ANTIBACTERIAL BODY WASH- benzalkonium chloride lotion
Vi-Jon

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.

Active ingredients

Benzalkonium chloride 0.13%

Purpose

Antibacterial

Use

to decrease bacteria on the skin

Warnings

For external use only

When using this product

- do not get into eyes. If contact occurs, rinse eyes thoroughly with water.

Keep out of reach of children

If swallowed, get medical help or contact a Poison Control Center right away.

Directions

- work a small amount into a lather
- scrub thoroughly
- rinse

Inactive ingredients

water, PEG-120 methyl glucose dioleate, cetrimonium chloride, glycerin, lauramidopropylamine oxide, cocamide MEA, fragrance, sodium sulfate, myristamidopropylamine oxide, sodioum chloride, citric acid, tetrasodium EDTA, methylchloroisothiazolinone, methylisothiazolinone, blue 1, red 33

principal display panel

Antibacterial Body Wash

21 FL OZ (621 mL)

Image not available.
Manufactured exclusively for private label distribution

ANTIBACTERIAL BODY WASH

benzalkonium chloride lotion

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:11344-337
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII:7N6JUD5X6Y)	BENZALKONIUM CHLORIDE	1.313 mg in 1 mL

Inactive Ingredients	
Ingredient Name	**Strength**
water (UNII: 059QF0KO0R)	
PEG-120 METHYL GLUCOSE DIOLEATE (UNII: YM0K64F20V)	
cetrimonium chloride (UNII: UC9PE95IBP)	
GLYCERIN (UNII: PDC6A3C0OX)	
LAURAMIDOPROPYLAMINE OXIDE (UNII: I6KX160QTV)	
COCO MONOETHANOLAMIDE (UNII: C80684146D)	
SODIUM SULFATE (UNII: 0YPR65R21J)	
MYRISTAMIDOPROPYLAMINE OXIDE (UNII: 3HSF539C9T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
EDETATE SODIUM (UNII: MP1J8420LU)	
methylchloroisothiazolinone (UNII: DEL7T5QRPN)	
METHYLISOTHIAZOLINONE (UNII: 229D0E1QFA)	
FD&C BLUE NO. 1 (UNII: H3R47K3TBD)	

D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:11344-337-56	621 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	10/15/2014	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph not final	part333A	10/15/2014	

Labeler - Vi-Jon (150931459)

Registrant - Vi-Jon (088520668)

Establishment

Name	Address	ID/FEI	Business Operations
Vi-Jon		088520668	manufacture(11344-337)

Revised: 7/2020 Vi-Jon