BIOPURE ANTI-BACTERIAL HAND WIPES HONEY WILDFLOWER SCENTbenzalkonium chloride liquid 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

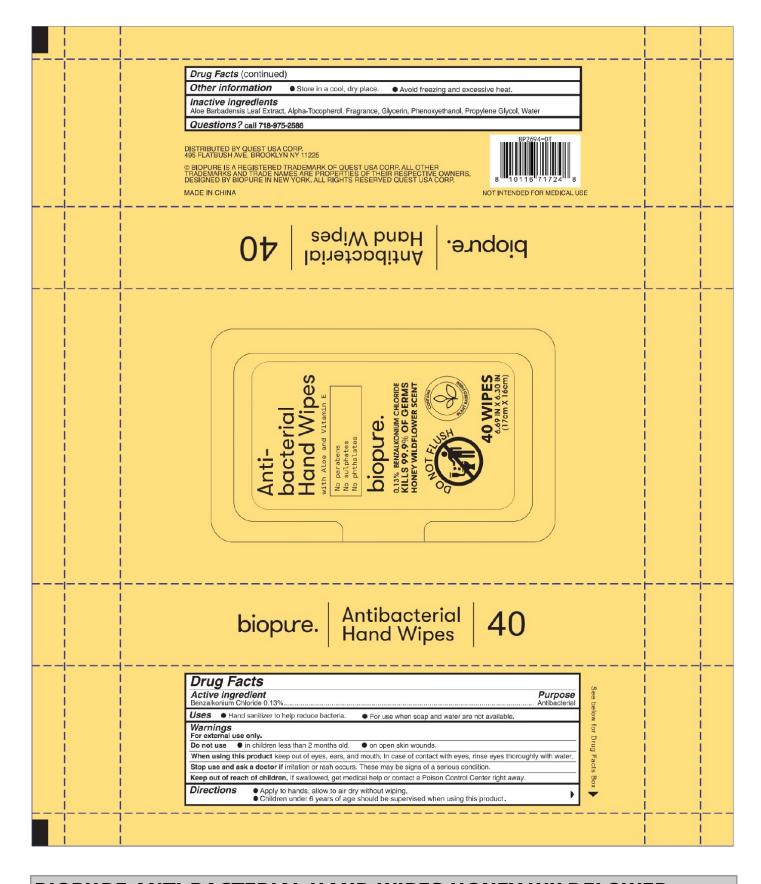
- Store in a cool, dry place.
- Avoid freezing and excessive heat.

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 liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:78691-023
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZ ALKONIUM CHLORIDE	1.3 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
ALOE VERA LEAF (UNII: ZY81Z83H0X)		
.ALPHATOCOPHEROL (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- 023-00	40 in 1 POUCH	08/07/2023		
1		3.8 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)	08/07/2023		

Labeler - Quest USA Corp (079869689)

Revised: 11/2023 Quest USA Corp