

ANTIBACTERIAL HAND WIPES- benzalkonium chloride cloth
Karza Med Group, LLC

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.

Anti Bacterial Soap





Add Water to Moisten Cloth. Rub Cloth to build Later. Air or Towel Dry

Inactive Ingredients: Cocmethyl Propyl Betain, Sodium Lauryl Ether Sulfate, Postassium Sorbate, Sodium Chloride, 1,2 Propendiol, Citric Acid, Sodium Myristate, Sodium Dodcyl Sulfate, Fatty Esther

Ethoxylates, Vitamin E, Aloe Vera.



For Hand Sanitizing and Decreasing Bacteria on Hands. If irritation occurs stop usage.

REQUIRED

DRUG FACTS:
Active Ingredient: 0.2% Benzalkonium chloride
Purpose: Antibacterial. Uses: For hand sanitizing to decrease bacteria on skin. Safe for repeated use.

WARNINGS: For external use only. Avoid Contact with Eyes. If in eyes, flush thoroughly with water. If irritations occur, stop use.


Inactive Ingredients: Cocamidopropyl Betaine, Sodium Lauryl Ether Sulfate, Potassium Sorbate, Sodium Chloride, 1,2 Propanediol, Citric Acid, Sodium Myristate, Sodium Laurate, Sodium Benzoate, Sodium Dodecyl Sulfate, Fatty Ester Ethoxylates, Vitamin E, Aloe Vera (Organic).

Cloth: 100% Polyester.


In Case of an Emergency: 1-800-424-9300 (Chemtec)

Manufactured in China
ISO 9001 accredited facility
Style #TS200

Distributed by:
Karza Med Group, LLC
New York, NY 10001
Phone: 845-344-1111



Air dry or wipe if needed



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Keep out of Reach of Children.

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ANTIBACTERIAL HAND WIPES

benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76749-020
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.02 g in 0.02 g

Inactive Ingredients

Ingredient Name	Strength
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
.ALPHA.-TOCOPHEROL (UNII: H4N855PNZ1)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
SODIUM MYRISTATE (UNII: 06BLC4V0IV)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM LAURATE (UNII: K146MR5EXO)	
SODIUM LAURETH SULFATE (UNII: BPV390UAP0)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	

Product Characteristics

Color		Score	
Shape	SQUARE (10 cm x 10 cm Cloth)	Size	
Flavor		Imprint Code	
Contains			

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76749-020-02	24 g in 1 BAG; Type 0: Not a Combination Product	07/15/2020	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	07/15/2020	

Labeler - Karza Med Group, LLC (117470622)

Establishment

Name	Address	ID/FEI	Business Operations
NINGBO RIWAY NONWOVENS TECH CO., LTD9		554165780	manufacture(76749-020)

Revised: 5/2020

Karza Med Group, LLC