

**MEDPURA BENZOYL PEROXIDE AQUEOUS BASE, ACNE TREATMENT GEL-**  
**benzoyl peroxide gel**  
**Akron Pharma Inc.**

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**MEDPURA 5% and 10% Benzoyl Peroxide**  
**Aqueous Base, Acne Treatment Gel**

Drug Facts

**Active ingredient**

Benzoyl peroxide 5%

Benzoyl peroxide 10%

**Purpose**

Acne treatment

**Use**

for the treatment of acne

**Warnings**

For external use only

Do not use this medication if you have very sensitive skin or 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 the 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.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Directions**

- **Sensitivity Test for a New User. Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, follow the directions stated below.**

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to 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 20°-25°C (68°-77°F)

### **Inactive ingredients**

carbomer, docusate sodium, methyl salicylate, purified water, propylene glycol, sodium hydroxide,

### **Questions or comments?**

call toll-free 1-877-225-6999

### **Manufactured for:**

Akron Pharma, Inc.  
Fairfield, NJ-07004  
[www.akronpharma.com](http://www.akronpharma.com)

# Benzoyl Peroxide 5% Gel

## Aqueous Base, Acne Treatment Gel



### Benzoyl Peroxide 5% Gel

Aqueous Base, Acne Treatment Gel

NDC 71399-9478-5

NET.WT 1.5 OZ (42.5g)

NDC 71399-9478-5

Benzoyl Peroxide 5% Gel



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**Akron Pharma**

Manufactured for:  
Akron Pharma, Inc.  
Fairfield, NJ- 07004  
www.akronpharma.com  
Rev.:04/24

Benzoyl Peroxide 5% Gel

NDC 71399-9478-5

**Drug Facts**

**Use** For the treatment of acne

**Warnings**

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

**For external use only**

**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 the 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

**Directions** Sensitivity Test for a New User: Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, or the small affected areas during the first 3 days, if no discomfort occurs, follow the directions stated below. Clean the skin thoroughly before applying the product. Over the entire affected area with a thin layer one to 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 ingredients** Carbomer, Cocousate Sodium, Methyl Salicylate, Propylene Glycol, Purified Water, Sodium Hydroxide

**Questions or comments?** call toll-free 1-877-225-6999

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Purpose** Acne treatment

**Drug Facts (continued)**

# Benzoyl Peroxide 5% Gel

## Aqueous Base, Acne Treatment Gel



### Benzoyl Peroxide 5% Gel

Aqueous Base, Acne Treatment Gel

NDC 71399-9478-6

NDC 71399-9478-6

Benzoyl Peroxide 5% Gel

NET.WT 2.1 OZ (60g)



**Akron Pharma**

Manufactured for:  
Akron Pharma, Inc.  
Fairfield, N.J. 07004  
www.akronpharma.com  
Rev.04/24

Benzoyl Peroxide 5% Gel

NDC 71399-9478-6

<b>Drug Facts</b>	<b>Stop use and ask a doctor if</b> ■ Irritation becomes severe less frequently or in a lower concentration. ■ Irritation may be reduced by using the product less frequently or in a lower concentration. ■ Irritation may be reduced by using the product less frequently or in a lower concentration.
<b>Warnings</b>	<b>Do not use if you</b> ■ Have very sensitive skin ■ Are sensitive to benzoyl peroxide ■ Use for external use only
<b>Directions</b> ■ Sensitive Test for a New User: Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, follow the directions stated below. ■ Clean the skin thoroughly before applying this product. ■ Cover the entire affected area with a thin layer one to three times daily. ■ Because excessive drying of the skin may occur, start with one application daily. ■ 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. ■ Irritation or sensitivity develops, stop use of both products and ask a doctor.	
<b>Other information</b> ■ Store at 20°-25°C (68°-77°F).	<b>Active ingredient</b> Benzoyl peroxide 5%
<b>Keep out of reach of children.</b> If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)	<b>Purpose</b> Acne treatment
<b>Questions or comments?</b> Call toll free 1-877-225-6999	<b>Drug Facts (continued)</b>

# Benzoyl Peroxide 5% Gel



# Aqueous Base, Acne Treatment Gel



## Benzoyl Peroxide 5% Gel

Aqueous Base, Acne Treatment Gel

NDC 71399-9478-9

NDC 71399-9478-9

Benzoyl Peroxide 5% Gel

NET.WT 3.2 OZ (90g)



**Akron Pharma**

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Akron Pharma, Inc.  
Fairfield, NJ- 07004  
www.akronpharma.com

Benzoyl Peroxide 5% Gel

NDC 71399-9478-9

**Drug Facts** (continued)

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Purpose** Acne treatment.

**Use for the treatment of acne**

**Warnings**

**Do not use if you** have very sensitive skin or are sensitive to benzoyl peroxide.

**For external use only**

**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 the 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.

**Other information** store at 20°-25°C (68°-77°F).

**Inactive ingredients** Carbomer, Docusate Sodium, Methyl Salicylate, Propylene Glycol, Purified Water, Sodium Hydroxide.

**Questions or comments?** call toll-free 1-877-225-6999



Maximum Strength

# Benzoyl Peroxide 10% Gel

## Aqueous Base, Acne Treatment Gel



Maximum Strength

# Benzoyl Peroxide 10% Gel

## Aqueous Base, Acne Treatment Gel

NDC 71399-9474-6

NET WT 2.1 OZ (60g)

NDC 71399-9474-6

Maximum Strength  
Benzoyl Peroxide  
10% Gel



**Akron Pharma**

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Akron Pharma, Inc.  
Fairfield, NJ- 07004  
www.akronpharma.com  
Rev.:04/24

Maximum Strength  
Benzoyl Peroxide  
10% Gel

NDC 71399-9474-6

**Drug Facts** (continued)

**Active ingredient**  
Benzoyl peroxide 10%

**Purpose**  
Acne treatment

**Warnings**  
Use for the treatment of acne

**For external use only**

**Do not use if you** have very sensitive skin, have sensitive skin, 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, stop use and use a sunscreen. Avoid contact with the eyes, lips, and mouth. Avoid contact with hair and dyed fabrics, which may be bleached by the 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.

**Other information** store at 20°-25°C (68°-77°F)

**Keep out of reach of children.** If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Directions** See **Warnings** for a New User. Apply product sparingly to one or two small affected areas during the first 3 days. If no discomfort occurs, then the directions state above. Clean the skin thoroughly before applying the product. Move the entire affected area with a thin layer of the product 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 both redness 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.

**Contains** benzoyl peroxide, diclofenac sodium, methyl salicylate, hydroxyethylcellulose, polyethylene glycol, carbomer, sodium hydroxide, and water.

**Questions or comments?** call toll-free 1-877-225-6999

Maximum Strength  
Benzoyl Peroxide  
10% Gel





## Inactive Ingredients

Ingredient Name	Strength
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0)	
<b>CARBOMER 980</b> (UNII: 4Q93RCW27E)	
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-9478-5	1 in 1 CARTON	09/13/2024	
1		42.5 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:71399-9478-6	1 in 1 CARTON	09/13/2024	
2		60 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:71399-9478-9	1 in 1 CARTON	09/13/2024	
3		90 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	09/13/2024	

## MEDPURA BENZOYL PEROXIDE AQUEOUS BASE, ACNE TREATMENT GEL

benzoyl peroxide gel

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:71399-9474
<b>Route of Administration</b>	TOPICAL		

## Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
<b>BENZOYL PEROXIDE</b> (UNII: W9WZN9A0GM) (BENZOYL PEROXIDE - UNII:W9WZN9A0GM)	BENZOYL PEROXIDE	100 mg in 1 mL

## Inactive Ingredients

Ingredient Name	Strength
<b>DOCUSATE SODIUM</b> (UNII: F05Q2T2JA0)	
<b>CARBOMER 980</b> (UNII: 4Q93RCW27E)	
<b>METHYL SALICYLATE</b> (UNII: LAV5U5022Y)	
<b>PROPYLENE GLYCOL</b> (UNII: 6DC9Q167V3)	
<b>WATER</b> (UNII: 059QF0KO0R)	
<b>SODIUM HYDROXIDE</b> (UNII: 55X04QC32I)	

## Product Characteristics

<b>Color</b>	white	<b>Score</b>	
<b>Shape</b>		<b>Size</b>	
<b>Flavor</b>		<b>Imprint Code</b>	
<b>Contains</b>			

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71399-9474-5	1 in 1 CARTON	09/13/2024	
1		42.5 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:71399-9474-6	1 in 1 CARTON	09/13/2024	
2		60 mL in 1 TUBE; Type 0: Not a Combination Product		
3	NDC:71399-9474-9	1 in 1 CARTON	09/13/2024	
3		90 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	09/13/2024	

**Labeler** - Akron Pharma Inc. (067878881)

