WET WIPE- benzalkonium chloride cloth
JIANGSU TERRA MEDICAL TECHNOLOGY 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.

WET WIPE

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Active Ingredient(s)

Benzalkonium chloride 1mg/pcs. Purpose: Antiseptic

Purpose

Antiseptic, WET WIPE

Use

WET WIPE to help reduce bacteria that potentially can cause disease. For use when soap and water are not available.

Warnings

For external use only. Flammable. Keep away from heat or flame

Do not use

- in children less than 2 months of age
- 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.

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Directions

- Wipe the surface of the skin and let it dry naturally
- Supervise children under 6 years of age when using this product to avoid swallowing.

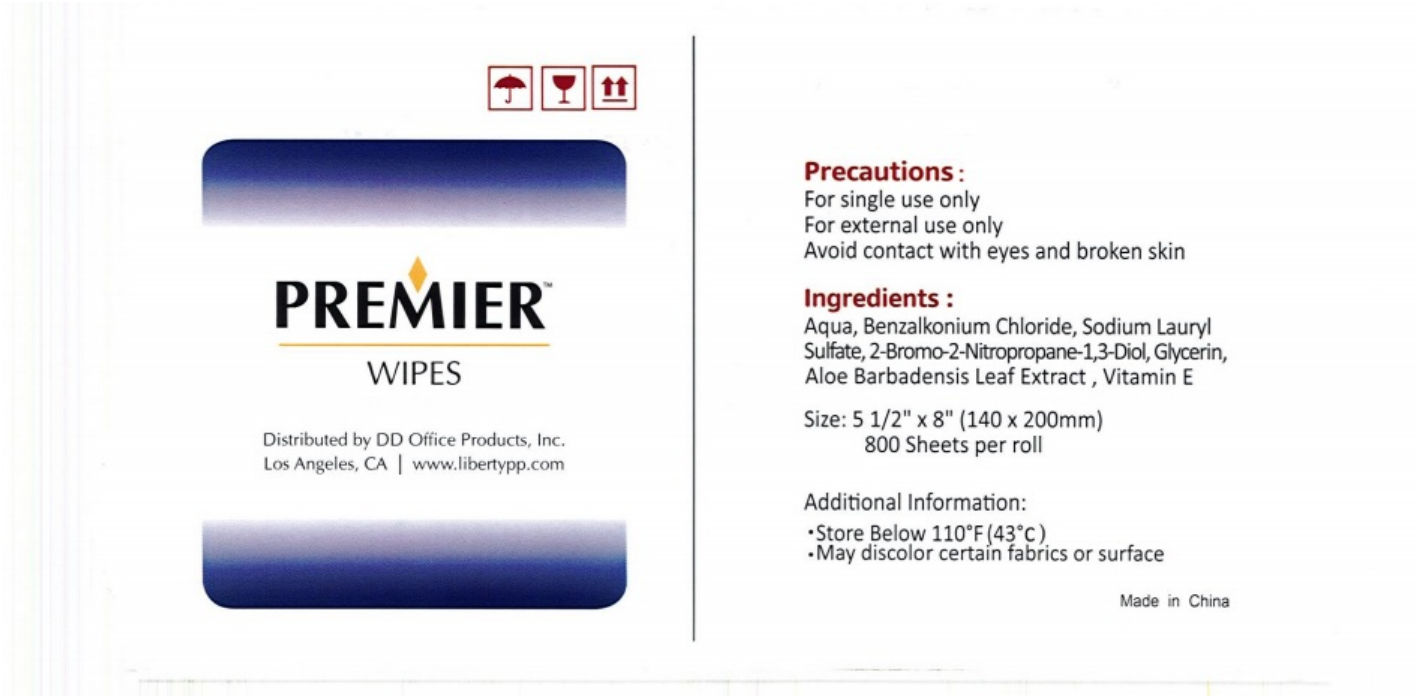
Other information

- Store between 15-30C (59-86F)
- Avoid freezing and excessive heat above 40C (104F)

Inactive ingredients

RO WATER
Sodium Lauryl Sulfate
2-Bromo-2-Nitropropane-1
Glycerin
Aloe Barbadensis Leaf Extract
Vitamin E

Package Label - Principal Display Panel



WET WIPE
benzalkonium chloride cloth

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69733-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1 mg

Inactive Ingredients	
Ingredient Name	**Strength**
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
BRONOPOL (UNII: 6PU1E16C9W)	
GLYCERIN (UNII: PDC6A3C0OX)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	

WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69733-101-01	800 in 1 BAG; Type 0: Not a Combination Product	03/30/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/30/2020	

Labeler - JIANGSU TERRA MEDICAL TECHNOLOGY CO LTD (421340473)

Establishment

Name	Address	ID/FEI	Business Operations
JIANGSU TERRA MEDICAL TECHNOLOGY CO LTD		421340473	manufacture(69733-101)

Revised: 8/2020 JIANGSU TERRA MEDICAL TECHNOLOGY CO LTD