

**BOB BARKER ANTIBACTERIAL UN-WRAPPED BODY- benzethonium chloride soap**  
**Bob Barker Company Inc.**

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**Bob Barker® Antibacterial un-wrapped body soap**

***Drug Facts***

**Active Ingredient**

Benzethonium Chloride 0.1%

**Purpose**

Antibacterial

**Uses**

For washing to decrease bacteria on skin.

**Warnings**

**For External Use Only.**

**Do not use** this product on infants under 6.

**When using this product** avoid contact with eyes. In case of eye contact, flush with water.

**Stop use and ask 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

**Inactive Ingredients**

Sodium Palmate, Calcium Carbonate, Water, Sodium Palm Kernelate, Talcum, Palm Acid, Glycerin, Sodium Chloride, Sodium Lauryl Ether Sulfate, Titanium Dioxide, Palm Kernel Acid, Fragrance, Tetrasodium EDTA, Tetrasodium Etidronate

**PRINCIPAL DISPLAY PANEL - 500 Bar Case**

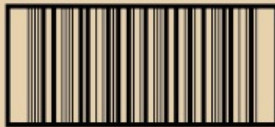
BobBarker®

ANTIBACTERIAL UNWRAPPED BODY SOAP

500 Bars-Total Net Weight 564.38 OZ. (35.27 LB) (16 KG)

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**MADE IN MALAYSIA  
MANUFACTURED FOR BOB BARKER COMPANY INC.,  
FUQUAY VARINA, NC 27526**





00744189005660

**ITEM # AU15-C**

**PO #**

**CARTON #**

**EXPIRATION DATE:**

**ITEM # AU15-C**  
**SIZE: # 1.5**



00744189005660

## BOB BARKER ANTIBACTERIAL UN-WRAPPED BODY

benzethonium chloride soap

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:53247-118
<b>Route of Administration</b>	CUTANEOUS		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZETHONIUM CHLORIDE</b> (UNII: PH41D05744) (BENZETHONIUM - UNII:1VU15B70BP)	BENZETHONIUM CHLORIDE	0.1 g in 100 g

## Inactive Ingredients

Ingredient Name	Strength
<b>Sodium Palmate</b> (UNII: S0A6004K3Z)	
<b>Calcium Carbonate</b> (UNII: H0G9379FGK)	
<b>Water</b> (UNII: 059QF0KO0R)	
<b>Sodium Palm Kernelate</b> (UNII: 6H91L1NXTW)	
<b>Talc</b> (UNII: 7SEV7J4R1U)	
<b>Palm Acid</b> (UNII: B6G0Y5Z616)	
<b>Glycerin</b> (UNII: PDC6A3C0OX)	
<b>Sodium Chloride</b> (UNII: 451W47IQ8X)	
<b>Sodium Laureth-3 Sulfate</b> (UNII: BPV390UAP0)	
<b>Titanium Dioxide</b> (UNII: 15FIX9V2JP)	
<b>Palm Kernel Acid</b> (UNII: 79P21R4317)	
<b>Edetate Sodium</b> (UNII: MP1J8420LU)	
<b>Etidronate Tetrasodium</b> (UNII: CZZ9T1T1X4)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-118-01	1000 in 1 CASE	06/01/2017	
1		11 g in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:53247-118-02	500 in 1 CASE	06/01/2017	
2		32 g in 1 APPLICATOR; Type 0: Not a Combination Product		
3	NDC:53247-118-03	144 in 1 CASE	06/01/2017	
3		75 g in 1 APPLICATOR; Type 0: Not a Combination Product		
4	NDC:53247-118-04	200 in 1 CASE	06/01/2017	
4		85 g in 1 APPLICATOR; Type 0: Not a Combination Product		

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph drug	M003	06/01/2017	

**Labeler** - Bob Barker Company Inc. (058525536)