

Clinical Trial Protocol

Iranian Registry of Clinical Trials

16 Sep 2026

The effect of progesterone supplementation on pregnancy rates in controlled ovarian stimulation and intrauterine insemination cycles: a randomized prospective trial

Protocol summary

Summary

The aim of this study was to evaluate the effect of vaginal progesterone as a luteal phase support on pregnancy rates in controlled ovarian hyper stimulation and intrauterine insemination cycles in couples with unexplained infertility or mild male factor infertility. This prospective randomized controlled trial was performed at the Shariati University Hospital, Tehran, Iran and private practice setting, Omid clinic and included 290 patients, who met the inclusion criteria. All the patients were undergone controlled ovarian hyper stimulation and intrauterine insemination. Men had at least two normal semen analyses before any treatment. 142 patients were randomized to start with a supported cycle and 148 patients started with an unsupported cycle. In supported cycles, the patients received vaginal progesterone once daily from the day after insemination until 12 weeks of pregnancy, and in non pregnant women for 14 days. No progesterone was given during unsupported cycles. The main outcome measures were clinical pregnancy rates per cycle.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201104266292N1**
Registration date: **2011-05-31, 1390/03/10**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2011-05-31, 1390/03/10

Registrant information

Name

Mahboobe Rahmani

Name of organization / entity

Tehran University of Medical Science, Shariati Hospital

Country

Iran (Islamic Republic of)

Phone

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Recruitment status

Recruitment complete

Funding source

Tehran University of Medical Sciences

Expected recruitment start date

2009-10-23, 1388/08/01

Expected recruitment end date

2010-10-23, 1389/08/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of progesterone supplementation on pregnancy rates in controlled ovarian stimulation and intrauterine insemination cycles: a randomized prospective trial

Public title

The effect of progesterone supplementation on pregnancy rates in controlled ovarian stimulation and intrauterine insemination cycles: a randomized prospective trial

Purpose

Treatment

Inclusion/Exclusion criteria

Inclusion criteria: All women included in the study were between the age 18 and 35 years and BMI of 18-28 kg/m², regular menses, and no PCOD (according to Rotterdam criteria), basal FSH < 10 IU/L and normal serum prolactin level and normal thyroid function. All couples had been trying unsuccessfully to conceive for at least one year before being enrolled in the trial. All women had bilateral tubal patency and normal uterine cavity, confirmed by hysterosalpinography (HSG), performed a maximum of 6 months before the start of the stimulation. Men had at least two semen analyses before any treatment. Normal semen analysis was defined by the threshold values of the World Health Organization criteria, ≥ 4% sperm had normal morphology according to the criteria of Kruger et al, or total motile sperm count ≥ 1 million following semen preparation. Exclusion criteria: abnormal prolactin serum level and abnormal thyroid function test, polycystic ovary syndrome according to Rotterdam criteria, diminished ovarian reserve (basal FSH level > 10 IU/mL, presence of resistant ovarian cyst (> 20 mm for > 1 month), hypogonadotropic hypogonadism or any contraindications for progesterone therapy.

Age

From **18 years** old to **35 years** old

Gender

Female

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: **300**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary IDs

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Tehran University of Medical Sciences

Street address

Keshavarz Blvd, Ghods Street

City

Tehran

Postal code

Approval date

2011-05-09, 1390/02/19

Ethics committee reference number

210/1434/90/9

Health conditions studied

1

Description of health condition studied

Female infertility, unspecified

ICD-10 code

N97.9

ICD-10 code description

inability to achieve a pregnancy

2

Description of health condition studied

Male infertility

ICD-10 code

N46

ICD-10 code description

Azoospermia NOS

Primary outcomes

1

Description

Clinical pregnancy rate

Timepoint

4 weeks after insemination

Method of measurement

Transvaginal sonography

Secondary outcomes

1

Description

Ovulation rate

Timepoint

The days 12, 14 cycles

Method of measurement

Transvaginal sonography

Intervention groups

1

Description

Intervention: application of progesterone 400 mg vaginal suppository Cyclogest once daily from the day of IUI until negative beta HCG or 12 week

Category

Treatment - Drugs

2

Description

Controi: no drug
Category
Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center
Shariati Hospital, Infertility Center
Full name of responsible person
Mahboobeh Rahmani
Street address
Gisha Brd, Amirabad Ave.
City
Tehran

Sponsors / Funding sources

1

Sponsor

Name of organization / entity
Tehran University of Medical Sciences
Full name of responsible person
Dr. Marzieh Aghahosseini
Street address
Gisha Brd, Amirabad Ave
City
Tehran
Grant name
Grant code / Reference number
Is the source of funding the same sponsor organization/entity?
Yes
Title of funding source
Tehran University of Medical Sciences
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
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Person responsible for updating data

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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