NEUTROGENA OIL FREE ACNE WASH- salicylic acid liquid
Johnson & Johnson Consumer Inc.

Neutrogena ® Oil-Free Acne Wash

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, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Betaine, Sodium Chloride, PEG-80 Sorbitan Laurate, Citric Acid, Disodium EDTA, C12-15 Alkyl Lactate, Benzalkonium

Chloride, Fragrance, Cocamidopropyl PG-Dimonium Chloride Phosphate, Glycerin, Aloe Barbadensis Leaf Extract, Chamomilla Recutita (Matricaria) Flower Extract, Propylene Glycol, Sodium Hydroxide, Yellow 5, Red 40

Questions?

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

Distributed by:

JOHNSON & JOHNSON

CONSUMER INC.

Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 269 mL Bottle Label

Oil-Free

Acne Wash

MICRO **CLEAR** ®

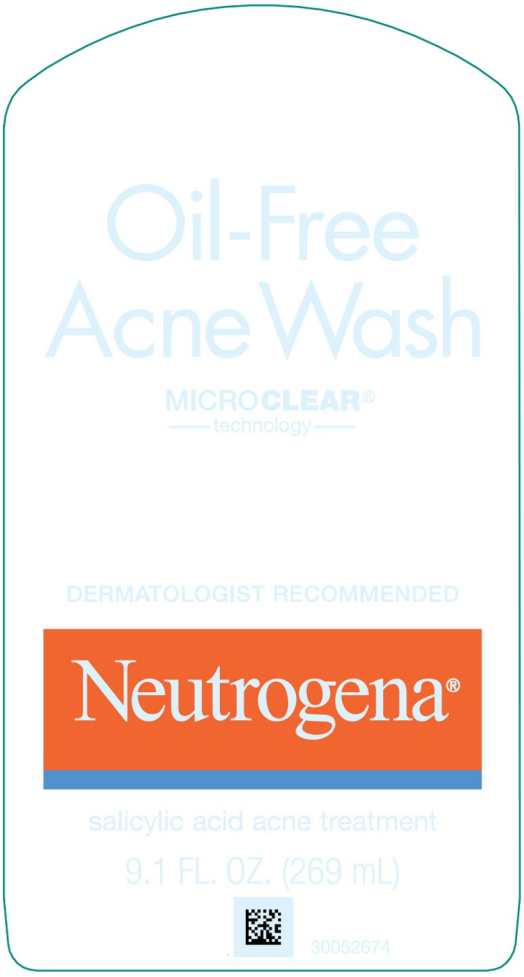
technology

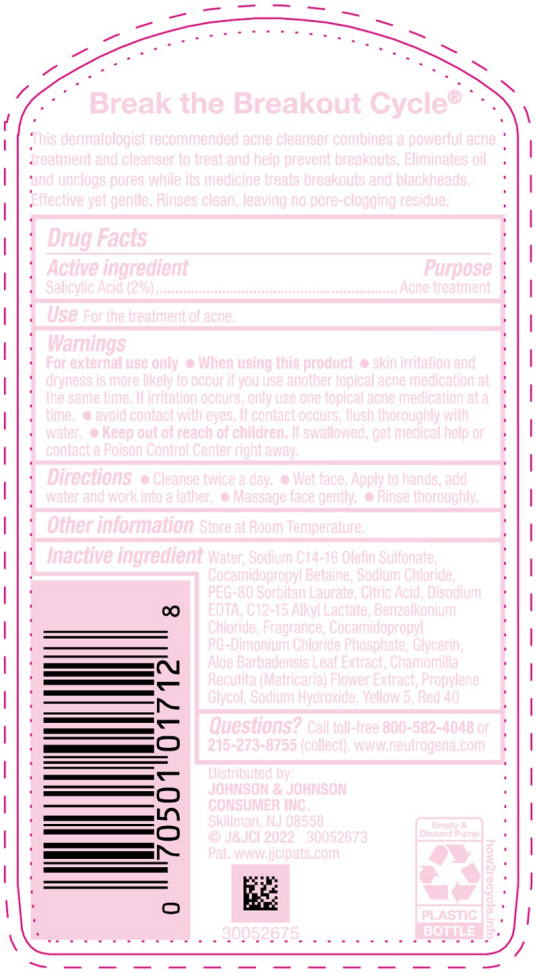
DERMATOLOGIST RECOMMENDED

Neutrogena ®

salicylic acid acne treatment

9.1 FL. OZ. (269 mL)





NEUTROGENA OIL FREE ACNE WASH

salicylic acid liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0770
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 mL

Inactive Ingredients	
Ingredient Name	**Strength**
WATER (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
PEG-80 SORBITAN LAURATE (UNII: 239B50Y732)	
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)	

EDETATE DISODIUM (UNII: 7FLD91C86K)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
GLYCERIN (UNII: PDC6A3C0OX)	
ALOE VERA LEAF (UNII: ZY81Z83H0X)	
CHAMOMILE (UNII: FGL3685T2X)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	
FD&C RED NO. 40 (UNII: WZB9127XOA)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0770-1	12 in 1 TRAY	07/29/2022	
1		14 mL in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:69968-0770-6	177 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/29/2022	
3	NDC:69968-0770-9	269 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product	07/29/2022	
4	NDC:69968-0770-2	2 in 1 CARTON	07/29/2022	
4		269 mL in 1 BOTTLE, PUMP; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	07/29/2022	

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 11/2023 Johnson & Johnson Consumer Inc.