

**PANOXYL- benzoyl peroxide soap**  
**Crown Laboratories**

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**Panoxyl Acne Treatment Bar**

***Active ingredient***

Benzoyl peroxide 10%

***Purpose***

Acne medication

***Use***

- for the treatment of acne

***Warnings***

**For external use only**

**Do not use if you**

- have very sensitive skin
- are sensitive to benzoyl peroxide

**When using 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.
- 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

**Stop use and ask a doctor if**

- irritation becomes severe

**If pregnant or breast-feeding,**

ask a health professional before use.

**Keep out of reach of children.**

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

**Directions**

- clean the skin thoroughly before applying this product
- work into a lather
- cover the entire affected area with a thin layer and rinse thoroughly one to three times daily
- rinse thoroughly and pat dry
- 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) [see USP Controlled Room Temperature].

**Inactive ingredients**

Aqua, Cetearyl Alcohol, Decyl Glucoside, Dextrin, Glycerin, Sodium Chloride, Sodium Cocoyl Isethionate, Tetrasodium Glutamate Diacetate

**Questions?**

call **1-833-279-6522**

**Panoxyl Acne Treatment Bar Carton**

**PanOxyl**®

**Acne Treatment Bar**

**for Face & Body**

**10% BENZOYL PEROXIDE**

**MAXIMUM STRENGTH**

Clears existing acne and helps prevent new breakouts

Maximum strength without a prescription

No. 1 DERMATOLOGIST RECOMMENDED BENZOYL PEROXIDE BRAND

Net wt. 4 oz (113 g)



## PANOXYL

benzoyl peroxide soap

### Product Information

|                                |                |                           |               |
|--------------------------------|----------------|---------------------------|---------------|
| <b>Product Type</b>            | HUMAN OTC DRUG | <b>Item Code (Source)</b> | NDC:0316-0264 |
| <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  | 12.6 g<br>in 113 g |

### Inactive Ingredients

| Ingredient Name   | Strength |
|---|----------|
| <b>ICODEXTRIN</b> (UNII: 2NX48Z0A9G)                      |          |
| <b>DECYL GLUCOSIDE</b> (UNII: Z17H97EA6Y)                 |          |
| <b>GLYCERIN</b> (UNII: PDC6A3C0OX)                        |          |
| <b>SODIUM CHLORIDE</b> (UNII: 451W47IQ8X)                 |          |
| <b>WATER</b> (UNII: 059QF0KO0R)                           |          |
| <b>SODIUM COCOYL ISETHIONATE</b> (UNII: 518XTE8493)       |          |
| <b>TETRASODIUM GLUTAMATE DIACETATE</b> (UNII: 5EHL50I4MY) |          |
| <b>CETOSTEARYL ALCOHOL</b> (UNII: 2DMT128M1S)             |          |

**Packaging**

| # | Item Code        | Package Description                                  | Marketing Start Date | Marketing End Date |
|---|------------------|--|----------------------|--------------------|
| 1 | NDC:0316-0264-04 | 113 g in 1 CARTON; Type 0: Not a Combination Product | 07/01/2023           |                    |
| 2 | NDC:0316-0264-01 | 43 g in 1 CARTON; Type 0: Not a Combination Product  | 02/23/2024           |                    |

**Marketing Information**

| Marketing Category | Application Number or Monograph Citation | Marketing Start Date | Marketing End Date |
|--------------------|--|----------------------|--------------------|
| OTC Monograph Drug | M006                                     | 07/01/2023           |                    |

**Labeler** - Crown Laboratories (119508400)

Revised: 1/2026

Crown Laboratories