

SANITIZING WET WIPES- alkyl dimethyl benzyl ammonium chloride cloth
Jiangsu Xiaolikang Medical Technology Co., Ltd.

Disclaimer: This drug has not been found by FDA to be safe and effective, and this labeling has not been approved by FDA. For further information about unapproved drugs, [click here](#).

Sanitizing Wet Wipes

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Active Ingredient(s)

Alkyl dimethyl benzyl ammonium chloride, 0.224mg/pcs

Dioctyl dimethyl ammonium chloride, 0.168mg/pcs

Didecyl dimethyl ammonium chloride, 0.07mg/pcs

Octyl decyl dimethyl ammonium chloride, 0.1mg/pcs

Purpose

Antiseptic, Sanitizing Wet Wipes

Use

Sanitizing Wet Wipes 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 air dry
- 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

Glycerin
 2-Bromo-2-Nitropropane-1
 IPBC
 PHMB(Polyaminopropyl Biguanide)
 Aloe Barbadensis Leaf Extract
 Vitamin E

Package Label - Principal Display Panel



SANITIZING WET WIPES

alkyl dimethyl benzyl ammonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:80337-201
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
QUATERNIUM-24 (UNII: 0T2NG1539G) (QUATERNIUM-24 - UNII:0T2NG1539G)	QUATERNIUM-24	0.1 mg
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.224 mg
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B) (DIDECYLDIMONIUM - UNII:Z7F472XQPA)	DIDECYLDIMONIUM CHLORIDE	0.07 mg

Inactive Ingredients

Ingredient Name	Strength
IODOPROPYNYL BUTYLCARBAMATE (UNII: 603P14DHEB)	
GLYCERIN (UNII: PDC6A3C0OX)	
DIOCTYLDIMONIUM CHLORIDE (UNII: 0X0RL40Y6H)	0.168 mg
POLYAMINOPROPYL BIGUANIDE (UNII: DT9D8Z79ET)	
BRONOPOL (UNII: 6PU1E16C9W)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
TOCOPHEROL (UNII: R0ZB2556P8)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:80337-201-01	80 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
unapproved drug other		03/30/2020	

Labeler - Jiangsu Xiaolikang Medical Technology Co., Ltd. (554546867)**Establishment**

Name	Address	ID/FEI	Business Operations
Jiangsu Xiaolikang Medical Technology Co., Ltd.		554546867	manufacture(80337-201)

Revised: 9/2020

Jiangsu Xiaolikang Medical Technology Co., Ltd.