



Magnum Compounding LLC
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**LET'S DO THIS
TOGETHER**

COMPOUNDED DRUG INFORMATION

Product Name: Minoxidil 5% Finasteride 0.01% Topical Solution

Formulation Type: Non-Sterile Compounded Preparation (Dermatologic, Alcohol-Free, PPG-Free)

Dispensed Volume: 30 mL

ACTIVE INGREDIENTS (per mL)

Ingredients	Concentration	Function
Minoxidil	5%	Helps support hair growth by promoting hair follicle activity and prolonging the hair cycle's growth phase.
Finasteride	0.01%	Inhibits the enzyme 5-alpha-reductase, reducing the conversion of testosterone to dihydrotestosterone (DHT). DHT contributes to progressive shrinking of susceptible hair follicles in androgenetic alopecia.

DESCRIPTION: Minoxidil 5% Finasteride 0.01% Topical Solution is a compounded medication applied directly to the scalp. It contains two active ingredients that work through different mechanisms to help manage certain types of hair loss. This medication is compounded specifically for an individual patient pursuant to a prescription. This formulation also contains excipients used to dissolve and deliver the active ingredients, including citric acid, ethoxy diglycol, dimethyl sulfoxide (DMSO), and propanediol. This is a compounded medication. Compounded medications are not FDA-approved, and FDA does not review compounded products for safety, effectiveness, or quality before they are marketed.

INDICATIONS

- This medication may be prescribed for the treatment of androgenetic alopecia (pattern hair loss) and other hair-loss conditions as determined by your healthcare provider. Hair growth occurs slowly. Improvement may require several months of consistent treatment, and individual results vary.
- Continued treatment is generally necessary to maintain the benefit. Hair loss may gradually resume after treatment is discontinued.

DIRECTIONS FOR USE

Use this medication exactly as prescribed by your healthcare provider and as stated on your prescription label.

If you have any questions or concerns regarding your medication, please do not hesitate to contact us.



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- Apply the prescribed amount directly to the affected areas of the scalp, not primarily to the hair.
- Apply to a clean, dry scalp unless otherwise directed.
- Gently spread the solution over the treatment area.
- Allow the medication to dry completely after application.
- Wash your hands thoroughly with soap and water immediately after applying.
- Avoid contact with the eyes, mouth, face, broken skin, or other unintended areas.
- Do not apply more medication or use it more frequently than prescribed. Using more does not necessarily improve results and may increase the risk of side effects.
- Do not share this medication with anyone else.

If the medication accidentally gets into your eyes, rinse thoroughly with water.

WHAT TO EXPECT?

Hair regrowth is gradual. Some patients may notice an initial temporary increase in hair shedding after beginning minoxidil therapy as follicles transition through the hair-growth cycle. Visible improvement may take several months. Response varies from person to person, and treatment does not produce hair regrowth in everyone. Do not discontinue treatment solely because you do not see immediate improvement unless instructed by your healthcare provider.

POTENTIAL SIDE EFFECTS

Common side effects may include:

- Scalp itching
- Redness or irritation
- Dryness or flaking
- Burning or stinging
- Changes in hair texture
- Unwanted hair growth on the face or other areas exposed to the medication

Contact your healthcare provider or pharmacist if irritation becomes persistent or severe.

IMPORTANT FINASTERIDE INFORMATION

Although finasteride is applied topically, some medication can be absorbed through the skin and enter the bloodstream.

Possible systemic adverse effects associated with finasteride may include:

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- Decreased sexual desire (libido)
- Erectile dysfunction
- Changes in ejaculation
- Testicular discomfort
- Breast tenderness or enlargement
- Mood changes, including depression or anxiety
- Fatigue or other persistent symptoms

Contact your healthcare provider if you develop concerning or persistent symptoms.

The FDA has received reports of systemic adverse effects in patients using compounded topical finasteride products. There is currently no FDA-approved topical finasteride product in the United States.

IMPORTANT PREGNANCY WARNING

Finasteride can cause harm to a developing male fetus.

Women who are pregnant or who may become pregnant should avoid direct skin contact with this medication.

After each application:

- Wash your hands thoroughly.
- Allow the scalp to dry completely.
- Avoid transferring wet medication to another person through direct contact.
- Prevent pregnant individuals or individuals who may become pregnant from touching the medication or the treated area before it has completely dried.

If accidental exposure occurs during pregnancy, contact a healthcare provider.

CONTACT YOUR HEALTHCARE PROVIDER

Stop using the medication and seek medical advice if you experience:

- Chest pain
- Rapid or irregular heartbeat
- Fainting or significant dizziness
- Swelling of the hands or feet
- Sudden unexplained weight gain
- Severe or persistent scalp irritation
- Significant mood or behavioral changes
- Persistent sexual side effects

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- Breast lump, pain, enlargement, or nipple discharge
- Signs of a serious allergic reaction

Seek immediate medical attention for severe symptoms.

WARNINGS AND PRECAUTIONS

Tell your healthcare provider before using this medication if you:

- Have heart disease or significant cardiovascular problems
- Have low blood pressure or experience frequent dizziness
- Have an irritated, infected, sunburned, or damaged scalp
- Are using other medications or medicated products on the scalp
- Have sudden, unexplained, or patchy hair loss
- Are pregnant, planning pregnancy, or breastfeeding

This medication is intended for external use only.

Keep out of reach of children and pets.

Do not swallow.

MISSED DOSE: If you miss an application, apply the next dose at the regularly scheduled time. Do not double the amount to make up for a missed application.

STORAGE: Store according to the instructions on your prescription label. Keep the container tightly closed and protect it from excessive heat, moisture, and direct light unless otherwise directed by the pharmacy. Keep out of reach of children and pets.

LEGAL & SAFETY DISCLAIMER: This compounded medication is prepared by Magnum Compounding LLC pursuant to a valid prescription from a licensed healthcare provider for an individual patient when commercially available options are not suitable. Compounded medications have not been evaluated or approved by the U.S. Food and Drug Administration (FDA) for safety, efficacy, or quality and are prepared in accordance with applicable standards, including USP <795> Non-Sterile Compounding guidelines. Magnum Compounding LLC does not diagnose medical conditions or determine treatment plans. Patients should use this medication only as directed by their prescribing healthcare provider and under appropriate medical supervision. If any unexpected symptoms, adverse reactions, or signs of hypersensitivity occur, discontinue use and seek medical attention promptly. Patients should consult their healthcare provider with any questions regarding use, risks, precautions, or duration of therapy.

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REFERENCES

- U.S. Food and Drug Administration (FDA). *FDA Alerts Health Care Providers, Compounders and Consumers of Potential Risks Associated with Compounded Topical Finasteride Products*. April 22, 2025.
- U.S. Food and Drug Administration. *PROPECIA® (finasteride) Prescribing Information*.
- DailyMed. *Minoxidil Topical Solution 5% - Drug Label Information*.

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