

SALICYLIC FOAM CLEANSER 20- salicylic foam cleanser 20 solution

Gazebo Wellness SKIN LLC

Skin Beauty Solutions Salicylic Foam, Removes dirt, dead skin, and oil from pores that can cause active acne.

Directions: Apply cleanser onto wet face and neck, massage and rinse with cool water, Use 3-5 times a week.

Inactive ingredients: Aqua (Water), Decyl Glucoside, Cocamidopropyl Betaine, Kosher Vegetable Glycerin, Phenoxyethanol, Xanthan Gum.

Active ingredient: Salicylic Acid.

Directions: Apply cleanser onto wet face and neck, massage and rinse with cool water, Use 3-5 times a week.

keep out of reach of children

Salicylic Foam Cleanser
20%

Removes dirt, dead skin, and oil from pores.

Distributed By: Skin Beauty Solutions Woodbury MN 55129

Content of labeling

Skin Beauty Solutions Salicylic Foam,
Removes dirt, dead skin, and oil from
pores that can cause active acne.



Ingredients: Salicylic Acid, Aqua (Water),
Decyl Glucoside, Cocamidopropyl Betaine,
Kosher Vegetable Glycerin,
Phenoxyethanol, Xanthan Gum
(Polysaccharide Gum), Tetrasodium Edta,
d-Calcium Pantothenate (Pro-Vitamin B5).

Salicylic Foam Cleanser

20%

Directions: Apply cleanser onto wet
face and neck, massage and rinse
with cool water, Use 3-5 times a week.

**Removes dirt,
dead skin, and
oil from pores.**

Distributed By:
Skin Beauty Solutions
Woodbury MN 55129

7.5oz / 225 ml



SALICYLIC FOAM CLEANSER 20

salicylic foam cleanser 20 solution

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:84785-0015
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	22.5 mg in 225 mL

Inactive Ingredients	
Ingredient Name	Strength
COCAMIDOPROPYL BETAINE (UNII: 5OCF3O11KX)	
XANTHAN GUM (UNII: TTV12P4NEE)	
GLYCERIN (UNII: PDC6A3C0OX)	
PHENOXYETHANOL (UNII: HIE492ZZ3T)	
WATER (UNII: 059QF0KO0R)	
DECYL GLUCOSIDE (UNII: Z17H97EA6Y)	

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:84785-0015-1	225 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/01/2024	

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M016	10/01/2024	

Labeler - Gazebo Wellness SKIN LLC (119609953)

Registrant - Gazebo Wellness SKIN LLC (119609953)

Establishment			
Name	Address	ID/FEI	Business Operations
Gazebo Wellness SKIN LLC		119609953	manufacture(84785-0015)

Revised: 12/2024

Gazebo Wellness SKIN LLC