

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

08 Jun 2026

### Comparison of the effects of Oral charcoal capsule with Aluminum Hydroxide Syrup on pruritus in hemodialytic patients in Gorgan 5th - Azar hospital.

#### Protocol summary

##### Summary

This study will conduct to compare oral Charcoal capsule with Aluminum Hydroxide Syrup effects on hemodialytic patients' pruritus in Gorgans' 5th Azar hospital. In this crossover clinical trail, 30 hemodialytic patients will select Gorgans' 5th Azar hospital via convenience sampling method. Then the subjects randomly will allocate to two groups (n= 15 per group). Inclusion criteria consist of: at least 2 weeks history of Uremic pruritus; Having Moderate or severe pruritus {base on measurement of Pruritus scale (MPS) or Visual Analog Scale (VAS)}; At least 6 months history of hemodialysis; Tree times in week and four hours for each session. Exclusion criteria : have dermal or non dermal causer pruritus. Each group receive oral charcoal capsule (6g daily in three doses) for two weeks as well as Aluminum Hydroxide syrup (30 cc; three times per day before each meal) for the same time (4 weeks intervention for groups). After 48 hours wash out period will replace treatments in two groups cross overly. Pruritus severity will be measured by visual analog scale (VAS) and measurement of pruritus scale (MPS).

#### General information

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT201107317164N1**  
Registration date: **2011-08-10, 1390/05/19**  
Registration timing: **retrospective**

Last update:

Update count: **0**

##### Registration date

2011-08-10, 1390/05/19

##### Registrant information

##### Name

Ali Abbasi

##### Name of organization / entity

Golestan medical Sciences university

##### Country

Iran (Islamic Republic of)

##### Phone

+98 17 1442 1664

##### Email address

abbasi\_a@goums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Vice chancellor for research of Golestan medical Sciences university

##### Expected recruitment start date

2007-06-12, 1386/03/22

##### Expected recruitment end date

2008-07-12, 1387/04/22

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Comparison of the effects of Oral charcoal capsule with Aluminum Hydroxide Syrup on pruritus in hemodialytic patients in Gorgan 5th - Azar hospital.

##### Public title

Effect of charcoal on uremic pruritus

##### Purpose

Treatment

##### Inclusion/Exclusion criteria

Inclusion criteria: At least 2 weeks history of Uremic pruritus; Having Moderate or severe pruritus {based on

measurement of Pruritus scale (MPS) or Visual Analog Scale (VAS)); At least 6 months history of hemodialysis; Tree time hemodialysis in week and four hours for each session. exclusion criteria: Having Dermal or non Dermal Causer Pruritus

**Age**

No age limit

**Gender**

Both

**Phase**

2-3

**Groups that have been masked**

No information

**Sample size**

Target sample size: 30

**Randomization (investigator's opinion)**

Randomized

**Randomization description****Blinding (investigator's opinion)**

Not blinded

**Blinding description****Placebo**

Not used

**Assignment**

Crossover

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

empty

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

Golestan medical Sciences university ethics committee

**Street address**

Golestan medical sciences and health services university, Shast kola street, Gorgan

**City**

Gorgan

**Postal code****Approval date**

2007-06-12, 1386/03/22

**Ethics committee reference number**

129299/35

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

Renal failure

**ICD-10 code**

N17-N19

**ICD-10 code description**

Renal failure

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

Uremic pruritus

**Timepoint**

Pre-intervention, after 2 weeks, post-intervention

**Method of measurement**

Visual analog scale (VAS) and measurement of pruritus scale (MPS)

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

Laboratory parameters

**Timepoint**

Pre-intervention, after 2 weekes, post intervention

**Method of measurement**

Serum

**Intervention groups****1****Description**

In the second group, subjects receive Aluminum Hydroxide syrup (30 cc; three times per day before each meal), for two weeks. Wash out period is 48 hours. In the second period treatment for tow weeks, use oral activated Charcoal capsule 6gr daily in three doses.

**Category**

Treatment - Drugs

**2****Description**

In the first group, Subjects receive oral activated Charcoal capsule 6gr daily in three doses, for two weeks. Wash out period is 48 hours. second period treatment (tow weeks), continue by Aluminum Hydroxide syrup (30 cc; three times per day before each meal).

**Category**

Treatment - Drugs

**Recruitment centers****1****Recruitment center****Name of recruitment center**

dialysis ward in 5 Azar hospital, Gorgan

**Full name of responsible person**

Mis Asyeh Khalili

**Street address**

5 Azar hospital, 5 Azar street, Gorgan, Iran

**City**

Gorgan

## Sponsors / Funding sources

### 1

#### Sponsor

**Name of organization / entity**

Golestan medical Sciences university, Vice chancellor for research

**Full name of responsible person**

Dr Arabi

**Street address**

Golestan medical sciences and health services university, Shast kola street, Gorgan

**City**

Gorgan

**Grant name****Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

**Title of funding source**

Golestan medical Sciences university, Vice chancellor for research

**Proportion provided by this source**

100

**Public or private sector**

*empty*

**Domestic or foreign origin**

*empty*

**Category of foreign source of funding**

*empty*

**Country of origin****Type of organization providing the funding**

*empty*

## Person responsible for general inquiries

**Contact****Name of organization / entity**

Golestan medical Sciences university

**Full name of responsible person**

Ali Abbasi

**Position**

Instructor & Master

**Other areas of specialty/work****Street address**

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## Person responsible for scientific

## inquiries

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## Sharing plan

**Deidentified Individual Participant Data Set (IPD)**

*empty*

**Study Protocol**

*empty*

**Statistical Analysis Plan**

*empty*

**Informed Consent Form**

*empty*

**Clinical Study Report**

*empty*

**Analytic Code**

*empty*

**Data Dictionary**

*empty*