

**UNIVERSITY MEDICAL PHARMACEUTICALS ANTIBACTERIAL HAND WIPES-
benzalkonium chloride cloth**

Jurong Dongfa General Merchandise Co Ltd

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.

**75127-003 University Medical Pharmaceuticals ANTIBACTERIAL HAND WIPES
0.13% Benzalkonium Chloride**

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antibacterial

USE

hand sanitizer to help reduce bacteria on the skin.

Warning

For external use only.

Do not use In or near the eyes In case of eye contact ,rinse eyes thoroughly with water.

Stop use and ask a doctor If Irritation or rash develops and persists more than 72 hours.

●Keep out of reach of children unless under adult supervision.

Directions

Wet hands thoroughly with wipe(s) and allow to air dry.

Discard after single use in trash bin. Do not flush.

To dispense peel back front label at tab enough to remove wipe(s) Press label firmly back In place to reseal pouch.

Inactive ingredients

Aqua, Glycerol, Aloe Barbadensis Leaf Juice, lanolin, Retinyl Palmitate, Allantoin,

Polysorbate-20, Disodium EDTA, Iodopropynyl Butylcarbamate, Methylisothiazolinone, Isopropyl Myristate.

Drug Facts (continued)

Directions

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Drug Facts (continued)

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DO NOT FLUSH



Drug Facts

Drug Facts (continued)

Active ingredient Benzalkonium Chloride 0.13%....	Purpose Antibacterial	Keep out of reach of children. If swallowed get medical help or contact a Poison Control Center right away.
Use ■ Hand sanitizer to help reduce bacteria on the skin		Do not use in or near the eyes. In case of eye contact, rinse eyes thoroughly with water.
Warnings For external use only. ▶		Stop use and ask a doctor if irritation or rash develops and persists more than 72 hours. ▼

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UNIVERSITY MEDICAL PHARMACEUTICALS ANTIBACTERIAL HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:75127-003
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
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
POLYSORBATE 20 (UNII: 7T1F30V5YH)	
LANOLIN (UNII: 7EV65EAW6H)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALLANTOIN (UNII: 344S277G0Z)	
VITAMIN A PALMITATE (UNII: 1D1K0N0VVC)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
GLYCERIN (UNII: PDC6A3C0OX)	
ISOPROPYL MYRISTATE (UNII: 0RE8K4LNJS)	
WATER (UNII: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:75127-003-01	200 in 1 CARTON	09/02/2020	
1		10 in 1 BAG		
1		45 g in 1 NOT APPLICABLE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	09/02/2020	

Labeler - Jurong Dongfa General Merchandise Co Ltd (528077612)**Establishment**

Name	Address	ID/FEI	Business Operations
Jurong Dongfa General Merchandise Co Ltd		528077612	manufacture(75127-003)