




Inactive ingredients water, caprylic/capric triglyceride, glycerin, sodium polyacrylate, hydroxyethyl acrylate/sodium acryloyldimethyl taurate copolymer, niacinamide, phenoxyethanol, xanthan gum, polyacrylate crosspolymer-6, allantoin, ethylhexylglycerin, zinc pca, sorbitan isostearate, t-butyl alcohol

Uses • for the treatment of acne

Active ingredients	Purpose
Salicylic acid 2%	Acne Treatment

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

Directions • cleanse skin thoroughly before applying serum • apply serum to face or affected areas twice daily
• because excessive drying may occur, start with one application daily

Warnings
For external use only. Do not use if allergic to salicylic acid. Using other topical acne treatments at the same time or immediately after following use of this product may increase dryness or irritation. If this occurs, only one medication should be used unless directed by a doctor.

ULTA ALL CLEAR SERUM			
salicylic acid gel			
Product Information			
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:79959-011
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
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)			1 mg in 1 g
TERT-BUTYL ALCOHOL (UNII: MD83SFE959)			0.042 mg in 1 g
PHENOXYETHANOL (UNII: HIE492ZZ3T)			9 mg in 1 g
SODIUM POLYACRYLATE (2500000 MW) (UNII: 05I15JN12J)			20 mg in 1 g
			0.5

XANTHAN GUM (UNII: TTV12P4NEE)	8.5 mg in 1 g
WATER (UNII: 059QF0KO0R)	766.2865 mg in 1 g
NIACINAMIDE (UNII: 25X51I8RD4)	10 mg in 1 g
MEDIUM-CHAIN TRIGLYCERIDES (UNII: C9H2L21V7U)	84.6 mg in 1 g
AMMONIUM ACRYLOYLDIMETHYLTaurate, DIMETHYLACRYLAMIDE, LAURYL METHACRYLATE AND LAURETH-4 METHACRYLATE COPOLYMER, TRIMETHYLOLPROPANE TRIACRYLATE CROSSLINKED (45000 MPA.S) (UNII: Q7UI015FF9)	1.995 mg in 1 g
ALLANTOIN (UNII: 344S277G0Z)	1 mg in 1 g
ZINC PIDOLATE (UNII: C32PQ86DH4)	1 mg in 1 g
SORBITAN ISOSTEARATE (UNII: 01S2G2C1E4)	0.814 mg in 1 g
GLYCERIN (UNII: PDC6A3C0OX)	63.1825 mg in 1 g
HYDROXYETHYL ACRYLATE/SODIUM ACRYLOYLDIMETHYL TAURATE COPOLYMER (100000 MPA.S AT 1.5%) (UNII: 86FQE96TZ4)	12.58 mg in 1 g

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:79959-011-01	1 in 1 CARTON	06/01/2024	
1		27.168 g in 1 TUBE; Type 0: Not a Combination Product		

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M006	06/01/2024	

Labeler - APR BEAUTY GROUP INC (204198006)

Establishment

Name	Address	ID/FEI	Business Operations
APR BEAUTY GROUP INC		204198006	manufacture(79959-011) , label(79959-011) , pack(79959-011)