

NEXT1 ANTI-BACTERIAL DEODORANT SPORT- benzalkonium chloride soap
Keefe Group

Next1™ Anti-bacterial Deodorant Sport Bar

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, Green 3, Red 33. 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™

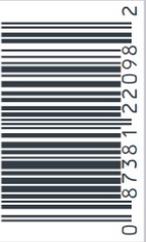
Anti-bacterial

Deodorant

Sport Bar

Contains no animal fat

NET WT 5 OZ (142g)

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NEXT1 ANTI-BACTERIAL DEODORANT SPORT

benzalkonium chloride soap

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:51870-003
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
WATER (UNII: 059QF0KO0R)	
SODIUM PALMATE (UNII: 50A6004K3Z)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	
GLYCERIN (UNII: PDC6A3C0OX)	
CAMEL (UNII: T9D99G2B1R)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SODIUM LAURYL SULFATE (UNII: 368GB5141J)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
COCONUT ACID (UNII: 40U37V505D)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

ETIDRONIC ACID (UNII: M2F465ROXU)	
EDETATE SODIUM (UNII: MP1J8420LU)	
FD&C GREEN NO. 3 (UNII: 3P3ONR6O1S)	
D&C RED NO. 33 (UNII: 9DBA0SBB0L)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:51870-003-05	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)