

**BIOPURE ANTIBACTERIAL HAND WIPES BENZALKONIUM CHLORIDE 0.12% -  
250 WIPES- benzalkonium chloride cloth  
QUEST USA CORP.**

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**BioPure Antibacterial Hand Wipes Benzalkonium Chloride 0.12% - 250 Wipes**

***Drug Facts***

***Active ingredient***

Benzalkonium Chloride 0.12%

***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, between 15°-30° (59°-86°F).
- Avoid freezing and excessive heat above 40°C (104°F).

## Inactive ingredients

Glycerin, Phenoxyethanol, Propylene Glycol, Purified Water

## Questions?

call 718-975-2586

## Package Labeling:

DISTRIBUTED BY QUEST USA CORP.  
495 FLATBUSH AVE, BROOKLYN NY 11225

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MADE IN CHINA

BPBZK250CWM



Not intended for medical use

BioPure®

ANTIBACTERIAL

# Hand Wipes

BENZALKONIUM CHLORIDE 0.12%

KILLS 99.9% OF GERMS

DO NOT FLUSH

250 Wipes

5in.x7in.(12.7cmx17.8cm)

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## BIOPURE ANTIBACTERIAL HAND WIPES BENZALKONIUM CHLORIDE 0.12% - 250 WIPES

benzalkonium chloride cloth

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:78691-009
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZALKONIUM CHLORIDE</b> (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.2 mg in 1 mL

### Inactive Ingredients

Ingredient Name	Strength
<b>GLYCERIN</b> (UNII: PDC6A3C0OX)	
<b>PHENOXYETHANOL</b> (UNII: HIE492Z3T)	

**PROPYLENE GLYCOL** (UNII: 6DC9Q167V3)

**WATER** (UNII: 059QF0K00R)

### Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:78691-009-00	250 in 1 CONTAINER	09/20/2020	
1		2.8 mL in 1 PATCH; Type 2: Prefilled Drug Delivery Device/System (syringe, patch, etc.)		

### Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	09/20/2020	

**Labeler** - QUEST USA CORP. (079869689)

### Establishment

Name	Address	ID/FEI	Business Operations
Zhejiang Furuisen Spunlaced Nonwovens Co., Ltd.		723796509	manufacture(78691-009)

Revised: 12/2025

QUEST USA CORP.