A Randomized, Double-Blind, Placebo-Controlled, Parallel Group, Multi-dose Study to Evaluate the Efficacy and Safety of VDPHL01 in Female Subjects with Androgenetic Alopecia

Status: Recruiting

Eligibility Criteria

Sex: Female
Age Group: 18 years and over
This study is NOT accepting healthy volunteers
Inclusion Criteria:

- women - 18-65 years of age - mild to moderate Androgenetic Alopecia (AGA)

Exclusion Criteria:

- history of organ transplant - history of heart disease

Conditions & Interventions

Interventions:
Drug: Placebo, Drug: Placebo, Drug: VDPHL01 BID, Drug: VDPHL01 QD
Conditions:
Dermatology (Skin, Hair & Nails)
Keywords:
AGA, Androgenetic Alopecia, dermatology, women's health

More Information

Description: The research study will compare VDPHL01 with placebo (no active ingredient) to learn about the safety and how VDPHL01 works for women who have androgenetic alopecia (AGA). AGA is a genetic disorder caused by an excessive (too much) hair follicle response to androgens resulting in hair loss. VDPHL01 is an extended release (ER) oral (taken by mouth) form of minoxidil.
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