

TULA ACNE CLEARING AND TONE CORRECTING GEL- tula acne clearing and tone correcting gel liquid

Tula Life 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.

DRUG FACTS

Active Ingredients

Salicylic Acid 2.0%

Purpose

Acne Treatment

Uses

- treats acne
- Helps prevent the development or new acne blemishes

Warnings

For external use only

Using other topical medications at the same time or immediately following use of this product may increase dryness or irritation of the skin. If this occurs, only one medication should be used unless directed by a doctor.

When using this product

- avoid contact with the eyes
- if contact occurs, rinse thoroughly with water

Keep out of reach of children

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

Directions

- clean the skin thoroughly before applying this product
- cover the entire affected area with a thin layer one to three times daily
- because excessive drying of the skin may occur, start with

one application daily, then gradually increase to two or three times daily if needed or as directed by a doctor

- if bothersome dryness or peeling occurs, reduce application to once a day or every other day

Inactive ingredients

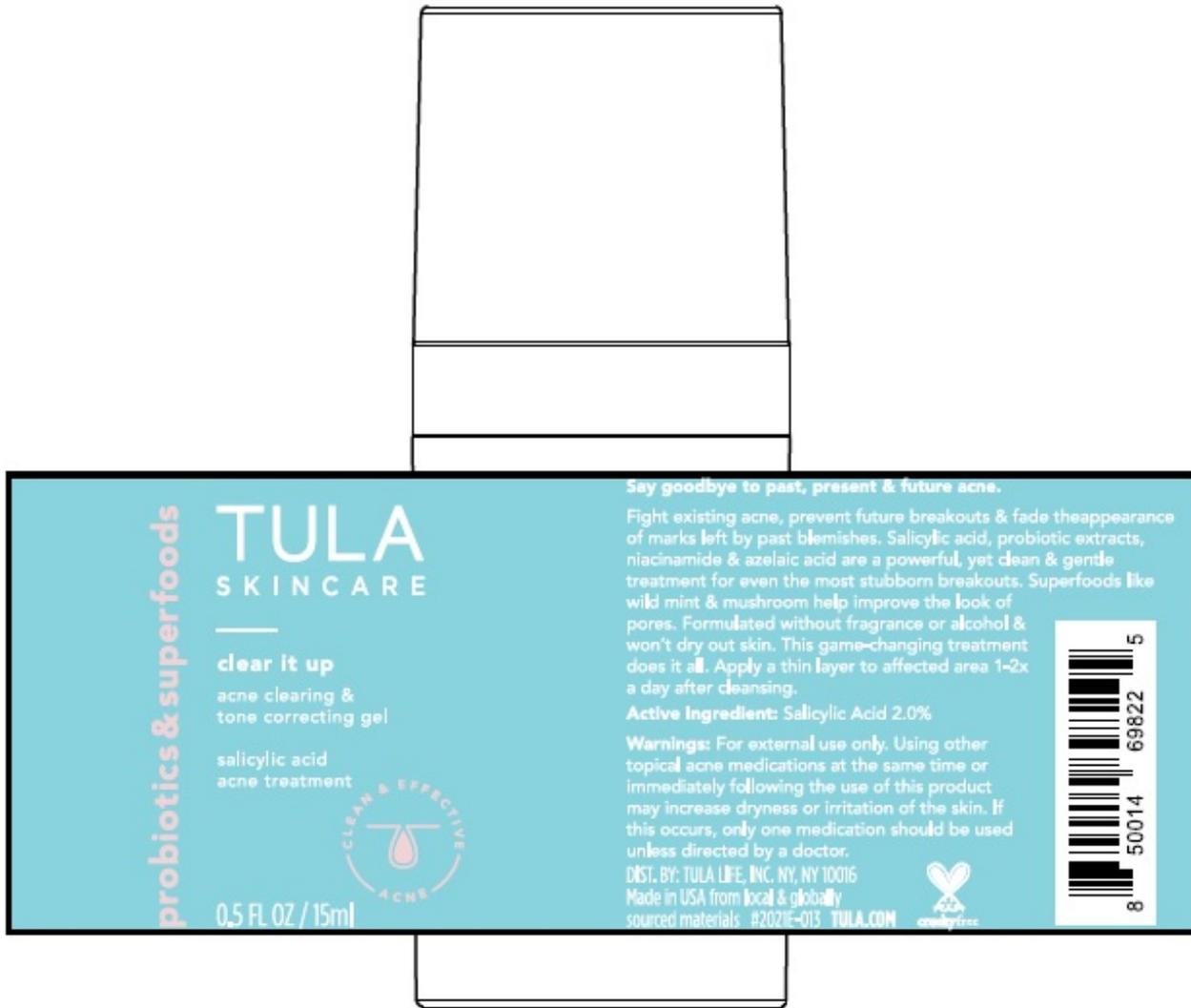
Water/Aqua/Eau, Sodium Polyacrylate, Sodium PCA, Glycerin, Polyacrylate Crosspolymer-6, Azelaic Acid, Lactococcus Ferment Lysate, Zinc PCA, Epilobium Angustifolium Flower/Leaf/Stem Extract, Niacinamide, Bisabolol, Allantoin, Sodium Hyaluronate, Lactic Acid, Sodium Benzoate, Sodium Chloride, Sodium Metabisulfite, Ethylhexylglycerol, Phenoxyethanol

Questions or Comments?

1-844-545-1236

Principal Display Panel

30 ml Package



TULA ACNE CLEARING AND TONE CORRECTING GEL

tula acne clearing and tone correcting gel liquid

Product Information

Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:72296-010
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
SALICYLIC ACID (UNII: O414PZ4LPZ) (SALICYLIC ACID - UNII:O414PZ4LPZ)	SALICYLIC ACID	2 g in 100 g

Inactive Ingredients

Ingredient Name	Strength
WATER (UNII: 059QF0KO0R)	
SODIUM POLYACRYLATE (8000 MW) (UNII: 285CYO341L)	
SODIUM PYRROLIDONE CARBOXYLATE (UNII: 469OTG57A2)	

GLYCERIN (UNII: PDC6A3C0OX)
AZELAIC ACID (UNII: F2VW3D43YT)
LACTOCOCCUS LACTIS (UNII: F1A0PSN10V)
ZINC (UNII: J41CSQ7QDS)
EPILOBIUM ANGUSTIFOLIUM LEAF (UNII: 7NV86426N2)
NIACINAMIDE (UNII: 25X51I8RD4)
.ALPHA.-BISABOLOL, (+)- (UNII: 105S6I733Z)
ALLANTOIN (UNII: 344S277G0Z)
LACTIC ACID (UNII: 33X04XA5AT)
SODIUM BENZOATE (UNII: OJ245FE5EU)
SODIUM CHLORIDE (UNII: 451W47IQ8X)
SODIUM METABISULFITE (UNII: 4VON5FNS3C)
ETHYLHEXYLGLYCERIN (UNII: 147D247K3P)
PHENOXYETHANOL (UNII: HIE492ZZ3T)
HYALURONATE SODIUM (UNII: YSE9PPT4TH)

Packaging				
#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:72296-010-30	1 in 1 CARTON	12/26/2022	
1		30 g in 1 TUBE; Type 0: Not a Combination Product		
2	NDC:72296-010-15	1 in 1 CARTON	12/26/2022	
2		15 g in 1 BOTTLE, PLASTIC; Type 0: Not a Combination Product		

Marketing Information			
Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC monograph final	part333D	12/26/2022	

Labeler - Tula Life LLC (080051358)