

**5% MINOXIDIL SPRAY.- 5% minoxidil spray spray**  
**Consilii LLC**

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**Active Ingredient**

Minoxidil 5%

**Purpose**

Hair Regrowth Treatment

**Use**

to regrow hair on the top of the scalp

**Warnings**

For external use only

Keep away from fire and flame

Avoid contact with eyes

**Do not use**

You have no family history of hair loss, hair loss is sudden and/or patchy  
you are under 18 years of age. Do not use it on babies and children

**When Using**

Do not apply on other parts of the body

void contact with eyes. In case of accidental contact, rinse eyes with a large amount of cool tap water

It takes time to regrow hair. You may need to use this product 2 times a day for a least 4 months before you see results, The amount of hair regrowth is different for each person

**Stop Use**

chest pain, rapid heart beat, faintness, or dizziness occurs

sudden, unexplained weight gain occurs

your hands or feet swell

scalp irritation or redness occurs

**Ask Doctor**

When Pregnant or breast-feeding

**Keep Out Of Reach Of Children**

If swallowed, get medical help or contact a Poison Control Center right away. (1-800-222-1222)

**Directions**

Apply 1ml(5 Sprays) twice a day.Once in the morning and another time in the evening before bed

Spray directly on top of the scalp in the area to be treated

Discontinuing use may result in hair loss

**Other information**

Before use, read all information on the carton

Store at controlled room temperature 20 to 25 C (68 to 77°F)

**Inactive ingredients**

Deionized Water,Propylene Glycol,Ethyl Alcohol,Potassium Sorbate,GABA

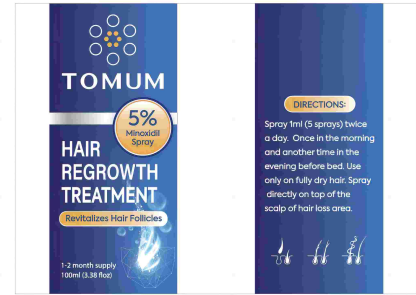
**Questions**

Tomumcs@gmail.com

**PRINCIPAL DISPLAY PANEL**

Size / Package size: 4.4\*4.4\*15.2cm

Bottle label size: 11.6\*8.4cm



## 5% MINOXIDIL SPRAY.

5% minoxidil spray spray

### Product Information

<b>Product Type</b>	HUMAN OTC DRUG	<b>Item Code (Source)</b>	NDC:83299-014
<b>Route of Administration</b>	TOPICAL		

### Active Ingredient/Active Moiety

Ingredient Name	Basis of Strength	Strength
MINOXIDIL (UNII: 5965120SH1) (MINOXIDIL - UNII:5965120SH1)	MINOXIDIL	5 g in 100 mL

### Inactive Ingredients

Ingredient Name	Strength
ALCOHOL (UNII: 3K9958V90M)	
POTASSIUM SORBATE (UNII: 1VPU26JZZ4)	
WATER (UNII: 059QF0KO0R)	
PROPYLENE GLYCOL (UNII: 6DC9Q167V3)	
.GAMMA.-AMINO BUTYRIC ACID (UNII: 2ACZ6IPC6I)	

## Packaging

#	Item Code	Package Description	Marketing Start Date	Marketing End Date
1	NDC:83299-014-01	100 mL in 1 BOTTLE; Type 0: Not a Combination Product	10/07/2023	

## Marketing Information

Marketing Category	Application Number or Monograph Citation	Marketing Start Date	Marketing End Date
ANDA	ANDA075357	10/07/2023	

**Labeler** - Consilii LLC (118891890)

## Establishment

Name	Address	ID/FEI	Business Operations
Consilii LLC		118891890	label(83299-014) , manufacture(83299-014)

Revised: 10/2023

Consilii LLC