

STERIZAR ADVANCED BARRIER CONTROL- benzalkonium chloride cloth
Unicorn Media Partners 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.

STERiZAR®
Advanced Barrier Control

Drug Facts

Active Ingredient(s)

Benzalkonium Chloride 1.6 %

Purpose

Antiseptic

Use

Antibacterial Hand Wipes for Rapid Kill of Bacteria on Skin. For Use When Soap and Water Not Available.

Warnings

For external use only. Keep away from heat. Flammable, Keep Away from Heat and Flame

Do not use

- *On open skin wounds*

When using this product keep out of eyes 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 sign of serious medical conditions.

Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away at 1-800-222-1222

Directions

- *Wipe hands for 30 seconds then leave to air dry.*
- *Always reseal wipe container after use.*

Other information

- *Store in cool dry area.*
- *Avoid freezing*
- *Store at 15-30 C (59-86F) Avoid freezing and excessive heat above 40C (104F)*

Inactive ingredients

Amines, C12-14 Alkyl dimethyl, Decyl Dimethyl Ammonium Chloride, N-oxides, Purified Water, Sodium Ethyl Hexyl Sulfate.

Imported and distributed by
Unicorn Media Partners LLC
41 E. Sunrise Hwy.
Lindenhurst, NY 11757

PRINCIPAL DISPLAY PANEL - 200 Wipe Bag Label

NDC 79310-305-26

STERiZAR®

Advanced Barrier Control

Antibacterial Hand Wipes

Specially formulated to kill
harmful bacteria in seconds

ALCOHOL
FREE

Kills 99.999% of germs

Ideal for killing bacteria on skin.
No need to rinse

Contents: 200 average

• HALAL MONITORING •
COMMITTEE - UK

NDC 79310-305-26

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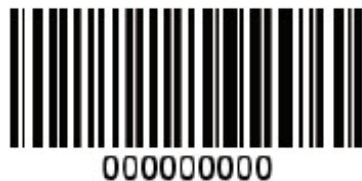
Contents: 200 average





Antibacterial Hand Wipes

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Batch No.:
Use by

Manufactured in the United Kingdom
Imported and distributed by
Unicorn Media Partners LLC
41 E. Sunrise Hwy.
Lindenhurst, NY 11757
Telephone: +1 631-496-3817
www.sterizarusa.com



benzalkonium chloride cloth

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79310-305
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
Benzalkonium Chloride (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	Benzalkonium Chloride	16 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
DIDECYLDIMONIUM CHLORIDE (UNII: JXN40O9Y9B)	
SODIUM ETHASULFATE (UNII: 12838560LI)	
WATER (UNII: 059QF0KO0R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79310-305-26	200 in 1 BAG	11/01/2020	
1		2.35 mL in 1 APPLICATOR; 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	11/01/2020	

Labeler - Unicorn Media Partners LLC (117525623)

Revised: 1/2022

Unicorn Media Partners LLC