SHOPPING ANNUITY PREMIUM ANTI-BACTERIAL HANDSOAP- benzalkonium chloride liquid 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.

SHOPPING ANNUITY® BRAND PREMIUM ANTIBACTERIAL HAND SOAP

Drug Facts

Active ingredient

Benzalkonium chloride 0.13%

Purpose

Antimicrobial

Use

For hand-washing 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.

Directions

- Pump into hands; wet as needed.
- Lather vigorously for at least 20 seconds.
- Rinse and dry thoroughly.

Inactive ingredients

Water, Lauramine Oxide, Glycerin, Cocamidopropyl Betaine, Cetrimonium Chloride, PEG-120 Methyl Glucose Dioleate, PEG/PPG-120/10 Trimethylolpropane Trioleate, Laureth-2,

Cocamide MEA, Citric Acid, Aloe Barbadensis Leaf Extract, Allantoin, Disodium EDTA, Sodium Chloride, Tetrasodium EDTA, Methylchloroisothiazolinone, Methylisothiazolinone, Magnesium Chloride, Magnesium Nitrate.

CONVERT SPENDING INTO EARNING

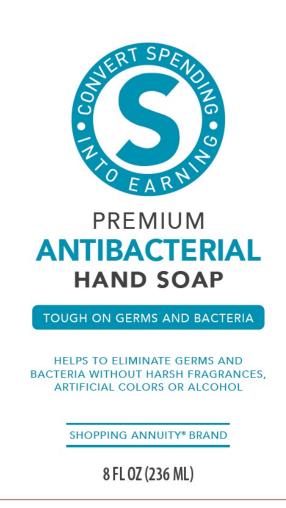
TOUGH ON GERMS AND BACTERIA

HELPS TO ELIMINATE GERMS AND BACTERIA WITHOUT HARSH FRAGRANCES, ARTIFICIAL COLORS OR ALCOHOL

Manufactured for and distributed by: Market America, Inc., 1302 Pleasant Ridge Road, Greensboro, NC 27409, USA

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Packaging







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SHOPPING ANNUITY PREMIUM ANTI-BACTERIAL HANDSOAP

benzalkonium chloride liquid

Product	Inform	ation
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Product Type HUMAN OTC DRUG Item Code (Source) NDC:76209-118

Route of Administration TOPICAL

l	Active Ingredient/Active Moiety			
l	Ingredient Name	Basis of Strength	Strength	
	BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	0.13 g in 100 mL	

Inactive Ingredients	
Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
LAURAMINE OXIDE (UNII: 4F6FC4MI8W)	
GLYCERIN (UNII: PDC6A3C0OX)	
COCAMIDOPROPYL BETAINE (UNII: 50CF3011KX)	
CETRIMONIUM CHLORIDE (UNII: UC9PE95IBP)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
TRIMETHYLOLPROPANE TRIOLEATE (UNII: S448BZW2BR)	
LAURETH-2 (UNII: D4D38LT1L5)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
ALLANTOIN (UNII: 344S277G0Z)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
EDETATE SODIUM (UNII: MP1J8420LU)	
METHYLCHLOROISOTHIAZOLINONE (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
MAGNESIUM CHLORIDE (UNII: 02F3473H9O)	
MAGNESIUM NITRATE (UNII: 77CBG3UN78)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:76209- 118-08	236 mL in 1 BOTTLE, PUMP; 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	nart2221	02/11/2021	

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Labeler - Market America, Inc. (797412236)

Registrant - Cemi International, Inc (015038336)

Establishment			
Name	Address	ID/FEI	Business Operations
Cemi International, Inc		015038336	manufacture(76209-118)

Revised: 3/2021 Market America, Inc.