

Clinical Trial Protocol

Iranian Registry of Clinical Trials

09 Jun 2026

Efficacy of intrauterine infusion of platelet_rich plasma on pregnancy outcome in patients with repeated implantation failure.

Protocol summary

Study aim

To evaluate the effectiveness of platelet-rich plasma in improvement of pregnancy and implantation rate in women with history of repeated implantation failure.

Design

Clinical trial with total of 120 eligible women that randomly assigned into two equal parallel groups (intervention and control) through Balanced Block Randomization. (Block size=4).

Settings and conduct

After endometrial preparation for embryo transfer in both groups, In intervention group, 48 hours before embryo transfer (ET), 20 ml of peripheral venous blood will be obtained by syringe pre-filled with 2ml of anticoagulant solution, and then will be centrifuged immediately at 170 g for 10 minutes to separate red blood cells. This plasma will be re-centrifuged at 340 g for 5 minutes to obtain platelet-rich plasma (PRP), that contained platelet 4-5 times more than peripheral blood. 0.5ml of PRP will be infused into the uterine cavity with IUI catheter and after 48 hours, ET will be done. In control group, ET will be done without intrauterine infusion of PRP. After 2 weeks of ET, serum Beta_human chorionic Gonadotropin (β HCG) will be measured in all patients. This research will be done at Infertility Department Of Shariati Hospital, Tehran, Iran

Participants/Inclusion and exclusion criteria

Infertile 20-40 years old women that failed to conceive after three or more embryo transfer and have at least one frozen good - quality blastocyst- stage embryo, body mass index below 30 kg/m², without Genetic, Chromosomal, Uterine, Immunologic, Hormonal and Hematologic disorders.

Intervention groups

In intervention group, 48 hours before embryo transfer, 0.5ml of platelet-rich plasma will be infused into the uterine cavity with IUI catheter. In control group embryo transfer undergoing without intrauterine infusion of platelet-rich plasma.

Main outcome variables

Clinical Pregnancy Implantation Rate

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20190503043454N1**

Registration date: **2019-07-06, 1398/04/15**

Registration timing: **registered_while_recruiting**

Last update: **2019-07-06, 1398/04/15**

Update count: **0**

Registration date

2019-07-06, 1398/04/15

Registrant information

Name

Parvaneh Lak

Name of organization / entity

Country

Iran (Islamic Republic of)

Phone

+98 21 2249 8504

Email address

lakparvaneh@yahoo.com

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-05-22, 1398/03/01

Expected recruitment end date

2019-08-23, 1398/06/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Efficacy of intrauterine infusion of platelet_rich plasma on pregnancy outcome in patients with repeated implantation failure.

Public title

Efficacy of platelet_rich plasma on embryo implantation

Purpose

Treatment

Inclusion/Exclusion criteria**Inclusion criteria:**

Infertile women between 20 -40 years old
History of at least 3 or more failure in embryo transfer
Have at least one frozen Blastocyst - stage embryo
Body Mass Index below 30 kg/m2

Exclusion criteria:

Congenital or acquire uterine abnormalities
Genetic or chromosomal abnormalities
Immunologic disorders
Hematologic disorders
Hormonal disorders

Age

From **20 years** old to **40 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **120**

Randomization (investigator's opinion)

Randomized

Randomization description

Balanced Block Randomization (Block size=4): We prepare four balls,two A and two B.for first patient ,a nurse who does not know our plan for both A and B groups, selects one ball randomly from basket.for next patient this selection were done through remaining three balls.for third patient from remaining two balls and the last ball for the fourth patient, and then another block was selected.therefore our blocks show one of these pattern. (BAAB) (BABA) (ABBA) (BBAA) (ABAB) (AABB)

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 Tehran University of Medical Sciences

Street address

Room 605, Floor 6 , Central Building Of Tehran University Of Medical Science., Keshavarz Blvd

City

Tehran

Province

Tehran

Postal code

1417653761

Approval date

2017-08-09, 1396/05/18

Ethics committee reference number

86388

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

Repeated Implantation Failure

ICD-10 code**ICD-10 code description****Primary outcomes****1****Description**

Clinical Pregnancy

Timepoint

4 weeks after embryo transfer

Method of measurement

To detect gestational sac with Fetal Heartbeat by Trans Vaginal Ultrasound.

2**Description**

The Implantation Rate

Timepoint

Four weeks after the embryo transfer

Method of measurement

The ratio of gestational sac determination on trans vaginal ultra Sonography to the number of transferred embryos.

Secondary outcomes

empty

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

Intervention group:48 hours before embryo transfer (ET), 20 ml of peripheral venous blood will be obtained by syringe pre-filled with 2ml of Acid Citrate A anticoagulant solution(Royagen),and then will be centrifuged

immediately at 170 g for 10 minutes to separate red blood cells. this plasma will be re-centrifuged at 340 g for 5 minutes to obtain platelet-rich plasma (PRP), that contained platelet 4-5 times more than peripheral blood. 0.5ml of PRP will be infused into the uterine cavity with IUI catheter and after 48 hours, ET will be done with IUI catheter (Takwin, Iran).

Category

Treatment - Other

2**Description**

Control group: ET will be done with IUI catheter without intrauterine infusion of PRP.

Category

N/A

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

Shariati hospital

Full name of responsible person

Lak Parvaneh

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Shariati Hospital, Jalal-e-Al-e-Ahmad Hwy

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Sponsors / Funding sources**1****Sponsor****Name of organization / entity**

Tehran University of Medical Sciences

Full name of responsible person

Mohammad Ali Sahraian

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TUMS Vice Chancellor for Research., Tehran
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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

Tehran 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

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

Tehran University of Medical Sciences

Full name of responsible person

Parvaneh Lak

Position

Fellowship of infertility

Latest degree

Specialist

Other areas of specialty/work

Gynecology and Obstetrics

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Person responsible for scientific inquiries**Contact****Name of organization / entity**

Tehran University of Medical Sciences

Full name of responsible person

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Person responsible for updating data

Contact

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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