

MARKET AMERICA ANTIBACTERIAL HAND SANITIZER- benzalkonium chloride spray
Market America, Inc.

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.

ANTIBACTERIAL HAND SANITIZER

Drug Facts

Active Ingredient

Benzalkonium Chloride 0.13%

Purpose

Antimicrobial

Use

- For hand sanitizing to decrease bacteria on the skin.

Warnings

For external use only

When using this product avoid contact with eyes. In case of contact with eyes, flush eyes with water.

Stop use and ask a doctor if irritation or rash develops and lasts.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Other Information

- May discolor certain fabrics or surfaces

Directions

- wet hands thoroughly with product and rub hands together until dry.
- children under 6 years of age should be supervised when using this product
- Not recommended for infants

Inactive ingredients

Ionized water, Urea

CONVERT SPENDING INTO EARNING

alcohol-free

triclosan-free &

fragrance free

Eliminates 99.99% of Germs

SHOPPING ANNUITY® BRAND

BAC-D

BACTERIAL DEFENSE

Manufactured for Market America, Inc.,
1302 Pleasant Ridge Rd., Greensboro, NC 27409

Packaging

alcohol-free
triclosan-free &
fragrance-free

CONVERT SPENDING
INTO EARNING

**ANTIBACTERIAL
HAND
SANITIZER**

Eliminates 99.99% of Germs

3.0 FL OZ (88.7ml)

BAC-D
BACTERIAL DEFENSE

310C01860902

SHOPPING ANNUITY® BRAND

Manufactured for Market America, Inc.,
1302 Pleasant Ridge Rd., Greensboro, NC 27409

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benzalkonium chloride spray

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:76209-117
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
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BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.3 mg in 1 mL
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Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
UREA (UNII: 8W8T17847W)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76209-117-03	88.7 mL in 1 BOTTLE, SPRAY; Type 0: Not a Combination Product	03/11/2021	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	03/11/2021	

Labeler - Market America, Inc. (797412236)

Registrant - Cemi International, Inc (015038336)

Establishment

Name	Address	ID/FEI	Business Operations
Cemi International, Inc		015038336	manufacture(76209-117)

Revised: 3/2021

Market America, Inc.