PANOXYL- salicylic acid soap
Crown Laboratories

Panoxyl Acne Cleansing Bar

Active ingredient

Salicylic Acid 2%

Purpose

Acne medication

Use

- for the treatment of acne

Warnings

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.

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

- wet face and work product into a lather
- cover the entire affected area with a thin layer 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

Other information

- Store at 20 ^{o}- 25 ^{o}C (68 ^{o}- 77 ^{o}F) [see USP Controlled Room Temperature].

Inactive ingredients

Aqua, Butyrospermum Parkii (Shea Butter), Cocamidopropyl Hydroxysultaine, Galactoarabinan, Glycerin, Mandelic Acid, Sodium Chloride, Sodium Cocoate*, Sodium Gluconate, Sodium Palm Kernelate*, Sodium Palmate, Tetrasodium EDTA, Tetrasodium Glutamate Diacetate, Titanium Dioxide (CI 77891), Zinc PCA *contains one or more of these ingredients

Questions?

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

Panoxyl Acne Treatmet Bar Carton

PanOxyl

Acne Cleansing Bar

for Face & Body

2% SALICYLIC ACID

ACNE TREATMENT BAR

DAILY CONTROL

Clears and helps prevent acne

Gentle, hypoallergenic formula

Helps exfoliate for a clearer complexion

DERMATOLOGIST RECOMMENDED

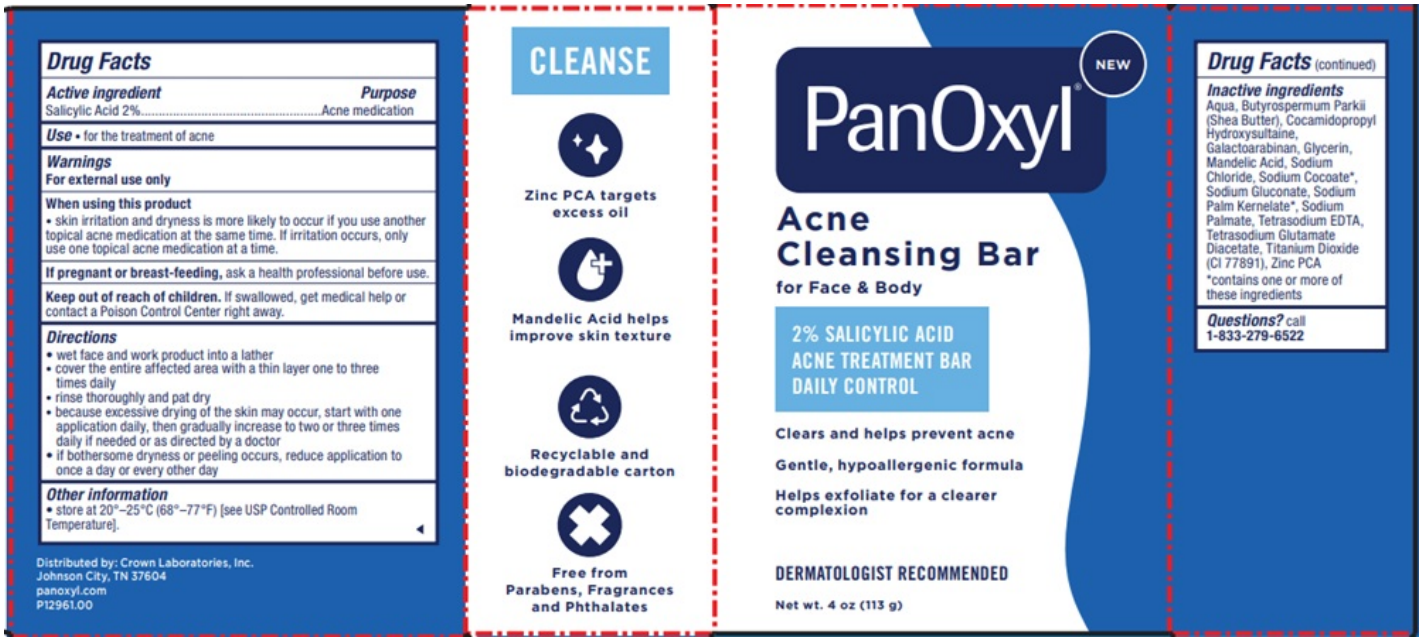
Net wt. 4 oz (113 g)

Distributed by: Crown Laboratories, Inc.

Johnson City, TN 37604

panoxyl.com

P12961.00



PANOXYL

salicylic acid soap

Product Information

Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0288
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	20 mg in 1 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
GALACTOARABINAN (UNII: SL4SX1O487)	
TETRASODIUM EDTA (UNII: MP1J8420LU)	
TITANIUM DIOXIDE (UNII: 15FIX9V2JP)	
BUTYROSPERMUM PARKII (SHEA) BUTTER (UNII: K49155WL9Y)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	
ZINC PCA (UNII: C32PQ86DH4)	
GLYCERIN (UNII: PDC6A3C0OX)	
TETRASODIUM GLUTAMATE DIACETATE (UNII: 5EHL50I4MY)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
MANDELIC ACID (UNII: NH496X0UJX)	
SODIUM COCOATE (UNII: R1TQH25F4I)	
SODIUM PALM KERNELATE (UNII: 6H91L1NXTW)	
SODIUM PALMATE (UNII: S0A6004K3Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0288-01	113 g in 1 CARTON; Type 0: Not a Combination Product	01/05/2026	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	01/05/2026	

Labeler - Crown Laboratories (119508400)

Revised: 1/2026

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