

NEXT 1 ANTI-BACTERIAL DEODORANT- benzalkonium chloride soap
Keefe Group

Next 1 TM Anti-bacterial Deodorant Bar Soap

Drug Facts

Active ingredient

Benzalkonium Chloride 0.1%

Purpose

Antibacterial

Uses

for washing to decrease bacteria on the skin

Warnings

For external use only

When using this product do not use in or near the eyes, in case of eye contact, flush with water

Stop use and ask a doctor if irritation or redness develop

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.

Inactive ingredients Sodium Palmate, Sodium Palm Kernelate, Water, Glycerin, Sodium Chloride, Sodium Lauryl Sulfate, Fragrance, Titanium Dioxide, Coconut Acid, Citric Acid, Etidronic Acid, Tetrasodium EDTA. May contain Caramel.

Distributed by:

Olivette Products, L.L.C.

Bridgeton, MO 63044

Phone: 1-800-325-8998

MADE IN COLOMBIA

- **Kills germs**
- **Odor protection**

Next 1 TM

Anti-bacterial

Deodorant

Bar Soap

Contains no animal fat

NET WT 5 OZ (142g)

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100% VEG

100% VEGETABLE

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NEXT 1 ANTI-BACTERIAL DEODORANT

benzalkonium chloride soap

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51870-004
Route of Administration	TOPICAL		
Active Ingredient/Active Moiety			
Ingredient Name		Basis of Strength	Strength
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7) (BENZALKONIUM - UNII: 7N6JUD5X6Y)		BENZALKONIUM CHLORIDE	0.1 g in 100 g
Inactive Ingredients			
Ingredient Name			Strength
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)			
SODIUM LAURYL SULFATE (UNII: 368GB5141J)			
EDETATE SODIUM (UNII: MP1J8420LU)			
CAMEL (UNII: T9D99G2B1R)			
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)			
GLYCERIN (UNII: PDC6A3C0OX)			
SODIUM CHLORIDE (UNII: 451W47IQ8X)			
COCONUT ACID (UNII: 40U37V505D)			

CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	
ETIDRONIC ACID (UNII: M2F465ROXU)	
WATER (UNII: 059QF0KO0R)	
SODIUM PALMATE (UNII: S0A6004K3Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51870-004-05	142 g in 1 BOX; Type 0: Not a Combination Product	06/15/2017	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	505G(a)(3)	06/15/2017	

Labeler - Keefe Group (839439601)

Revised: 1/2024

Keefe Group