NEUTROGENA ACNE PROOFING CLEANSER- 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™ gel cleans er

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 lather. Massage face gently. Rinse thoroughly.

Inactive ingredients

Water, Cocamidopropyl Hydroxysultaine, Sodium Methyl Cocoyl Taurate, Sodium Chloride, Glycerin, Sodium Hydrolyzed Potato Starch Dodecenylsuccinate, Fragrance, Sodium Hydroxide, PEG-120 Methyl Glucose Dioleate, Panthenol, C12-15 Alkyl Lactate, Disodium EDTA, Cocamidopropyl PG-Dimonium Chloride Phosphate, Benzalkonium Chloride

Other information

Store at room temperature.

Questions?

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

Distributed by: **JOHNSON & JOHNSON CONSUMER INC.** Skillman, NJ 08558

PRINCIPAL DISPLAY PANEL - 170 g Bottle Label

NEW

Acne Proofing™ gel cleanser

clears breakouts and helps defend against new ones to make skin more acne resistant

 $\begin{array}{c} CLEARDEFEND \\ TECHNOLOGY^{TM} \end{array}$

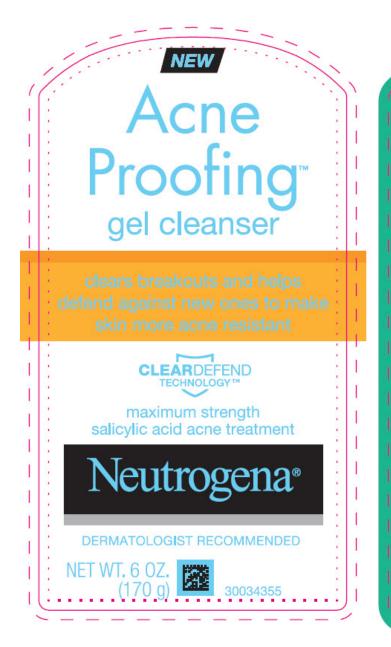
maximum strength salicylic acid acne treatment

Neutrogena[®]

DERMATOLOGIST RECOMMENDED

NET WT. 6 OZ. (170 g)

30034355





NEUTROGENA ACNE PROOFING CLEANSER

salicylic acid gel

Product Information

Product Type HUMAN OTC DRUG Item Code (Source) NDC:69968-0110

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)	
Glycerin (UNII: PDC6A3C0OX)	
Cocamidopropyl Hydroxysultaine (UNII: 62V75NI93W)	
Sodium Methyl Cocoyl Taurate (UNII: JVL98CG53G)	
Sodium Chloride (UNII: 451W47IQ8X)	
PEG-120 Methyl Glucose Dioleate (UNII: YM0K64F20V)	
Benzalkonium Chloride (UNII: F5UM2KM3W7)	
Sodium Hydroxide (UNII: 55X04QC32I)	
COCAMIDO PRO PYL PRO PYLENE GLYCOL-DIMO NIUM CHLO RIDE PHO SPHATE (UNII: H2KVQ74JM4)	
C12-15 Alkyl Lactate (UNII: GC844VRD7E)	
Panthenol (UNII: WV9CM0O67Z)	
EDETATE DISODIUM ANHYDROUS (UNII: 8 NLQ36 F6 MM)	

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l	# Item Code	Package Description	Marketing Start Date	Marketing End Date		
l	1 NDC:69968-0110-1	14 g in 1 TUBE; Type 0: Not a Combination Product	10 / 0 1 / 2 0 1 7			
l	2 NDC:69968-0110-6	170 g in 1 BOTTLE; Type 0: Not a Combination Product	10 /0 1/20 17			

Marketing Inform	Marketing Information					
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date			
OTC MONOGRAPH FINAL	part333D	10 / 0 1 / 2 0 1 7				

Labeler - Johnson & Johnson Consumer Inc. (002347102)

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