NEUTROGENA ACNE PROOFING DAILY SCRUB- salicylic acid gel
Johnson & Johnson Consumer Inc.

Disclaimer: Most OTC drugs are not reviewed and approved by FDA, however they may be marketed if they comply with applicable regulations and policies. FDA has not evaluated whether this product complies.

Neutrogena Acne Proofing 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 Rome Temperature

Inactive ingredients

Water, Sodium C14-16 Olefin Sulfonate, Cocamidopropyl Hydroxysultaine, Sorbitol Cellulose, Sodium Hydrolyzed Potato Starch Dodecenylsuccinate, Acrylates Crosspolymer-4, Panthenol, Sodium Hydroxide, Fragrance, Microcrystalline Wax, C12-15 Alkyl Lactate, Benzalkonium Chloride, Disodium EDTA, Cocamidopropyl PG-Dimonium Chloride Phosphate, Talc, Yellow 5 Lake, Red 30

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 - 119 g Tube Label

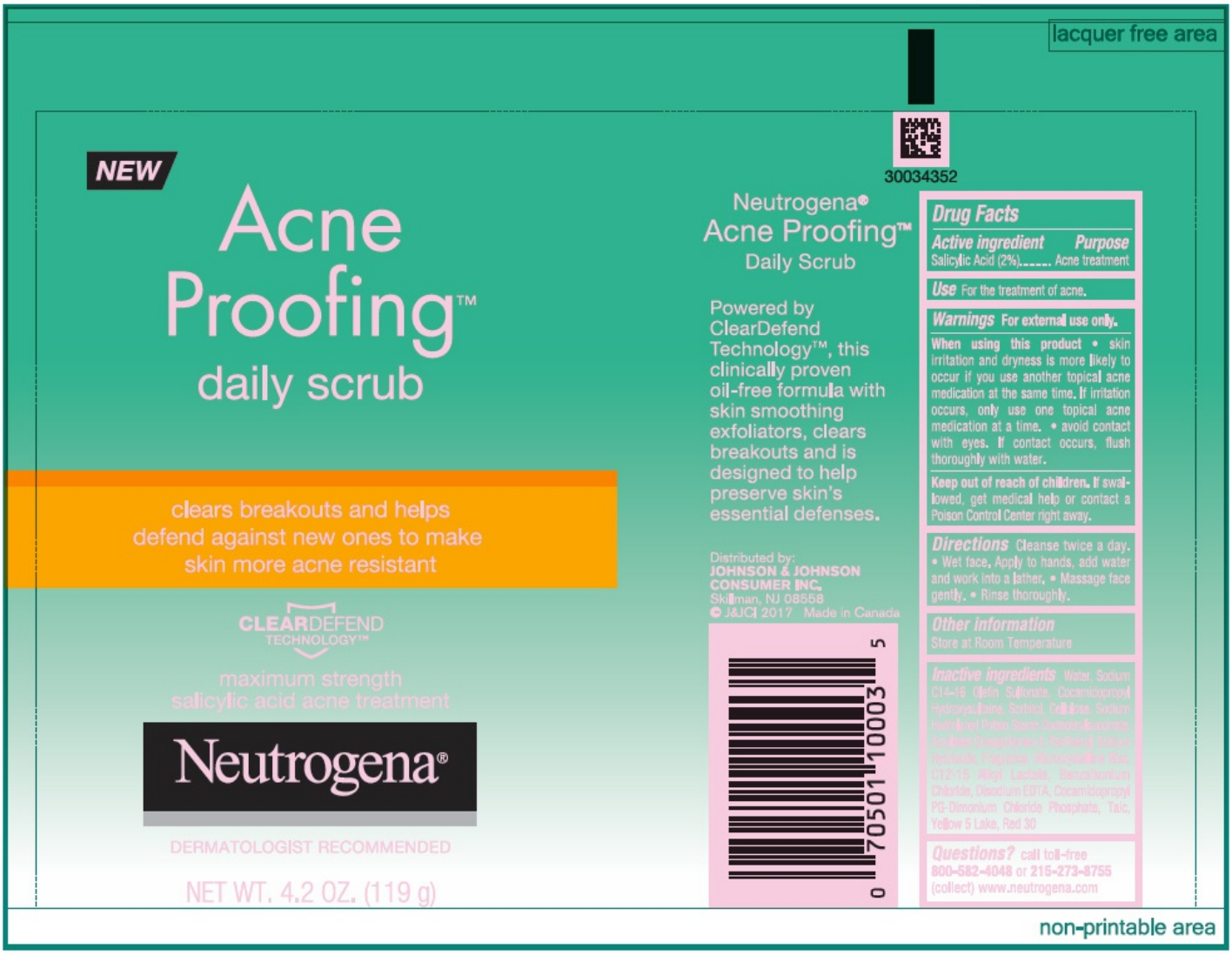
NEW

Acne
Proofing™
daily scrub
clears breakouts and helps
defend against new ones to make
skin more acne resistant

CLEARDEFEND
TECHNOLOGY ™

maximum strength
salicylic acid acne treatment

Neutrogena ®
DERMATOLOGIST RECOMMENDED
NET WT . 4.2 OZ. (119 g)



NEUTROGENA ACNE PROOFING DAILY SCRUB

salicylic acid gel

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:69968-0112
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)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	

SORBITOL (UNII: 506T60A25R)	
POWDERED CELLULOSE (UNII: SMD1X3XO9M)	
PANTHENOL (UNII: WV9CM0O67Z)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
MICROCRYSTALLINE WAX (UNII: XOF597Q3KY)	
C12-15 ALKYL LACTATE (UNII: GC844VRD7E)	
BENZALKONIUM CHLORIDE (UNII: F5UM2KM3W7)	
D&C RED NO. 30 (UNII: 2S42T2808B)	
COCAMIDOPROPYL PROPYLENE GLYCOL-DIMONIUM CHLORIDE PHOSPHATE (UNII: H2KVQ74JM4)	
EDETATE DISODIUM ANHYDROUS (UNII: 8NLQ36F6MM)	
TALC (UNII: 7SEV7J4R1U)	
FD&C YELLOW NO. 5 (UNII: I753WB2F1M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:69968-0112-4	119 g in 1 TUBE; Type 0: Not a Combination Product	10/01/2017	05/22/2024

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	10/01/2017	05/22/2024

Labeler - Johnson & Johnson Consumer Inc. (118772437)

Revised: 6/2023

Johnson & Johnson Consumer Inc.