

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

10 Jun 2026

### Comparison of Efficacy of Topical Anagain 4% Solution Versus Topical Minoxidil 5% Solution in Androgenetic Alopecia

#### Protocol summary

##### Study aim

This study aims to compare the efficacy of topical Anagain 4% versus topical Minoxidil 5% in the treatment of androgenetic alopecia.

##### Design

Community based, parallel group, single blinded, RCT

##### Settings and conduct

In this study 38 patient with moderate to severe AGA are enrolled after explaining thoroughly the course and purpose of study. Patients are randomly assigned one of the two groups, one receiving topical Anagain 4% and the other receiving topical Minoxidil 5% solution twice daily for 6 months.

##### Participants/Inclusion and exclusion criteria

Inclusion criteria A. Clinical diagnosis of AGA B. Age range of 18 - 55 years, C. Patients who haven't received topical or systemic treatments for AGA within 3 months prior to the study. D. Patient who gave consent for the study Exclusion criteria A. Those with other causes of hair loss such as fungal infection such as taenia capitis, inflammatory or scarring alopecia, hyper-androgenism or other hormonal disorders B. Hypersensitive to the medications C. A history of severe systemic disease (renal, cardiovascular hepatic and lungs). D. Pregnant or breastfeeding patients

##### Intervention groups

Intervention group 1: Patients in the first group will be treated with Topical Anagain 4% (1 cc twice daily).  
Intervention group 2: Patients in the second group will receive topical Minoxidil 5% (1 cc twice daily) for 6 months.

##### Main outcome variables

Number of hairs, Change in photographic pictures and patient satisfaction score.

#### General information

##### Reason for update

##### Acronym

#### IRCT registration information

IRCT registration number: **IRCT20210823052264N11**

Registration date: **2024-09-29, 1403/07/08**

Registration timing: **registered\_while\_recruiting**

Last update: **2024-09-29, 1403/07/08**

Update count: **0**

#### Registration date

2024-09-29, 1403/07/08

#### Registrant information

##### Name

Najia Ahmed

##### Name of organization / entity

PNS shifa

##### Country

Pakistan

##### Phone

+92 81 2864092

##### Email address

najiaomer@yahoo.com

#### Recruitment status

**Recruitment complete**

#### Funding source

#### Expected recruitment start date

2024-07-01, 1403/04/11

#### Expected recruitment end date

2024-12-31, 1403/10/11

#### Actual recruitment start date

empty

#### Actual recruitment end date

empty

#### Trial completion date

empty

#### Scientific title

Comparison of Efficacy of Topical Anagain 4% Solution Versus Topical Minoxidil 5% Solution in Androgenetic Alopecia

**Public title**

Comparison of Efficacy of Topical Anagain 4% Solution Versus Topical Minoxidil 5% Solution in Androgenetic Alopecia

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Clinical diagnosis of AGA score 2-5 on Norwood-Hamilton Classification Age range of 18 - 55 years Patients who haven't received topical or systemic treatments for AGA within 3 months prior to the study. Patient who gave consent for the study.

**Exclusion criteria:**

Those with other causes of hair loss such as fungal infection such as taenia capitis, inflammatory or scarring alopecia, hyper-androgenism or other hormonal disorders Hypersensitive to the medications A history of severe systemic disease (renal, cardiovascular hepatic and lungs) Pregnant or breastfeeding patients

**Age**

From **18 years** old to **55 years** old

**Gender**

Both

**Phase**

3

**Groups that have been masked**

- Participant
- Data analyser

**Sample size**

Target sample size: **92**

**Randomization (investigator's opinion)**

Randomized

**Randomization description**

Randomization done by lottery method in which chits were kept in a jar and patient will be asked to pick one deciding their treatment regimen as per interventional group 1 and 2

**Blinding (investigator's opinion)**

Double blinded

**Blinding description**

Researcher who is assessing the response is blind and the patients are blinded. Participant/patient picks up a chit from the jar and decides the treatment method as per interventional group 1 and 2

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Institutional Ethical Review Board (IERB) certificate - CMH Quetta

**Street address**

CMH Quetta

**City**

Quetta

**Postal code**

08762

**Approval date**

2024-08-30, 1403/06/09

**Ethics committee reference number**

CMH QTA-IERB/33/2024

**Health conditions studied****1****Description of health condition studied**

Androgenetic Alopecia

**ICD-10 code**

L64.9

**ICD-10 code description**

Androgenic alopecia, unspecified

**Primary outcomes****1****Description**

Total hair density and total number of vellus and terminal hairs

**Timepoint**

before start of study and after 1,2,3 and 6 months after start of study.

**Method of measurement**

Photographic pictures

**2****Description**

Patient satisfaction score

**Timepoint**

After 6 months

**Method of measurement**

Questionnaire

**Secondary outcomes****1****Description**

Any side effect after applying minoxidil or anagain topically

**Timepoint**

1,2,3 and 6 months

**Method of measurement**

interviewing the patients and clinical examination

## Intervention groups

### 1

#### Description

Intervention group: Participants will be given Topical Anagain 4% solution. They will be shown the correct method of application of the drug which they will apply twice daily on the scalp, 12 hours apart for 6 months.

#### Category

Treatment - Drugs

### 2

#### Description

Control group: Participants will be given Topical Minoxidil 5% solution. They will be shown the correct method of application of the drug which they will apply twice daily on the scalp, 12 hours apart for 6 months.

#### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Combined Military Hospital Quetta

##### Full name of responsible person

Nosheen Abbas

##### Street address

CMH Quetta

##### City

Quetta

##### Postal code

08762

##### Phone

+92 335 0250058

##### Email

nausheenabbas1122@gmail.com

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Combined Military Hospital Quetta

##### Full name of responsible person

Nosheen Abbas

##### Street address

CMH Quetta

##### City

Quetta

##### Postal code

08762

##### Phone

+92 335 0250058

##### Email

nausheenabbas1122@gmail.com

#### Grant name

CMH Quetta

#### Grant code / Reference number

#### Is the source of funding the same sponsor organization/entity?

Yes

#### Title of funding source

Combined Military Hospital Quetta

#### Proportion provided by this source

100

#### Public or private sector

Public

#### Domestic or foreign origin

Domestic

#### Category of foreign source of funding

*empty*

#### Country of origin

#### Type of organization providing the funding

Academic

## Person responsible for general inquiries

#### Contact

##### Name of organization / entity

Combined Military Hospital Quetta

##### Full name of responsible person

Nosheen Abbas

##### Position

Registrar

##### Latest degree

Subspecialist

##### Other areas of specialty/work

Dermatology

##### Street address

CMH Quetta

##### City

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##### Province

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## Person responsible for scientific inquiries

#### Contact

##### Name of organization / entity

Combined Military Hospital Quetta

##### Full name of responsible person

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##### Latest degree

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nausheenabbas1122@gmail.com

## Person responsible for updating data

### Contact

**Name of organization / entity**  
Combined Military Hospital Quetta  
**Full name of responsible person**  
Nosheen Abbas  
**Position**  
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## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be a plan to make this available

### Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

### Statistical Analysis Plan

Undecided - It is not yet known if there will be a plan to make this available

### Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

### Clinical Study Report

Undecided - It is not yet known if there will be a plan to make this available

### Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

### Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available