NEUTROGENA ALL IN 1 ACNE CONTROL DAILY SCRUB- salicylic acid liquid Kenvue Brands LLC

Neutrogena® All-in-1 Acne Control daily scrub

Drug Facts

Active ingredient

Salicylic Acid 2%

Purpose

Acne treatment

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.
- avoid contact with eyes. If contact occurs, flush thoroughly with water

Keep out of reach of children. If swallowed, get medical help or contact a Poison Control Center right away.

Directions

Cleanse twice a day. Wet face. Apply to hands, add water and work into a lather. Massage face gently. Rinse thoroughly.

Other Information

Store at Room Temperature.

Inactive ingredients

water, cetyl alcohol, PPG-15 stearyl ether, cellulose, glycerin, Polysorbate 60, steareth-21, potassium cetyl phosphate, xanthan gum, fragrance, disodium EDTA, menthyl lactate, sodium hydroxide

Questions?

Call toll- free **800-582-4048 or 215-273-8755** (collect). www.neutrogena.com

Distributed by:

Kenvue Brands LLC

Summit, NJ 07901

PRINCIPAL DISPLAY PANEL - 124 mL Tube Label

All-in-1

Acne

Control

daily scrub

works on acne's

past, present & future

- exfoliates past acne marks
- clears present breakouts
- helps prevent future breakouts

Neutrogena ®

salicylic acid acne treatment

4.2 FL. OZ. (124mL)



NEUTROGENA ALL IN 1 ACNE CONTROL DAILY SCRUB

salicylic acid liquid

Product Information Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0092 Route of Administration TOPICAL

Active Ingredient/Active Moiety Ingredient Name Basis of Strength SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ) SALICYLIC ACID (UNII: 0414PZ4LPZ) SALICYLIC ACID 20 mg in 1 mL

Inactive Ingredients			
Ingredient Name	Strength		
WATER (UNII: 059QF0KO0R)			
CETYL ALCOHOL (UNII: 936JST6JCN)			
PPG-15 STEARYL ETHER (UNII: 1II18XLS1L)			
POWDERED CELLULOSE (UNII: SMD1X3XO9M)			
STEARETH-21 (UNII: 53J3F32P58)			

POLYSORBATE 60 (UNII: CAL22UVI4M)	
POTASSIUM CETYL PHOSPHATE (UNII: 03KCY6P7UT)	
XANTHAN GUM (UNII: TTV12P4NEE)	
EDETATE DISODIUM (UNII: 7FLD91C86K)	
MENTHYL LACTATE, (-)- (UNII: 2BF9E65L7I)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
GLYCERIN (UNII: PDC6A3C0OX)	

Packaging			
# Item Code	Package Description	Marketing Start Date	Marketing End Date
1 NDC:69968- 0092-4	124 mL in 1 TUBE; Type 0: Not a Combination Product	06/01/2016	

Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
OTC Monograph Drug	M006	06/01/2016			

Labeler - Kenvue Brands LLC (118772437)

Revised: 4/2025 Kenvue Brands LLC