

**BIOPURE ANTIBACTERIAL HAND WIPE JUICY WATERMELON- benzalkonium chloride cloth  
Quest USA Corp**

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**Biopure Antibacterial Hand Wipe Juicy Watermelon**

**Drug Facts**

**Active ingredient**

Benzalkonium Chloride 0.13%

**Purpose**

Antibacterial

**Uses**

- Hand sanitizer to help reduce bacteria.
- For use when soap and water are not available.

**Warnings**

**For external use only.**

**Do not use**

- on children less than 2 months old.
- on open skin wounds.

**When using this product**

keep out of eyes, ears, and mouth. In case of contact with eyes, rinse eyes thoroughly with water.

**Stop use and ask a doctor**

if irritation or rash occurs. These may be signs of a serious condition.

**Keep out of reach of children.**

If swallowed, get medical help or contact a Poison Control Center right away.

**Directions**

- Apply to hands, allow to air dry without wiping.
- Children under 6 years of age should be supervised when using this product.

**Other Information**

- Store in a cool, dry place.
- Avoid freezing and excessive heat.

## Inactive ingredients

Aloe Barbadensis Leaf Extract, Alpha-Tocopherol, Fragrance, Glycerin, Phenoxyethanol, Propylene Glycol, Water

## Questions?

call 718-975-2586

## Anti-Bacterial Hand Wipe-Juicy Watermelon



## BIOPURE ANTIBACTERIAL HAND WIPE JUICY WATERMELON

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:78691-034
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

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

## Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
GLYCERIN (UNII: PDC6A3C00X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78691-034-00	50 mL in 1 CANISTER; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)	01/09/2026	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/09/2026	

**Labeler** - Quest USA Corp (079869689)

**Registrant** - Quest USA Corp (079869689)

Revised: 1/2026

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