

STERIZAR ADVANCED BARRIER CONTROL- benzalkonium chloride spray
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(s)

Hand Sanitizer Foam. For Hand Sanitizing to Decrease Bacteria On Skin. Safe for Repeated Use. 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

- *Squeeze gently approximately 2 ml (one shot) on physically clean hands.*
- *Rub into the hands skin for at least 30 second ensuring that all areas are covered until dry.*

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 - 100 mL Bottle Label

NDC 79310-304-23

STERiZAR®

Advanced Barrier Control

Hand Foam

Sanitizer

Specially formulated to kill
harmful bacteria in seconds

ALCOHOL

FREE

Kills 99.999% of germs

Remains effective after application for at
least 6 hours using Advanced Barrier Control.
No need to rinse. Moisturizing foam that
is kind to your hands.

3.4 FL OZ (100 mL)

• HALAL MONITORING •
COMMITTEE - UK

NDC 79310-304-23

STERiZAR[®]

Advanced Barrier Control



Hand Foam Sanitizer

Specially formulated to kill
harmful bacteria in seconds

**ALCOHOL
FREE**
Kills 99.999% of germs

Remains effective after application for at

least 6 hours using Advanced Barrier Control.
No need to rinse. Moisturizing foam that
is kind to your hands.

3.4 FL OZ (100 mL)



Hand Foam Sanitizer

Drug Facts	
Active Ingredient(s) Benzalkonium Chloride: 1.6%	Purpose Antiseptic
Use(s) Hand Sanitizer Foam. For Hand Sanitizing to Decrease Bacteria On Skin. Safe for Repeated Use. 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 signs 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: • Squeeze gently approximately 2 ml (one shot) on physically clean hands. • Rub into the hands skin for at least 30 seconds ensuring that all areas are covered until dry.	
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.	



000000000

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 spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79310-304
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: 059QF0K00R)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79310-304-25	50 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/01/2020	
2	NDC:79310-304-23	100 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/01/2020	
3	NDC:79310-304-22	500 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/01/2020	
4	NDC:79310-304-21	5000 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	11/01/2020	

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: 11/2020

Unicorn Media Partners LLC