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

Biopure Anti-bacterial Hand Wipes Fresh Morning Breeze Scent

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:

Drug Facts (continued)	
Other information	● Store in a cool, dry place. ● Avoid freezing and excessive heat.
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Questions? call 718-975-2586	

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

MADE IN CHINA

BP2692-DT

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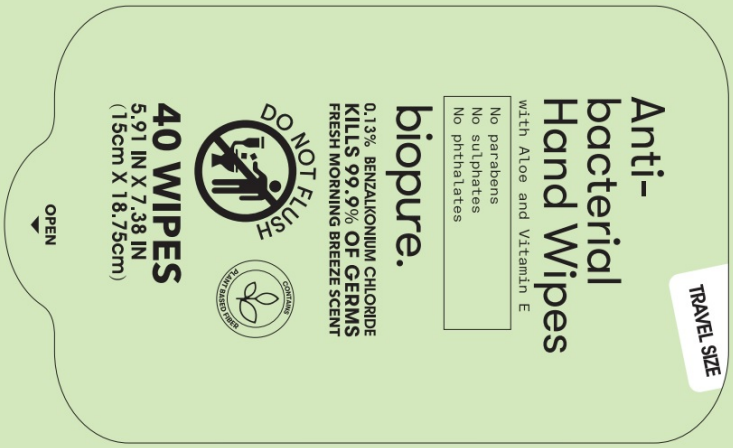
NOT INTENDED FOR MEDICAL USE

biopure. | Antibacterial Hand Wipes | 40



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See below for Drug Facts Box

BIOPURE ANTI-BACTERIAL HAND WIPES FRESH MORNING BREEZE SCENT

benzalkonium chloride liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78691-022
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (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: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	**Item Code**	**Package Description**	**Marketing Start Date**	**Marketing End Date**
1	NDC:78691-022-00	40 in 1 POUCH	08/07/2023	
1		3.9 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)	08/07/2023	

Labeler - Quest USA Corp (079869689)

Revised: 11/2023 Quest USA Corp