

## **BOB BARKER ANTIBACTERIAL UNWRAPPED BODY- chloroxylenol soap**

**Bob Barker Company 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.*

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### **Bob Barker® Antibacterial unwrapped body soap**

#### ***Drug Facts***

#### **Active Ingredient**

Chloroxylenol 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, Palm (Elaeis Guineensis) Oil, Sodium Lauryl Sulfate, Glycerin, Sodium Chloride, Titanium Dioxide, Fragrance, Tetrasodium EDTA, Etidronic Acid

#### **PRINCIPAL DISPLAY PANEL - 500 Bar Case**

BobBarker®

ANTIBACTERIAL UNWRAPPED BODY SOAP

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

**BobBarker**<sup>®</sup>

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00744189005660

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





00744189005660

ITEM # AU15-C

PO #

CARTON #

EXPIRATION DATE:

ITEM # AU15-C  
SIZE: # 1.5**BOB BARKER ANTIBACTERIAL UNWRAPPED BODY**

chloroxylenol soap

**Product Information**

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:53247-116
<b>Route of Administration</b>	CUTANEOUS		

**Active Ingredient/Active Moiety**

Ingredient Name	Basis of Strength	Strength
CHLOROXYLENOL (UNII: 0F32U78V2Q) (CHLOROXYLENOL - UNII:0F32U78V2Q)	CHLOROXYLENOL	0.1 g in 100 g

**Inactive Ingredients**

Ingredient Name	Strength
Sodium Palmate (UNII: S0A6004K3Z)	
Calcium Carbonate (UNII: H0G9379FGK)	
Water (UNII: 059QF0KO0R)	
Sodium Palm Kernelate (UNII: 6H91L1NXTW)	
PALM OIL (UNII: 5QUO05548Z)	
Glycerin (UNII: PDC6A3C0OX)	
Sodium Chloride (UNII: 451W47IQ8X)	
Titanium Dioxide (UNII: 15FIX9V2JP)	
Edetate Sodium (UNII: MP1J8420LU)	
Etidronic Acid (UNII: M2F465ROXU)	

**Packaging**

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:53247-116-01	1000 in 1 CASE	06/01/2017	
1		11 g in 1 APPLICATOR; Type 0: Not a Combination Product		
2	NDC:53247-116-02	500 in 1 CASE	06/01/2017	
2		32 g in 1 APPLICATOR; Type 0: Not a Combination Product		

3	NDC:53247-116-03	144 in 1 CASE	06/01/2017	
3		75 g in 1 APPLICATOR; Type 0: Not a Combination Product		
4	NDC:53247-116-04	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 NOT FINAL	part333E	06/01/2017	

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

Revised: 9/2019

Bob Barker Company Inc.