10 ACNE MED- benzoyl peroxide gel Face Reality, Inc.

10% Acne Med

Drug Facts

Active ingredient

Benzoyl Peroxide 10%

Purpose

Acne Treatment

Use

for the treatment of acne

Warnings

For external use only

Do not use

- if you have very sensitive skin
- if you are sensitive to benzoyl peroxide

When using this product

- skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time.
- avoid unnecessary sun exposure and use a sunscreen
- avoid contact with eyes, lips and mouth
- avoid contact with hair and dyed fabrics, which may be bleached by this product
- skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Irritation may be reduced by using the product less frequently or in a lower concentration.

Stop use and ask a doctor if

irritation becomes severe

Keep out of reach of children.

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

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one or three times daily
- because excessive drying of the skin may occur, start with one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor
- if bothersome dryness or peeling occurs, reduce application to once a day or every other day
- if going outside, apply sunscreen after using this product. If irritation or sensitivity develops, stop use of both products and ask a doctor.

Other information

• Store at room temperature. Keep away from heat or direct sunlight.

Inactive ingredients

Purified Water, Glycerin, Propylene Glycol, Carbomer, Sodium Hydroxide.

Questions or Comments?

1-866-477-3077

Package Labeling:



Tone Indicates Non-Print Areas

Active Ingredient

Purpose

Benzoyl Peroxide 10% For the treatment of Acne

Warnings

For external use only

Do not use ■ if you have very sensitive skin ■ if you are sensitive to benzoyl peroxide

When using this product ■ skin irritation and dryness is more likely to occur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medication at a time. ■ avoid unnecessary sun exposure and use a sunscreen ■ avoid contact with eyes, lips and mouth ■ avoid contact with hair and dyed fabrics, which may be bleached by this product ■ skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Irritation may be reduced by using the product less frequently or in a lower concentration.

Stop use and ask a doctor if ■ irritation becomes severe

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

■ **Directions:** Apply a thin layer to affected area as directed. Use for 15 to 30 minutes a day at first and increase wearing time gradually until it can be tolerated at night.

Inactive Ingredients

Purified Water, Glycerin, Propylene Glycol, Carbomer, Sodium Hydroxide.

This Unit Not Labeled For Retail Sale
QUESTIONS OR COMMENTS?
1-866-477-3077



Distributed by: Face Reality, Inc. Danville, CA 94526 facerealityskincare.com face reality

10% acne med

1.5 FL OZ (44 mL)

10 ACNE MED

benzoyl peroxide gel

Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:70707-151
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
BENZOYL PEROXIDE (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII: W9WZN9A0GM)	BENZOYL PEROXIDE	100 mg in 1 mL	

Inactive Ingredients		
Ingredient Name	Strength	
WATER (UNII: 059QF0KO0R)		
GLYCERIN (UNII: PDC6A3C0OX)		
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)		
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)		
SODIUM HYDROXIDE (UNII: 55X04QC32I)		

F	Packaging				
#	tem Code	Package Description	Marketing Start Date	Marketing End Date	
1	NDC:70707-151- 15	1 in 1 CARTON	10/15/2011		
1		44 mL in 1 TUBE; Type 0: Not a Combination Product			

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	10/15/2011	

Labeler - Face Reality, Inc. (602958071)

Revised: 10/2023 Face Reality, Inc.