SOVEREIGN SILVER FOAMING FACIAL CLEANSER- salicylic acid/acne treatment gel gel

Natural Immunogenics Corporation

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.

SOVEREIGN SILVER Foaming Facial Cleanser Exfoliating Wash 1% Salicylic Acid/Acne treatment Gel

Salicylic Acid 1%

Acne treatment

USES for treatment of acne.Penetrates pores to eliminate most acne blemishes, blackheads and whiteheads.

Warnings

For external use only.

When using this product. skin irritation and dryness is more likely to occcur if you use another topical acne medication at the same time. If irritation occurs, only use one topical acne medicarion a a time.

Stop use and ask a doctor if. skin irritation occurs or gets worse.

Keep out of reach of children. If swallowed, get medical help or contact Poison Control Center right away.

Directions Take a quarter sized amount of the product in wet hands and work into a fine lather. Apply to wet face and massage for 60 seconds. Rinse with warm water and pat the skin dry. Use twice daily.

Inactive Ingredients Capryloyl/Caproyl Methyl Glucamide, Citric Acid, Coco-Glucoside, Cocos Nucifera(Coconut)Fruit Extract, Glycerin, Hamamelis Virginiana (Witch Hazel) Leaf Extract, Jojoba Esters, Lactobacillus, Lactobacillus Ferment, Lauroyl/Myristoyl Methyl Glucamide, Propanediol, Salix Alba(Willow) Bark Extract, Silver, Sodium Citrate, Sunfloweroyl Methylglucamide, Water, Xanthan Gum.



salicylic acid/acne treatment gel gel

Product Information				
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:52166-016	
Route of Administration	TOPICAL			

Active Ingredient/Active Moiety			
Ingredient Name	Basis of Strength	Strength	
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	1 g in 59 mL	

Inactive Ingredients				
Ingredient Name	Strength			
GLYCERIN (UNII: PDC6A3C0OX)				
LIMOSILACTOBACILLUS FERMENTUM (UNII: 2C1F12C6AP)				
XANTHAN GUM (UNII: TTV12P4NEE)				
PROPANEDIOL (UNII: 5965N8W85T)				
SILVER (UNII: 3M4G523W1G)				
COCONUT JUICE (UNII: AMN6S4M09G)				
LAUROYL/MYRISTOYL METHYL GLUCAMIDE (UNII: SC667B999P)				
CAPRYLOYL/CAPROYL METHYL GLUCAMIDE (UNII: 0451R360HR)				
COCO GLUCOSIDE (UNII: ICS790225B)				
SALIX ALBA BARK (UNII: 205MXS71H7)				
SODIUM CITRATE (UNII: 1Q73Q2JULR)				
SUNFLOWER OIL (UNII: 3W1JG795YI)				
CITRIC ACID MONOHYDRATE (UNII: 2968PHW8QP)				
HYDROGENATED JOJOBA OIL/JOJOBA OIL, RANDOMIZED (IODINE VALUE 64-70) (UNII: 96YYQ5TK1K)				
HAMAMELIS VIRGINIANA TOP (UNII: UDA30A2JJY)				

Packaging					
	# Item Code Package Description		Marketing Start Date	Marketing End Date	
		NDC:52166-016- 01	59 mL in 1 BOTTLE; Type 0: Not a Combination Product	07/24/2023	

Marketing Information				
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date	
OTC monograph final	part333D	07/24/2023		

Labeler - Natural Immunogenics Corporation (048744085)

Establishment

Name	Address	ID/FEI	Business Operations
Inspec Solutions		081030372	manufacture(52166-016)

Revised: 9/2023 Natural Immunogenics Corporation