BIOPURE ANTI-BACTERIAL HAND WIPES HONEY WILDFLOWER SCENT-
benzalkonium chloride cloth
Quest USA Corp

Biopure Anti-bacterial Hand Wipes Honey Wildflower 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:

BIOPURE ANTI-BACTERIAL HAND WIPES HONEY WILDFLOWER SCENT

benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:78691-033
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:78691-033-00	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