

ACNE CONTROL SUPERCLEAN BODY WASH AHA BHA FORMULA- salicylic acid gel

Bentley Laboratories, LLC

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.

Geologie Acne Control Superclean Body Wash AHA+ BHA Formula

DRUG FACTS

Active Ingredients..... Salicylic Acid 2%

Purpose Acne Treatment

Uses: For the treatment of acne. Penetrates pores to eliminate most acne blemishes, whiteheads & blackheads. Helps prevent new acne blemishes, whiteheads & blackheads.

Directions: Lather with water and massage onto affected areas. Let sit for 30 to 60 seconds to maximize results. Rinse clean with warm water.

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.

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

INACTIVE INGREDIENTS: WATER/AQUA/EAU, SODIUM C14-16 OLEFIN SULFONATE, COCAMIDOPROPYL HYDROXYSULTAINE, DISTEARETH-75 IPDI, PROPANEDIOL, SODIUM LAUROAMPHOACETATE, GLYCERIN, GLYCOLIC ACID, COCAMIDOPROPYL BETAINE, PHENOXYETHANOL, SODIUM CHLORIDE, GLUCO-NOLACTONE, SODIUM HYDROXIDE, SODIUM GLUCONATE

Geologie Clear System Package Labeling

Geologie

CLEAR SYSTEM

Acne Control

Superclean

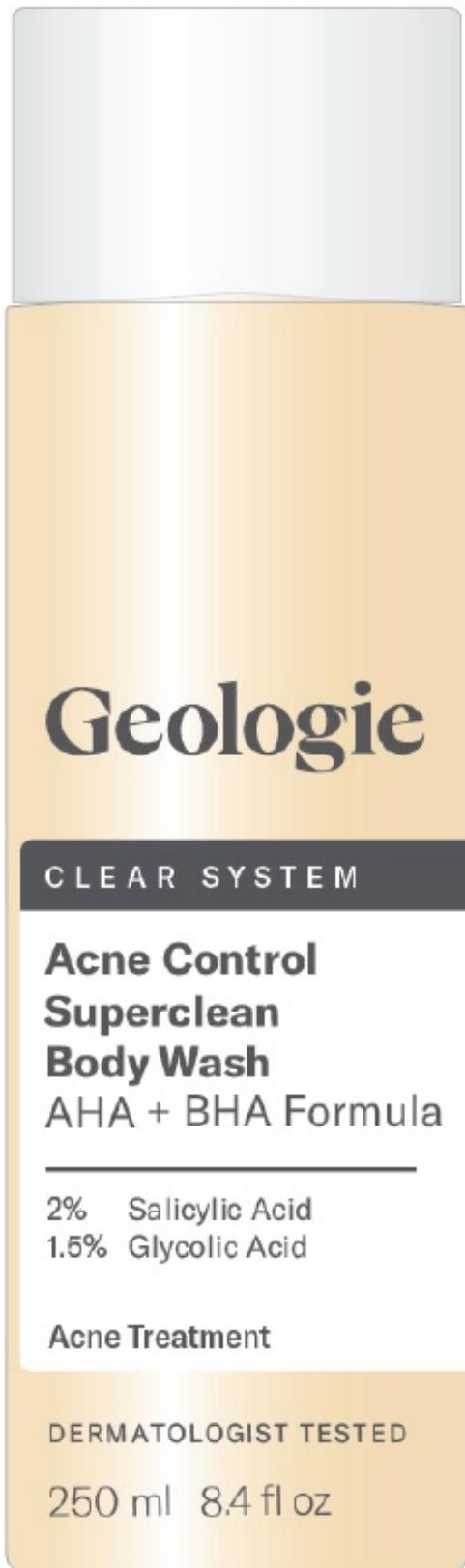
Body Wash

250 mL 8.4 fl oz

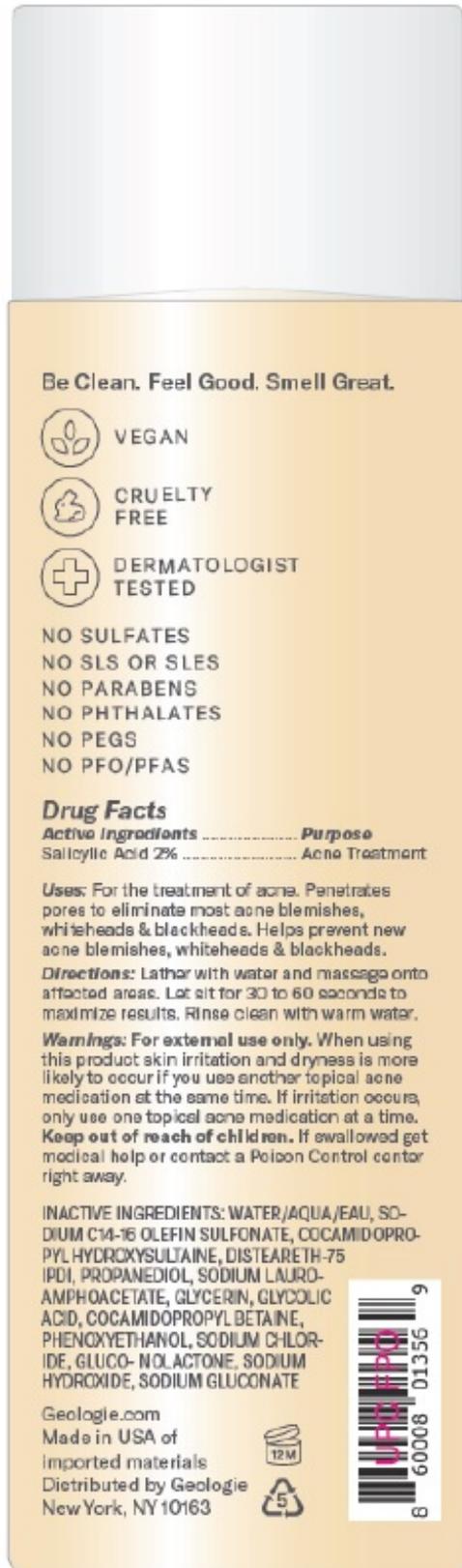
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FRONT



BACK

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ACNE CONTROL SUPERCLEAN BODY WASH AHA BHA FORMULA
salicylic acid gel

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:54111-170
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: 0414PZ4LPZ) (SALICYLIC ACID - UNII:0414PZ4LPZ)	SALICYLIC ACID	2.0 g in 100 mL

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM C14-16 OLEFIN SULFONATE (UNII: O9W3D3YF5U)	
COCAMIDOPROPYL HYDROXYSULTAINE (UNII: 62V75NI93W)	
DISTEARETH-75 ISOPHORONE DIISOCYANATE (UNII: 5365FJ30SC)	
PROPANEDIOL (UNII: 5965N8W85T)	
SODIUM LAUROAMPHOACETATE (UNII: SLK428451L)	
GLYCERIN (UNII: PDC6A3C0OX)	
GLYCOLIC ACID (UNII: 0WT12SX38S)	
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
SODIUM CHLORIDE (UNII: 451W47IQ8X)	
GLUCONOLACTONE (UNII: WQ29KQ9POT)	
SODIUM HYDROXIDE (UNII: 55X04QC32I)	
SODIUM GLUCONATE (UNII: R6Q3791S76)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:54111-170-50	250 mL in 1 BOTTLE; Type 0: Not a Combination Product	03/01/2023	

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	03/01/2023	

Labeler - Bentley Laboratories, LLC (068351753)

Registrant - Geologie (081158549)