ANTIBACTERIAL GOLD- benzalkonium chloride soap Kareway Product, 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.

Pure-Aid Antibacterial Soap Gold

Active Ingredient

Benzalkonium Chloride 0.1%

Purpose

Antibacterial

Use

For washing to decrease bacteria on the skin

Warnings

For external use only

- Avoid contact with eyes. In case of eye contact, flush with water.
- Stop use and ask a doctor if irritation and redness develops.

Keep out of reach of children.

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

Directions

- Wet bar with water
- Lather vigorously and wash skin
- Rinse and dry thoroghly

Inactive Ingredient

butylated hydroxytoluene, etidronate tetrasodium, fragrance, glycerine, orange HO, PEG-6 methyl ether, pentasodium pentetate, sodium chloride, sodium laureth-5 sulfate, sodium palm kernelate, sodium palmate, sorbitol, titanium dioxide, water

Package label











Drug Facts	Drug Facts(continued)
Active Ingredient Purpose Benzalkonium	Inactive Ingredients butylated hydroxytoluene, etidronate tetrasodium, fragrance, glycerine,
OSE FOR Washing to decrease bacterial on the skin	orange HO, PEG-6 methyl ether, pentasodium pentetate, sodium chloride,
Warnings For external use only	sodium laureth-5 sulfate, sodium palm kernelate, sodium palmate, sorbitol, titanium dioxide, water
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eye contact, flush with water.

Stop use and ask a doctor if irritation and redness develops.

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

Directions = Wet bar with water

Lather vigorously and wash skin

Rinse and dry thoroughly

KARE WAY

Exclusively distributed by: Kareway Product Inc. 2550 S. Dominguez Hills Dr. Compton, CA 90220 Made in Indonesia



ANTIBACTERIAL GOLD

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:67510-0116
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZALKO NIUM CHLO RIDE (UNII: F5UM2KM3W7) (BENZALKO NIUM - UNII:7N6 JUD5X6 Y)	BENZALKONIUM CHLORIDE	0.1 g in 100 g

Inactive Ingredients		
Ingredient Name	Strength	
GLYCERIN (UNII: PDC6A3C0OX)		
PALMITIC ACID (UNII: 2V16EO95H1)		
PEG-6 METHYL ETHER (UNII: WXH089JZ5E)		
PENTASO DIUM PENTETATE (UNII: 961TOZ5L7T)		
SODIUM CHLORIDE (UNII: 451W47IQ8X)		
SO DIUM PALMITATE (UNII: JQ43KP6296)		
SO DIUM PALM KERNELATE (UNII: 6 H9 1L1NXTW)		
SORBITOL (UNII: 506T60A25R)		
ETIDRO NATE TETRASO DIUM (UNII: CZZ9T1T1X4)		
WATER (UNII: 059QF0KO0R)		
BUTYLATED HYDRO XYTO LUENE (UNII: 1P9 D0 Z171K)		
TITANIUM DIO XIDE (UNII: 15FIX9 V2JP)		
FD&C YELLOW NO. 6 (UNII: H77VEI93A8)		

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:67510-0116-3	3 in 1 CELLO PACK	07/10/2020	
1	100 g in 1 BOX; 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	part333E	07/10/2020	

Labeler - Kareway Product, Inc. (121840057)

Revised: 7/2020 Kareway Product, Inc.