

BENZALKONIUM CHLORIDE- antibacterial hand wipes cloth
PreCare Corp.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Antibacterial Hand Wipes

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 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°C (59°F-86°F).
- Avoid freezing and excessive heat above 40°C (104°F).

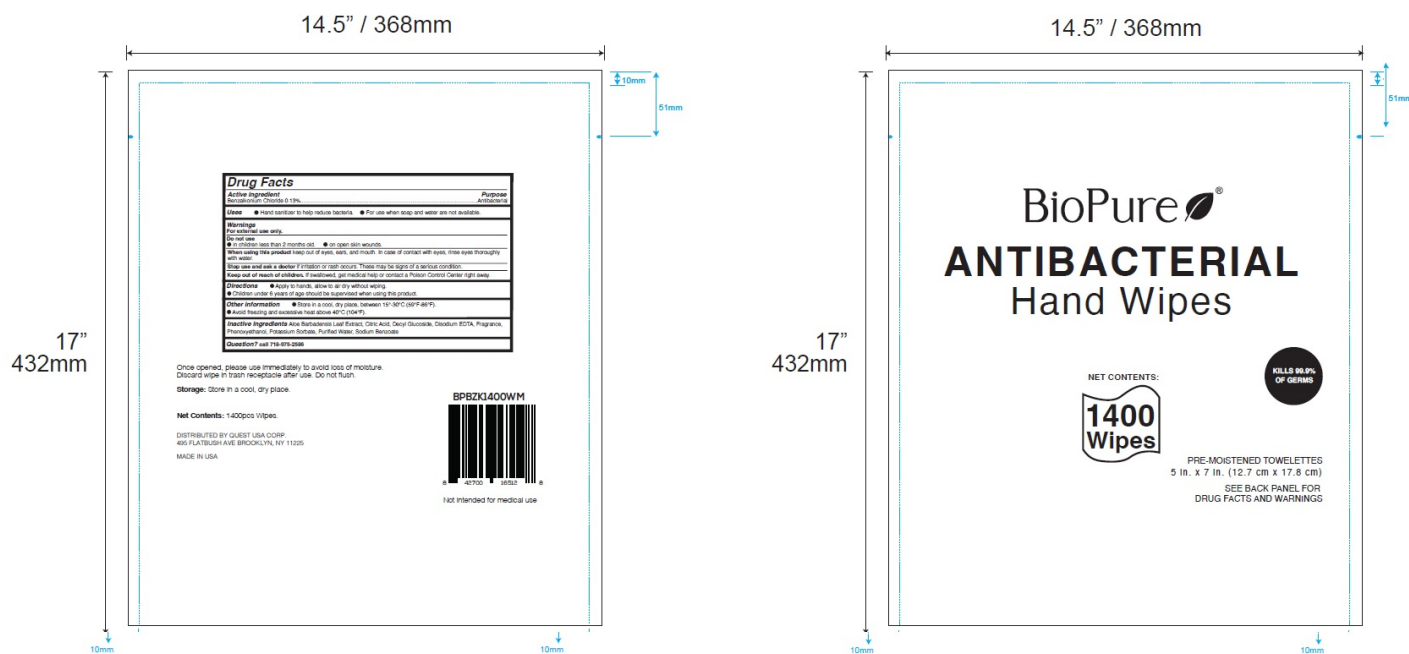
Inactive ingredients

Aloe Barbadensis Leaf Extract, Citric Acid, Decyl Glucoside, Disodium EDTA, Fragrance, Phenoxyethanol, Potassium Sorbate, Purified Water, Sodium Benzoate

Questions?

call 718-975-2586

Principal Display Panel



BENZALKONIUM CHLORIDE

antibacterial hand wipes cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:53118-140
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

Inactive Ingredients

Ingredient Name	Strength
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53118-140-04	1400 in 1 CONTAINER; Type 0: Not a Combination Product	02/14/2022	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	02/14/2022	

Labeler - PreCare Corp. (858442403)

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PreCare Corp.