NEUTROGENA SKIN POLISHING ACNE CLEANSER- salicylic acid cream 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® SKIN POLISHING ACNE CLEANSER

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

Wet face. Squeeze into hands. Apply to face and massage gently. Rinse thoroughly. If excessive drying or peeling occurs, reduce application to every other day.

Other information

Sunburn Alert

This product contains an alpha hydroxy acid (AHA) that may increase your skin's sensitivity to the sun and particularly the possibility of sunburn. Use a sunscreen, wear protective clothing and limit sun exposure while using this product and for a week afterwards. Store at room temperature.

Inactive ingredients

Water, Cetyl Alcohol, PPG-15 Stearyl Ether, Glycolic Acid, Glycerin, Polysorbate 60, Steareth-21, C12-15 Alkyl Lactate, Cetyl Lactate, Potassium Cetyl Phosphate, Sodium Hydroxide, Xanthan Gum, Cocamidopropyl PG-Dimonium Chloride Phosphate, Disodium EDTA, Menthol, Fragrance, Benzalkonium Chloride, Panthenol

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 - 150 mL Tube Label

NEW FORMULA

NEUTROGENA® SKIN POLISHING ACNE CLEANSER

salicylic acid acne treatment

5.1 FL. OZ. (150 mL)



salicylic acid cream

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:69968-0320	
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)		
Cetyl Alcohol (UNII: 936JST6JCN)		
PPG-15 Stearyl Ether (UNII: 1II18 XLS1L)		
Glycolic Acid (UNII: 0WT12SX38S)		
Glycerin (UNII: PDC6A3C0OX)		
Polysorbate 60 (UNII: CAL22UVI4M)		
Steareth-21 (UNII: 53J3F32P58)		
Sodium Hydroxide (UNII: 55X04QC32I)		
C12-15 Alkyl Lactate (UNII: GC844VRD7E)		
Benzalkonium Chloride (UNII: F5UM2KM3W7)		
Edetate Disodium Anhydrous (UNII: 8NLQ36F6MM)		
Cetyl Lactate (UNII: A7EVH2RK4O)		
Potassium Cetyl Phosphate (UNII: 03KCY6P7UT)		
Cocamidopropyl Propylene Glycol-Dimonium Chloride Phosphate (UNII: H2KVQ74JM4)		
Menthol, Unspecified Form (UNII: L7T10 EIP3A)		
Panthenol (UNII: WV9CM0O67Z)		
Xanthan Gum (UNII: TTV12P4NEE)		

l	Packaging			
l	# Item Code	Package Description	Marketing Start Date	Marketing End Date
l	1 NDC:69968-0320-5	150 mL in 1 TUBE; Type 0: Not a Combination Product	12/0 1/20 16	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC MONOGRAPH FINAL	part333D	12/0 1/20 16		

Labeler - Johnson & Johnson Consumer Inc. (002347102)