ACNE FACE AND BODY- benzoyl peroxide shampoo, suspension
Face Reality, Inc.

Acne Face and Body Scrub

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 medication if you have very sensitive skin or if you are sensitive to benzoly peroxide.

Keep out of reach of children.

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, Cellulose Acetate, Disodium Cocoamphodiacetate, Propylene Glycol, Carbomer, Sodium Hydroxide, Fragrance.

Package Labeling:

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Distributed by:
Face Reality, Inc.
San Leandro, CA 94577
www.facerealityskincare.com

face reality™
SKINCARE

acne face
and body scrub
for acne-prone skin

6.0 FL OZ (180 mL)

ACNE FACE AND BODY

benzoyl peroxide shampoo, suspension

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:70707-101
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	25 mg in 1 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
AMMONIUM LAURYL SULFATE (UNII: Q7AO2R1M0B)	
GLYCERIN (UNII: PDC6A3C0OX)	
CELLULOSE ACETATE (UNII: 3J2P07GVB6)	
DISODIUM COCOAMPHODIACETATE (UNII: 18L9G3U51M)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
CARBOXYPOLYMETHYLENE (UNII: 0A5MM307FC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:70707-101-06	180 mL in 1 BOTTLE; Type 0: Not a Combination Product	12/30/2016	
2	NDC:70707-101-01	180 mL in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product	09/15/2011	09/15/2011

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	09/15/2011	

Labeler - Face Reality, Inc. (602958071)

Revised: 12/2023 Face Reality, Inc.