# CLARIFYING FACE AND BODY WASH- benzoyl peroxide liquid Acneology Treatment Center, LLC

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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#### Clarifying Face & Body Wash

#### **Drug Facts**

#### **Active Ingredient**

Benzoyl Peroxide 2.5%

#### **Purpose**

For the treatment of acne

#### Uses

Adults and children 12 years and over:

- For the treatment of acne
- For acne-prone or oily skin

#### Warnings

**For external use only.** Avoid contact with eyes and eyelids. This product may cause irritation. Mild irritation may be reduced by using the product less frequently. If irritation becomes severe, discontinue use and consult a doctor.

#### Do not use

this product if you have very sensitive skin or if you are sensitive to benzoyl peroxide.

### Keep out of reach of children

• If swallowed, get medical help or contacting Poison Control Center right away.

#### **Directions**

Use 1 to 3 times daily on affected skin, work into lather, rinse off thoroughly.

#### Other Information

- May bleach hair or fabrics.
- Store at room temperature.
- Keep away from heat or direct sunlight.

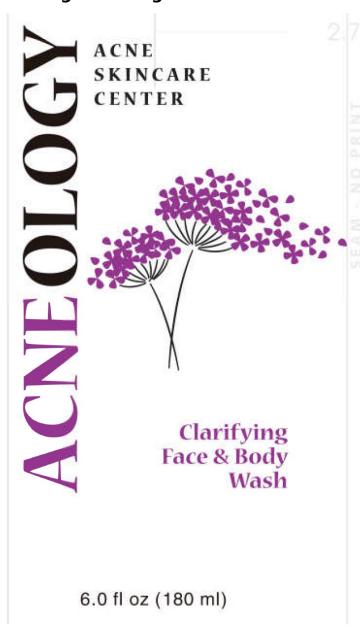
#### Inactive Ingredients

Purified Water, Ammonium Lauryl Sulfate, Glycerin, Disodium Cocoamphodiacetate, Propylene Glycol, Carbomer, Sodium Hydroxide.

#### **Questions or Comments?**

Call Toll-Free 800-439-4166

#### Package Labeling:



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Distributed by Acneology Skincare Center, San Jose, CA 95131 www.acneologyskincare.com

#### **CLARIFYING FACE AND BODY WASH**

benzoyl peroxide liquid

#### **Product Information**

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:73610-000
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZ N9A0GM) (BENZ OYL PEROXIDE - UNII: W9WZ N9A0GM)	BENZOYL PEROXIDE	25 mg in 1 mL

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)		
GLYCERIN (UNII: PDC6A3C0OX)		
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:73610-000- 00	180 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/16/2020	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	03/16/2020		

## Labeler - Acneology Treatment Center, LLC (047568655)

Revised: 9/2021 Acneology Treatment Center, LLC