

PANOXYL CLARIFYING EXFOLIANT- salicylic acid liquid
Crown Laboratories, Inc.

PanOxyl Clarifying Exfoliant

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

- shake well before using
- clean the skin thoroughly before applying this product
- using a cotton pad, cover the entire affected area with a thin layer one to three times daily
- 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 ° - 25 °C (68 ° - 77 °F) [see USP Controlled Room Temperature].

Inactive ingredients

Betaine, Butylene Glycol, Chamomilla Recutita (Matricaria) Flower Extract, Glycerin, Hydroxyacetophenone, Pentyleneglycol, Phenoxyethanol, Potassium Sorbate, Propanediol, Purified Water, Sodium Benzoate, Sodium Citrate Dihydrate, Sorbic Acid, Spirulina Platensis (Blue Algae) Extract

Questions or comments?

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

Panoxyl Clarifying Exfoliant Bottle

new

NDC 0316-0296-04

DERMATOLOGIST RECOMMENDED

PanOxyl®

Clarifying Exfoliant

2% Salicylic Acid

- Clears and Helps Prevent Acne
- Unclogs and Minimizes the Appearance of Pores
- Gentle Alcohol-Free Formula
- Blue Algae and Antioxidants Help Calm Irritation and Redness

4 fl oz (118 mL)

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 PANOXYL is a registered trademark of Crown Laboratories, Inc.

 Manufactured by: Crown Laboratories, Inc., Johnson City, TN 37604

 P12117.00

PANOXYL CLARIFYING EXFOLIANT

salicylic acid liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:0316-0296
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
BETAINE (UNII: 3SCV180C9W)	
BUTYLENE GLYCOL (UNII: 3XUS85K0RA)	
MATRICARIA CHAMOMILLA FLOWERING TOP (UNII: 3VNC7T6Z02)	
GLYCERIN (UNII: PDC6A3C0OX)	
HYDROXYACETOPHENONE (UNII: G1L3HT4CMH)	
PENTYLENE GLYCOL (UNII: 50C1307PZG)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
PROPANEDIOL (UNII: 5965N8W85T)	
WATER (UNII: 059QF0KO0R)	

SODIUM BENZOATE (UNII: OJ245FE5EU)	
SORBIC ACID (UNII: X045WJ989B)	
TRISODIUM CITRATE DIHYDRATE (UNII: B22547B95K)	
SPIRULINA PLATENSIS (UNII: 9L3TIH1UUE)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:0316-0296-04	118 mL in 1 BOTTLE; Type 0: Not a Combination Product	06/02/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	06/01/2023	

Labeler - Crown Laboratories, Inc. (079035945)

Establishment

Name	Address	ID/FEI	Business Operations
Crown Laboratories, Inc.		079035945	manufacture(0316-0296)

Revised: 10/2023

Crown Laboratories, Inc.