

Clinical Trial Protocol

Iranian Registry of Clinical Trials

02 Jun 2026

Effect of GnRH-a in endometrial preparation for frozen-thawed embryo transfer cycles in patient with repeated implantation failure.

Protocol summary

Study aim

This study was conducted to investigate the effects of GnRH agonist administration on the implantation rate in the FET cycles of women with RIF, prior to estrogen-progesterone preparation of the endometrium.

Design

The case group (n = 34) received 0.1 mg/day of the GnRH agonist (Variopeptyl, VarianDarou, Iran), subcutaneous, from day 21 of the cycle preceding the actual FET cycle. On the second day of the cycle, the dose of GnRH agonist was reduced to 0.05 mg and 6 mg/day oral estradiol valerate (2 mg, Aburaihan Co., Tehran, Iran) was also started. When the endometrial thickness reached to 7.5 mm, vaginal supplementation of Cyclogest® pessaries (Cox Pharmaceuticals, Barnstaple, UK) at 400 mg twice daily was started and the GnRH agonist was also stopped. The control group (n = 33), received 6 mg/day oral estradiol valerate (2 mg, Aburaihan Co., Tehran, Iran) from the second day of the cycle without the GnRH agonist. In the two groups, frozen-thawed embryos were transferred on the fourth day of progesterone treatment.

Settings and conduct

In this randomized clinical trial, 67 infertile women with history of idiopathic RIF (at least two implantation failures), undergoing FET cycles in Yazd Reproductive Sciences Institute.

Participants/Inclusion and exclusion criteria

67 infertile women with history of idiopathic RIF (at least two implantation failures), undergoing FET cycles included. All women with endometrial polyp, uterine myoma, and uterine anomaly were excluded from the study

Intervention groups

67 infertile women with history of idiopathic RIF (at least two implantation failures), undergoing FET cycles included.

Main outcome variables

The clinical outcomes) including chemical and clinical

pregnancy, in addition to implantation rates, were compared between the two groups.

General information

Reason for update

Update clinical trial based on the latest changes in method and recruitment and trial completion dates

Acronym

IRCT registration information

IRCT registration number: **IRCT201708292604N3**

Registration date: **2017-09-18, 1396/06/27**

Registration timing: **registered_while_recruiting**

Last update: **2021-04-19, 1400/01/30**

Update count: **1**

Registration date

2017-09-18, 1396/06/27

Registrant information

Name

Robab Davar

Name of organization / entity

Yazd Research and Clinical Center for Infertility

Country

Iran (Islamic Republic of)

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+98 35 1824 7085

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Recruitment status

Recruitment complete

Funding source

Vice Chancellor for research of Shahid Sadoughi University of Medical Sciences, Yazd, Iran

Expected recruitment start date

2017-08-16, 1396/05/25

Expected recruitment end date

2017-11-16, 1396/08/25
Actual recruitment start date
2017-08-01, 1396/05/10
Actual recruitment end date
2017-11-30, 1396/09/09
Trial completion date
2018-01-30, 1396/11/10

Scientific title
Effect of GnRH-a in endometrial preparation for frozen-thawed embryo transfer cycles in patient with repeated implantation failure.

Public title
Effect of GnRH-a in endometrial preparation for frozen-thawed embryo transfer cycles in patient with repeated implantation failure.

Purpose
Treatment

Inclusion/Exclusion criteria
Inclusion criteria:
infertile women with at least 2 previous failed implantation.
Exclusion criteria:
Endometrial polyp myoma uterine anomaly

Age
From **18 years** old to **42 years** old

Gender
Female

Phase
2-3

Groups that have been masked
No information

Sample size
Target sample size: **67**
Actual sample size reached: **67**

Randomization (investigator's opinion)
Randomized

Randomization description
Patients are divided into two groups of 34 patients according to permutation block method of control and intervention. Therapeutic tasks within the blocks are determined in such a way that they are random, but the desired allocation ratio is achieved in each block. There are 8 blocks in each, is 8 numbers in order to reaches sample size to 64 and one block of 4 is considered. Generate random codes using random block allocation method which will be generated with the help of Random allocation software version 1. The first person eligible to enter the study is given number one and so on until the last person eligible for number 67 is given. Using a table generated by random allocation software by number, individuals are placed in the control and intervention group. The output of Random allocation software version 1 is specified as A and B in the specified blocks, which describe the number of blocks and the number of patients in each block.

Blinding (investigator's opinion)
Not blinded

Blinding description

Placebo
Not used

Assignment
Parallel
Other design features

Secondary Ids
empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics Committee of Yazd Reproductive Sciences
institute shahid sadoughi university

Street address

Yazd Reproductive Sciences Institute

City

Yazd

Province

Yazd

Postal code

8916877391

Approval date

2017-05-21, 1396/02/31

Ethics committee reference number

IR.SSU.RSI.REC.1396.1

Health conditions studied

1

Description of health condition studied

repeated implantation failure

ICD-10 code

N98.9

ICD-10 code description

complication associated with artificial fertilization
,unspecified

Primary outcomes

1

Description

chemical pregnancy

Timepoint

2 weeks after embryo transfer

Method of measurement

serum BHCG

2

Description

clinical pregnancy

Timepoint

2 wk after positive β hCG

Method of measurement

the detection of a fetal heartbeat in sonography

3

Description

Implantation rate

Timepoint

2 wk after positive β hCG

Method of measurement

the number of gestational sacs per embryos transferred

Secondary outcomes

empty

Intervention groups

1

Description

In case group , patients receive daily subcutaneous injections of 0.1 mg of GnRH-a(diphereline) starting from day 21 of previous cycles.The dose is reduce to 0.05 mg from the second day of cycle when daily oral estradiol valerate 6mg is also started.Daily vaginal progesterone 400mg twice daily is started when endometrial thickness is > 7.5 mm and GnRH_a is also stopped.

Category

Treatment - Drugs

2

Description

In control group oral estradiol 6mg daily is started from day 2 cycle without diphereline. Daily vaginal progesterone 400mg twice daily is started when endometrial thickness is > 7.5 mm

Category

Treatment - Drugs

Recruitment centers

1

Recruitment center

Name of recruitment center

Yazd Reproductive Sciences Institute

Full name of responsible person

Dr.Robab Davar

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Yazd Safaeye Bouali Ave.

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Yazd Reproductive Sciences Institute, Shahid Sadoughi University

Full name of responsible person

Dr.Robab Davar

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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

Yazd Reproductive Sciences Institute, Shahid Sadoughi University

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

Yazd Reproductive Sciences Institute

Full name of responsible person

Dr.Robab Davar

Position

Professor

Latest degree

Subspecialist

Other areas of specialty/work

Gynecology and Obstetrics

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Person responsible for scientific inquiries

Contact

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Full name of responsible person
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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Yes - There is a plan to make this available

Statistical Analysis Plan

Yes - There is a plan to make this available

Informed Consent Form

Yes - There is a plan to make this available

Clinical Study Report

Yes - There is a plan to make this available

Analytic Code

Yes - There is a plan to make this available

Data Dictionary

Yes - There is a plan to make this available

Title and more details about the data/document

participant data set able to be shared

When the data will become available and for how long

2 months after publication

To whom data/document is available

Journal that published the results

Under which criteria data/document could be used

A person/office that is legally in charge

From where data/document is obtainable

Yazd Reproductive Sciences Institute 983538247085
saeideh_dashti@yahoo.com

What processes are involved for a request to access data/document

Submission of legal order

Comments