

ANTIBACTERIAL HAND WIPE- antibacterial hand wipe cloth
Fujian Yifa Healthcare Products Co., Ltd.

BIOPURE Anti-Bacterial Hand Wipes-Island Vallina

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

Use

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

keep out of eyes,ears, and mouth.In case of contact with eyes,rinse eyes thoroughly with water.

Stop Use

If irritation or rash occurs.These may be signs of a serious condition.

Ask 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

1. Apply to hands, allow to air dry without wiping;
2. Children under 6 years of age should be supervised when using this product.

Other information

1. Store in a cool, dry place;
2. Avoid freezing and excessive heat.

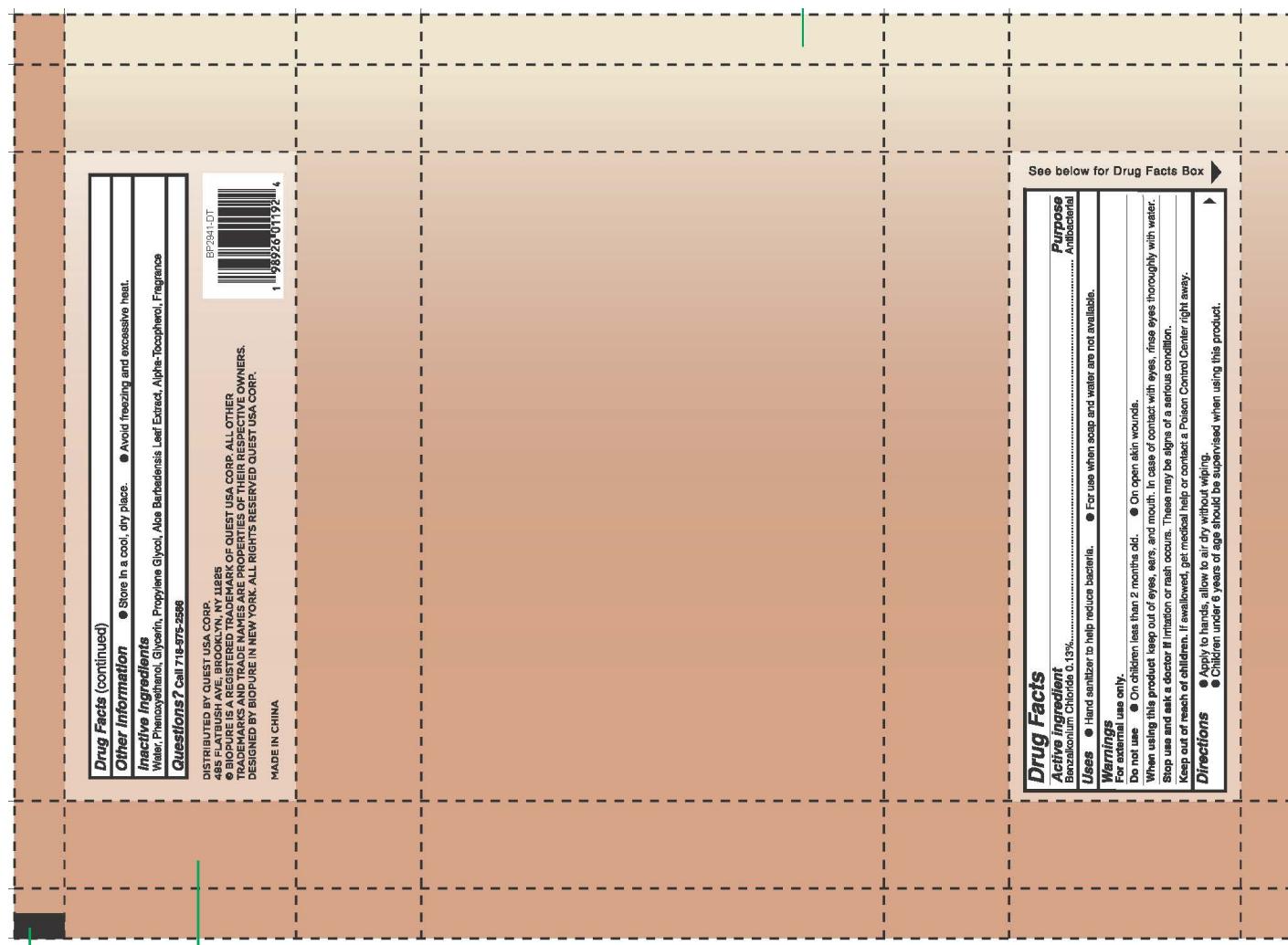
Inactive ingredients

Aloe Barbadensis Leaf Extract, Alpha-Tocopherol, Fragrance, Glycerin, Phenoxyethanol, Propylene Glycol, Water.

Questions

call 718-975-2586

PRINCIPAL DISPLAY PANEL-75543-306-00



biopure.

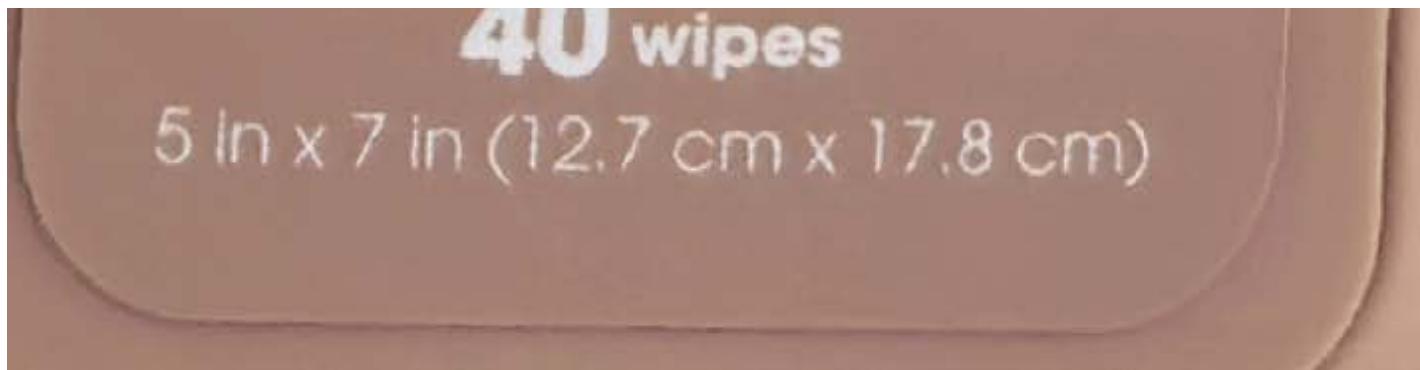
ANTI-BACTERIAL HAND WIPES

WITH ALOE AND VITAMIN E
0.13% BENZALKONIUM CHLORIDE
KILLS 99.9% OF GERMS

ISLAND VANILLA

NO PARABENS | NO SULPHATES | NO PHTHALATES





ANTIBACTERIAL HAND WIPE

antibacterial hand wipe cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75543-306
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.2 g in 40

Inactive Ingredients

Ingredient Name	Strength
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
ALOE BARBADENSIS LEAF (UNII: ZY81Z83H0X)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KOOR)	
GLYCERIN (UNII: PDC6A3C00X)	
ALPHA-TOCOPHEROL (UNII: H4N855PNZ1)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75543-306-00	40 in 1 PACKAGE; Type 0: Not a Combination Product	01/26/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M003	01/26/2026	

