

Clinical Trial Protocol

Iranian Registry of Clinical Trials

26 May 2026

The Effect of Letrozole and Dexamethasone on pregnancy outcome in Poly cystic ovarian syndrome patients candidate for intra uterine insemination: Randomized Clinical trial

Protocol summary

Study aim

The Effect of Letrozole and Dexamethasone on ovulation and pregnancy outcome in patients with polycystic Ovarian Syndrome

Design

A prospective , randomized, single blind clinical trial with two arm parallel group design consisted of 100 infertile poly cystic ovarian syndrome patients. Randomisation was done using computer with concealed randomization sequence. Phase of trial is 1-2

Settings and conduct

This study was done on patients who referred to Milad infertility center . Group A received Letrozole from 3th day of their cycle for 5 days and group B received 5 mg Letrozole from 3th day of their cycle for 5 days and 0.5 mg Dexamethazon for 10 days. Serum hormones level (LH, Estradiol , Androstenedion, testosterone) and folliculogenesis were assessed in all patients. When the dominant follicles reached 18 mm in diameter ovulation was triggered with 5000Iu IM hCG. IUI was performed 36 hours later. Pregnancy test was done after 14 days. Statistician analyzed data without any information about the intervention.

Participants/Inclusion and exclusion criteria

Inclusion criteria: women at the age of 20-35 years old; having at least one patent Fallopian tube; normal sperm analysis of spouse. Exclusion criteria: smoking; underlying disease (heart, kidney, liver or Endocrine disease)

Intervention groups

Group A received Letrozole tablet from third day of cycle for 5 days and group B received Letrozole tablet from third day of cycle for 5 days and Dexamethazon tablet 0.5 mg for 10 days. Serum hormones level (LH, Estradiol , Androstenedion, testosterone) and folliculogenesis were monitored in patients. Ovulation was triggered with hCG (5000Iu) when the dominant follicles reached 18

mm in diameter. Intra uterine insemination (IUI) was performed 36 hours later.

Main outcome variables

hormones including luteinizing hormone, estradiol , androstenedion and testosterone; pregnancy rate .

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20170315033085N4**

Registration date: **2018-07-29, 1397/05/07**

Registration timing: **retrospective**

Last update: **2018-07-29, 1397/05/07**

Update count: **0**

Registration date

2018-07-29, 1397/05/07

Registrant information

Name

malihe afiat

Name of organization / entity

Mashhad medical school

Country

Iran (Islamic Republic of)

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+98 51 3802 2608

Email address

afiatm@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2013-02-03, 1391/11/15

Expected recruitment end date

2014-04-09, 1393/01/20
Actual recruitment start date
2013-04-09, 1392/01/20
Actual recruitment end date
2014-04-19, 1393/01/30
Trial completion date
empty

Scientific title
The Effect of Letrozole and Dexamethasone on pregnancy outcome in Poly cystic ovarian syndrome patients candidate for intra uterine insemination: Randomized Clinical trial

Public title
letrozole and Dexametazon effect in Poly cystic ovarian syndrome patients

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
AGE: 25-35 years at least one patent Fallopien tube normal sperm analysis of spouse presence of poly-cystic ovarian syndrome
Exclusion criteria:
Smoking underlying disease (heart, kidney, liver or Endocrine)

Age
From **20 years** old to **35 years** old

Gender
Female

Phase
1-2

Groups that have been masked

- Data analyser

Sample size
Target sample size: **106**
Actual sample size reached: **100**

Randomization (investigator's opinion)
Randomized

Randomization description
Simple randomization

Blinding (investigator's opinion)
Single blinded

Blinding description
Data was sent in form of two Group (A and B) to the person responsible for data analysis and analyzer wasn't not aware of the intervention.

Placebo
Not used

Assignment
Parallel

Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Mashhad University of Medical Sciences

Street address

Ghoraishi department, Daneshgah Blvd

City

Mashhad

Province

Razavi Khorasan

Postal code

91897-95-96

Approval date

2013-02-02, 1391/11/14

Ethics committee reference number

IR.MUMS.REC.1391.803

Health conditions studied

1

Description of health condition studied

polycystic ovarian syndrome

ICD-10 code

E28.2

ICD-10 code description

Polycystic ovarian syndrome

Primary outcomes

1

Description

Luteinizing hormone

Timepoint

Third and 12th day of menstrual cycle

Method of measurement

laboratory Test

2

Description

Testestron

Timepoint

Third and 12th day of menstrual cycle

Method of measurement

laboratory Test

3

Description

Anderestandion

Timepoint

Third and 12th day of menstrual cycle

Method of measurement

laboratory Test

4

Description

Esteradiol

Timepoint

Third and 12th day of menstrual cycle

Method of measurement

laboratory Test

5

Description

Pregnancy

Timepoint

14 days after Intrauterine insemination

Method of measurement

laboratory Test, pregnancy test

Secondary outcomes

1

Description

Dominant follicles

Timepoint

12th day of Menestural cycle

Method of measurement

vaginal Sonography

2

Description

Endometrium thickness

Timepoint

12th day of Menestural cycle

Method of measurement

vaginal Sonography

Intervention groups

1

Description

Control group: group A received letrozole tablet (abouraihan , Iran) 5 mg from day 3 of menstrual cycle for 5 days .They were monitored hormonal and Sonographical on day 3 and day 12 of cycle. Ovulation was triggered with injection 5000 lu HCG (karma, Germany) when the dominant follicle reached 18 mm in diameter. A single IUI was performed 36 hours later. The Luteal phase was supported with the progesterone suppository (Cyclogest, 400 mg , Actavis, UK) . pregnancy test was don 14 day later.

Category

Treatment - Drugs

2

Description

Intervention group: group B received letrozol 5 mg(Abouraihan-Iran) from 3th day of menstrual cycle for 5 days plus Dexamethazon tablet 0.5 mg (Iran hormone) for 10 days. They were monitored hormonal and Sonographical on day 3 and day 12 of cycle. Ovulation was triggered with injection 5000 lu HCG (karma, Germany) when the dominant follicle reached 18 mm in

diameter. A single IUI was performed 36 hours later. The Luteal phase was supported with the progesterone suppository (Cyclogest, 400 mg , Actavis, UK) . pregnancy test was don 14 day later.

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Milad Hospital

Full name of responsible person

Dr Malihe Afiat

Street address

Emamreza hospital, Ebne Sina Blvd,Shariati Square

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Email

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Mashhad University of Medical Sciences

Full name of responsible person

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Obstetrics department, Imam Reza hospital, Shariati Square

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Grant name

Grant code / Reference number

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

Yes

Title of funding source

Mashhad University of Medical Sciences

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

Phone

+92 51 38022608

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+98 21 6658 1616

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Web page address**Person responsible for general inquiries****Contact****Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

Malihe Afiat

Position

Assistance Professor

Latest degree

Specialist

Other areas of specialty/work

Gynecology and Obstetrics

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Web page address**Person responsible for updating data****Contact****Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

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Position

Assistance Professor

Latest degree

Specialist

Other areas of specialty/work

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Full name of responsible person

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Position

Assistance Professor

Latest degree

Specialist

Other areas of specialty/work

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Imam Reza hospital, Shariat square

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Postal code

91897-57565

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

No - There is not a plan to make this available

Justification/reason for indecision/not sharing IPD

No more information

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