

BENZOYL PEROXIDE- benzoyl peroxide lotion
Innovida Pharmaceutique Corporation

Active Ingredient

Benzoyl peroxide 5%

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.

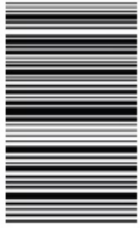
Directions

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).

NDC: 71800-044-09



INN^oVIDA
PHARMACEUTIQUE CORPORATION

Benzoyl Peroxide

Acne Medication, Lotion,
Benzoyl Peroxide 5%

TOPICAL ADMINISTRATION

1 oz (29.5 mL)

Drug Facts

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

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

BENZOYL PEROXIDE

benzoyl peroxide lotion

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:71800-044
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
BENZOYL PEROXIDE (UNII: W9WZ N9A0GM) (BENZOYL PEROXIDE - UNII:W9WZ N9A0GM)	BENZOYL PEROXIDE	50 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:71800-044-09	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