BIOTIN- biotin liquid
Shenzhen yuehaotong Technology Co.,Ltd

Salicylic acid 2%

wart remover

Apply twice daly(moring and night as a man and once as a woman.Afer apicaion, proeed with an acive masage for 3-5 minutes unthe product is fuy absorbed

Forevemaluseony Avoid coniac wth eves li coniac ocus nse ihorouthly wt waier Keep out ofreach oi chidrenifvou ale preanant.nursing. otiaveanyunderving medca condion.consutyou healthcare provider before using the product

For external use only.

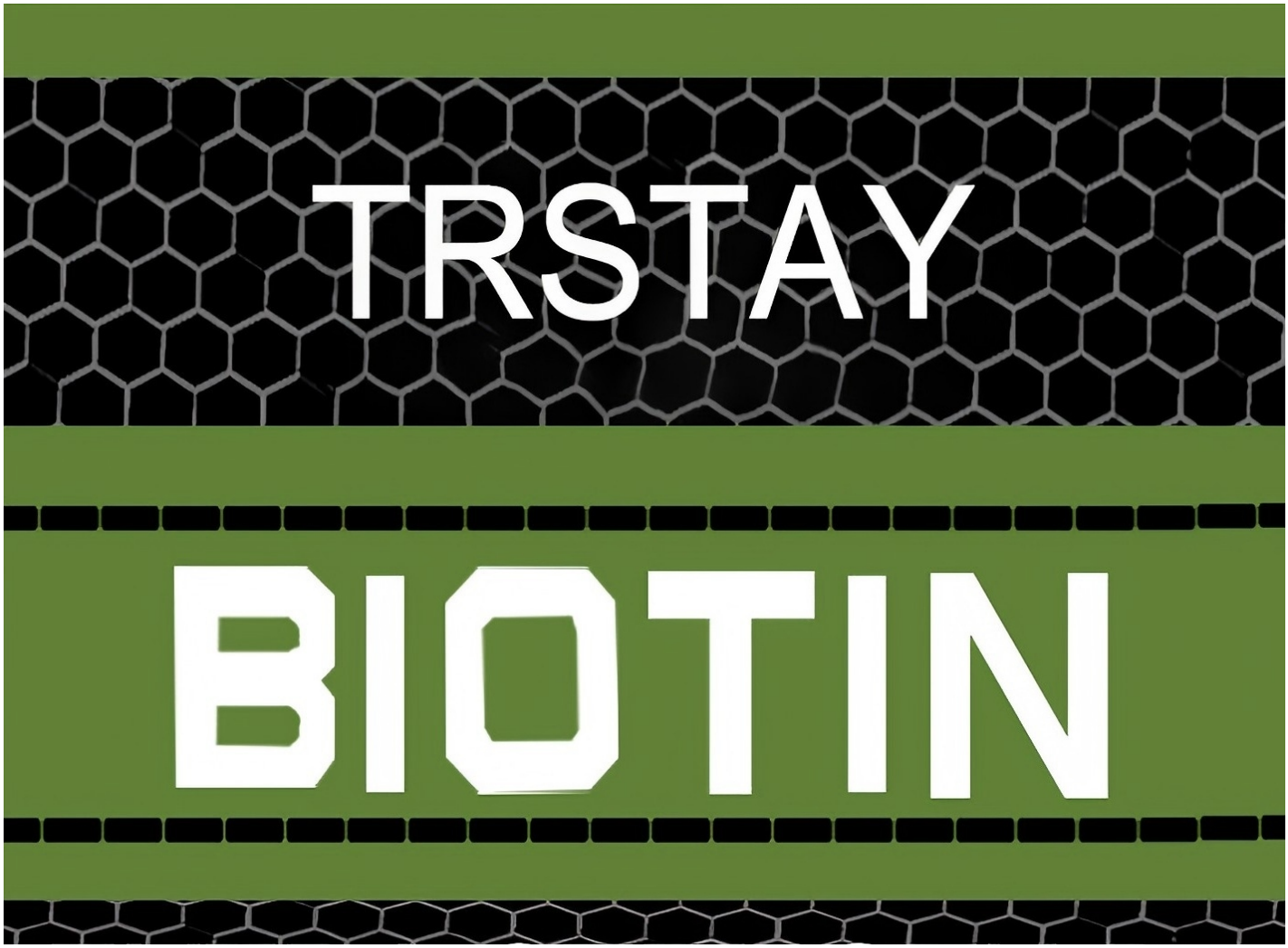
Discontinue use if any irritation or adverse reaction occurs

When using this product, avoid contact with eyes. If contact occurs, rinse thoroughly with waterDiscontinue use if any irritation or adverse reaction occurs.

on broken or irritated scalp

If sxralowed, get medical help or contact a Poison Conirol Cenler ngiht away.

Acrylates copolymer, Ethyl acetate



BIOTIN

biotin liquid

Product Information			
Product Type	HUMAN OTC DRUG	**Item Code (Source)**	NDC:87492-002
Route of Administration	TOPICAL		

Active Ingredient/Active Moiety		
Ingredient Name	**Basis of Strength**	**Strength**
MINOXIDIL (UNII: 5965120SH1) (MINOXIDIL - UNII:5965120SH1)	MINOXIDIL	5 mg in 100 mL

Inactive Ingredients	
Ingredient Name	**Strength**
SESAME OIL (UNII: QX10HYY4QV)	

CHLORPHENESIN (UNII: I670DAL4SZ)	
PALMITOYL TETRAPEPTIDE-7 (UNII: Q41S464P1R)	
PANTOTHENIC ACID (UNII: 19F5HK2737)	
PALMITOYL TRIPEPTIDE-1 (UNII: RV743D216M)	

Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:87492-002-01	5 mL in 1 BOX; Type 0: Not a Combination Product	03/07/2026	03/07/2031
2	NDC:87492-002-02	15 mL in 1 BOX; Type 0: Not a Combination Product	03/07/2026	03/07/2031
3	NDC:87492-002-03	30 mL in 1 BOX; Type 0: Not a Combination Product	03/07/2026	03/07/2031
4	NDC:87492-002-04	50 mL in 1 BOX; Type 0: Not a Combination Product	03/07/2026	03/07/2031
5	NDC:87492-002-05	100 mL in 1 BOX; Type 0: Not a Combination Product	03/07/2026	03/07/2031

Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
OTC Monograph Drug	M032	03/07/2026	03/07/2031

Labeler - Shenzhen yuehaotong Technology Co.,Ltd (602067669)

Establishment

Name	Address	ID/FEI	Business Operations
Shenzhen yuehaotong Technology Co.,Ltd		602067669	manufacture(87492-002)

Revised: 3/2026 Shenzhen yuehaotong Technology Co.,Ltd