## CT07 BENZOYL PEROXIDE TREATMENT- benzoyl peroxide gel Pharmco Laboratories, Inc.

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# CT07 Benzoyl Peroxide Treatment Gel 2.5% DRUG FACTS

## **Active Ingredient**

2.5% Benzoyl Peroxide

## **Purpose**

Acne Treatment

## **USES**

For the treatment of acne

### WARNINGS

For external use only.

## <u>Do not use if you</u>

- Have very sensitive skin.
- Are sensitive to benzoyl peroxide.

## When using this product

- Skin irritation & 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 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
  lower concentration.
- If going outside, apply sunscreen after using this product.
- Stop use and ask a doctor if irritation becomes severe.
- Keep out of reach of children.
- If swallowed get medical help or call a poison control center immediately.
- Keep away from excessive heat or heat sources.

### **DIRECTIONS**

• Clean the skin thoroughly before applying this product.

- Clover the entire affected area with a thin layer one to three times per day.
- Because excessive drying 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 and 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 both products and ask doctor.

## OTHER INGREDIENTS

Carbomer, Edetate Disodium, Glycerine, Propylene Glycol, Saccharide Isomerate, Sodium Hydroxide, Water

## OTHER INFORMATION

Store at 15 - 25°C (59 - 77°F) Protect from heat. Keep container tightly closed.

## PRINCIPAL DISPLAY PANEL - 60 g Tube Carton

clearup

Anti-acne Clearing Gel

Micronized Benzoyl Peroxide

2.5%

PREVENTS & TREATS BREAKOUTS

MADE IN THE USA ADVANCED FORMULA GEL TEXTURE

60g / 2.1 oz



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PREVENTS & TREATS BREAKOUTS



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- · If going outside, apply sunscreen after using this product.
- · Stop use and ask a doctor if irritation becomes severe.
- · Keep out of reach of children.
- If swallowed get medical help or call a poison control center immediately.
- · Keep away from excessive heat or heat sources.

#### **DIRECTIONS**

- Clean the skin thoroughly before applying this product.
- · Cover the entire affected area with a thin layer
- one to three times a day.
- Because excessive drying 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 and 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 both products and ask a doctor.

#### OTHER INGREDIENTS

Carbomer, Edetate Disodium, Glycerine, Propylene Glycol, Saccharide Isomerate, Sodium Hydroxide, Water

#### OTHER INFORMATION

Store at 15 - 25°C (59 - 77°F) Protect from heat. Keep container tightly closed. Manufactured For Clearup® Philippines clearup.co

## CT07 BENZOYL PEROXIDE TREATMENT

benzoyl peroxide gel

### **Product Information**

Product Type HUMAN OTC DRUG Item Code (Source) NDC:58400-024

Route of Administration TOPICAL

## **Active Ingredient/Active Moiety**

Ingredient Name	<b>Basis of Strength</b>	Strength
Benzoyl Peroxide (UNII: W9WZ N9A0GM) (Benzoyl Peroxide -	Benzoyl Peroxide	25 mg in 1 g

Inactive Ingredients				
Ingredient Name	Strength			
CARBOMER HOMOPOLYMER, UNSPECIFIED TYPE (UNII: 0A5MM307FC)				
Edetate Disodium (UNII: 7FLD91C86K)				
GLYCERIN (UNII: PDC6A3C0OX)				
Propylene Glycol (UNII: 6DC9Q167V3)				
Saccharide Isomerate (UNII: W8K377W98I)				
Sodium Hydroxide (UNII: 55X04QC32I)				
Water (UNII: 059QF0KO0R)				

ı	Packaging					
	#	Item Code	Package Description	Marketing Start Date	Marketing End Date	
l	1	NDC:58400-024- 02	60 g in 1 TUBE; Type 0: Not a Combination Product	07/01/2024		

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC monograph drug	M006	07/01/2024			

## Labeler - Pharmco Laboratories, Inc. (096270814)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Pharmco Laboratories, Inc.		119071697	MANUFACTURE(58400-024) , LABEL(58400-024) , PACK(58400-024)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Pharmco Laboratories, Inc.		119071691	MANUFACTURE(58400-024) , LABEL(58400-024) , PACK(58400-024)

<b>Establishment</b>			
Name	Address	ID/FEI	Business Operations
Pharmco Laboratories, Inc.		096270814	MANUFACTURE(58400-024) , LABEL(58400-024) , PACK(58400-024)

Revised: 6/2024 Pharmco Laboratories, Inc.