

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

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



250 Wipes

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

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benzalkonium chloride cloth

Product Information

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

Active Ingredient/Active Moiety

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

Inactive Ingredients

Ingredient Name	Strength
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:78691-009-00	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)

Revised: 12/2023

QUEST USA CORP.