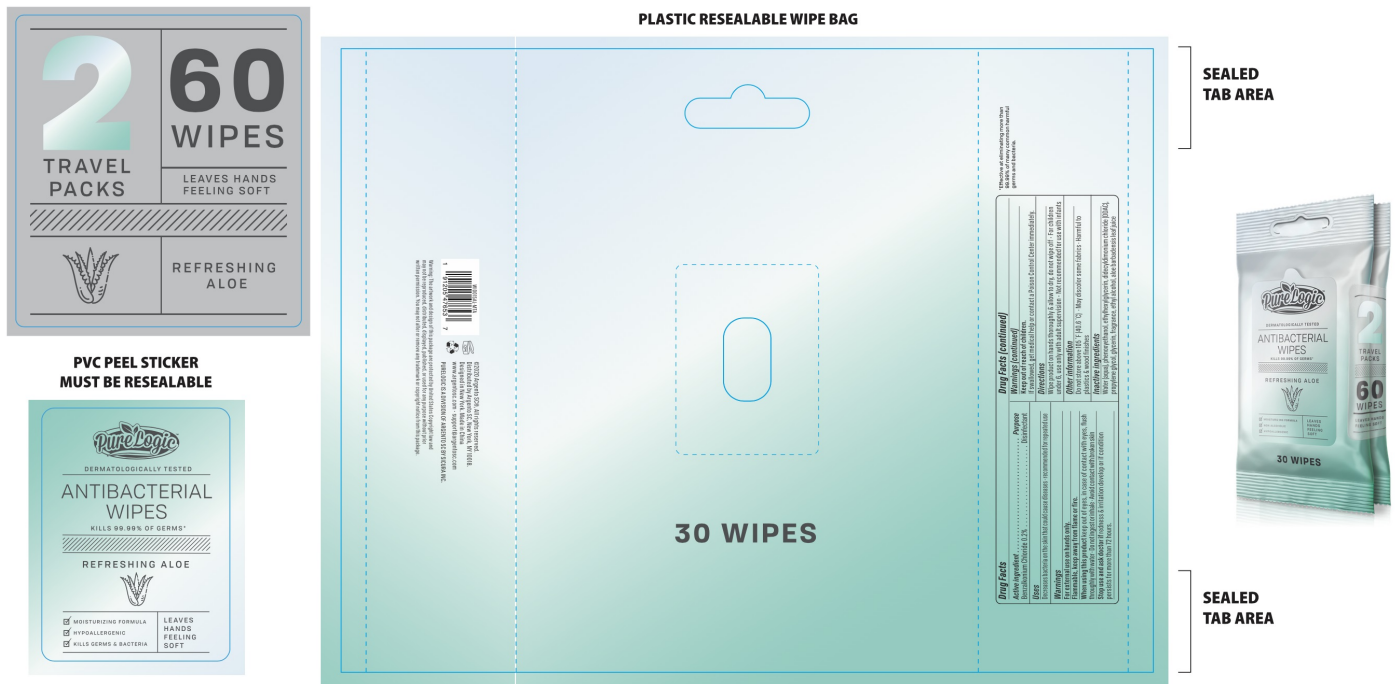
Do not store above 105F (40.6C)

May discolor some fabrics

Harmful to plastics and wood finishes

inactive ingredients

Purified water, Phenoxyethanol, Ethylhexylglycerin, Didecylmonium Chloride, Propylene Glycol, Glycerin, Fragrance, Aloe Barbadensis Leaf Juice



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Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:77731-012
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.006 g

Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0K00R)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:77731-012-01	2 in 1 PACKAGE	07/23/2020	
1		30 in 1 PACKET; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/23/2020	

Labeler - Argento sc by sicura inc. (168718778)

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