PANOXYL ACNE GEL WASH- salicylic acid gel
Crown Laboratories, Inc.

PanOxyl Acne Gel Wash

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 in the morning and at night
- 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 day

Other information

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

Inactive ingredients

1,2-Hexanediol, Allantoin, Aloe Barbadensis Leaf Juice, Aqua, Betaine, Camellia Sinensis Leaf Extract, Caprylyl Glycol, Citric Acid, Cocamidopropyl Hydroxysultaine, Decyl Glucoside, Glycerin, PEG-150 Distearate, Potassium Sorbate, Saccharide Isomerate, Sodium Benzoate, Sodium C14-16 Olefin Sulfonate, Sodium Chloride, Sodium Citrate, Sodium Hydroxide, Spirulina Platensis Extract, Zinc PCA

Questions?

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

PanOxyl Acne Gel Wash Label

NEW

PanOxyl

Acne Gel Wash

for Face

2% SALICYLIC ACID

ACNE TREATMENT WASH

Clears and helps prevent acne

Gentle gel-to-foam formula

Zinc PCA helps target excess oil

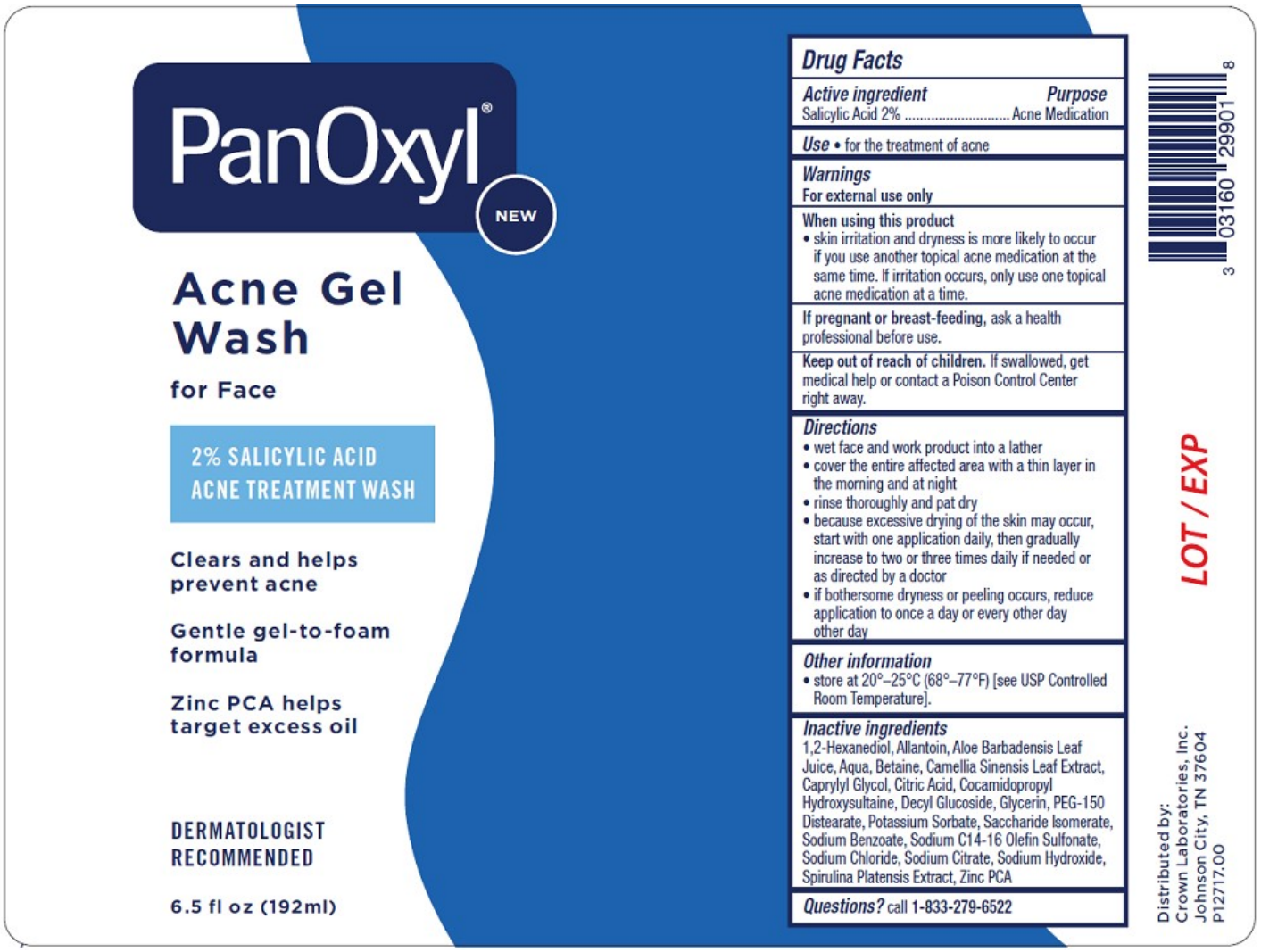
DERMATOLIGST RECOMMENDED

6.5 fl oz (192ml)

Distributed by: Crown Laboratories, Inc.

Johnson City, TN 37604

P12717.00



PANOXYL ACNE GEL WASH

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:0316-0299
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
1,2-HEXANEDIOL (UNII: TR046Y3K1G)	
GREEN TEA LEAF (UNII: W2ZU1RY8B0)	
CAPRYLYL GLYCOL (UNII: 00YIU5438U)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

SACCHARIDE ISOMERATE (UNII: W8K377W98I)	
SODIUM CITRATE (UNII: 1Q73Q2JULR)	
GLYCERIN (UNII: PDC6A3C0OX)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
BETAINE (UNII: 3SCV180C9W)	
CITRIC ACID (UNII: 2968PHW8QP)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
SPIRULINA PLATENSIS (UNII: 9L3TIH1UUE)	
WATER (UNII: 059QF0KO0R)	
SODIUM BENZOATE (UNII: OJ245FE5EU)	
ALOE BARBADENSIS LEAF JUICE (UNII: RUE8E6T4NB)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
PEG-150 DISTEARATE (UNII: 6F36Q0I0AC)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
ZINC PCA (UNII: C32PQ86DH4)	
ALLANTOIN (UNII: 344S277G0Z)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0299-01	192 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/01/2025	
2	NDC:0316-0299-03	89 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/01/2025	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	07/01/2025	

Labeler - Crown Laboratories, Inc. (119508400)

Establishment

Name	Address	ID/FEI	Business Operations
Crown Laboratories, Inc.		119508400	manufacture(0316-0299)

Revised: 1/2026 Crown Laboratories, Inc.