

BENZOYL PEROXIDE- benzoyl peroxide lotion
Innovida Pharmaceutique Corporation

Active Ingredient

Benzoyl peroxide USP 10%

Purpose

Acne treatment

Uses

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

- keep away from eyes, lips and mouth
- avoid unnecessary sun exposure and use a sunscreen · avoid contact with hair or dyed fabric, including carpet and clothing which may be bleached by this product
- skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Mild irritation may be reduced by using the product less frequently or in a lower concentration. If irritation becomes severe, discontinue use; if irritation still continues, consult a doctor
- using other topical acne medication at the same time or immediately following the use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Avoid contact with the eyes. If contact occurs, flush thoroughly with water.

Directions

Cleanse skin thoroughly before applying medication. Cover the entire affected area with a thin layer 1-3 times daily. If bothersome dryness or peeling occurs, reduce application to once a day.

Other information

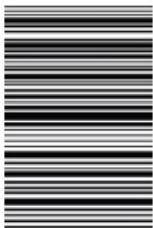
Keep tightly closed. Avoid storing at extreme temperatures (below 40° F and above 100° F).

Inactive Ingredients

carbomer, disodium EDTA, laureth-4, sodium hydroxide, titanium dioxide, water

Product label

NDC: 71800-046-09



INNVIDA
PHARMACEUTIQUE CORPORATION

Benzoyl Peroxide

Acne Medication, Lotion,
Benzoyl Peroxide 10%

TOPICAL ADMINISTRATION

1 oz (29.5 mL)

Drug Facts	Purpose
Active ingredient Benzoyl peroxide 10%.....	Acne treatment
Uses • 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 • keep away from eyes, lips and mouth • avoid unnecessary sun exposure and use a sunscreen • avoid contact with hair or dyed fabric, including carpet and clothing which may be bleached by this product • skin irritation may occur, characterized by redness, burning, itching, peeling, or possibly swelling. Mild irritation may be reduced by using the product less frequently or in a lower concentration. If irritation becomes severe, discontinue use; if irritation still continues, consult a doctor. • using other topical acne medication at the same time or immediately following the use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.	
Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away. Avoid contact with the eyes. If contact occurs, flush thoroughly with water.	
Directions Cleanse skin thoroughly before applying medication. Cover the entire affected area with a thin layer 1-3 times daily. If bothersome dryness or peeling occurs, reduce application to once a day.	
Other information Keep tightly closed. Avoid storing at extreme temperatures (below 40° F and above 100° F).	
Inactive ingredients: carbomer, disodium EDTA, laureth-4, sodium hydroxide, titanium dioxide, water	

Mfd. for: Innovida Pharmaceutique Corporation, Phoenix, AZ 85040, Tel.: 888-462-4166

INN^oVIDA
PHARMACEUTIQUE CORPORATION

NDC: 71800-046-09

Benzoyl Peroxide

Acne Medication, Lotion, Benzoyl Peroxide 10%

TOPICAL ADMINISTRATION

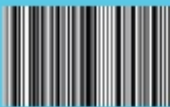
1 oz (29.5 mL)



Benzoyl Peroxide

Acne Medication, Lotion, Benzoyl Peroxide 10%

INN^oVIDA
PHARMACEUTIQUE CORPORATION



Tel: 888-462-4166

Mtd. for:
Phonvda Pharmaceutiqe Corporation
Phonvdx, AZ 85040

Drug Facts
Active ingredient
Benzoyl peroxide 10%
Uses - for the treatment of acne
Warnings - Do not use this medicine if you are allergic to benzoyl peroxide.
- Do not use this medicine if you have very sensitive skin or if you have very dry skin.
When using this product - Keep away from eyes, face and mouth. - At the first sign of irritation, stop use and use your moisturizer. - Avoid contact with hair and pedicure. - Avoid contact with pedicure. - Avoid contact with hair and pedicure. - Avoid contact with hair and pedicure.
Directions - Cleanse the skin thoroughly before applying. - Apply a thin layer to the affected areas. - Cover the entire affected area with a thin layer. - Rinse thoroughly with water. - Avoid contact with the eyes. If contact occurs, flush thoroughly with water.
Keep out of reach of children. - If swallowed, get medical help or contact Poison Control Center. - Do not use if the seal is broken. - Avoid contact with the eyes. If contact occurs, flush thoroughly with water.
Other important information - Avoid starting at extreme temperatures. - Avoid use if you are pregnant or breastfeeding. - Avoid use if you are pregnant or breastfeeding.
Inactive ingredients: carbonyl diethanolamine, EDTA, laureth-4, sodium hydroxide, triethanolamine, water.

BENZOYL PEROXIDE

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71800-046
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
CARBOXYMETHYLCELLULOSE SODIUM, UNSPECIFIED (UNII: K679OBS311)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
LAURETH-4 (UNII: 6HQ855798J)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
WATER (UNII: 059QF0KO0R)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:71800-046-09	1 in 1 CARTON	02/10/2025	
1		29.5 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	02/10/2025	

Labeler - Innovida Pharmaceutique Corporation (080892908)

Revised: 2/2025

Innovida Pharmaceutique Corporation