

BIOPURE ANTI-BACTERIAL HAND WIPES FRESH MORNING BREEZE SCENT-
benzalkonium chloride cloth
Quest USA Corp

Biopure Anti-bacterial Hand Wipes Fresh Morning Breeze

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

- in 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

Package Labeling:

biopure.
**Anti-bacterial
Hand Wipes**
with Aloe and Vitamin E

0.13% BENZALKONIUM CHLORIDE
KILLS 99.9% OF GERMS
FRESH MORNING BREEZE

20 WIPES
2.75 IN X 5.91 IN
(7 cm X 15 cm)



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Drug Facts (continued)

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Directions

- Apply to hands, allow to air dry without wiping.
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Other Information

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

Drug Facts (continued)

Inactive Ingredients
Water, Propylene Glycol, Glycerin, Phenoxyethanol, Fragrance, Aloe Barbadensis Leaf Extract, Tocopheryl Acetate

Questions? Call 718-975-2586

DISTRIBUTED BY QUEST USA CORP.
485 FLATBUSH AVE, BROOKLYN NY 11225
MADE IN CHINA

BP2781-DT



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NOT INTENDED FOR MEDICAL USE.



LOT# 8004213
MFG: 01/2026
EXP: 01/2028

BIOPURE ANTI-BACTERIAL HAND WIPES FRESH MORNING BREEZE SCENT

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78691-032
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 g
Inactive Ingredients			
Ingredient Name			Strength
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)			

.ALPHA.-TOCOPHEROL ACETATE, DL- (UNII: WR1WPI7EW8)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78691-032-00	20 g 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

Quest USA Corp