

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

28 Jun 2026

### Randomised clinical trial of comparison between with and without atraumatic hysteroscopy on the improvement of endometrial receptivity on the shariati hospitals patient in 1394

#### Protocol summary

##### Summary

The aim of study :Compare Clinical pregnancy rates for controlled ovarian stimulation in IVF cycles with and without hysteroscopy 250 patients with primary infertility with regular menses under the age of 40 years who meet all the inclusion criteria After providing adequate explanations by the investigators and after patients acceptance will be entered to the study. The patients will be divided between two group randomly. Trans vaginal sonography (TVS) will be done for all patients in 1-3 cycle days . Ovarian stimulation will be done with antagonist regimen in two groups with recombinant 225 IU FSH daily, the dose will adjusted according to the ovarian response in TVS. The GNRH antagonist will start when the follicles become 14 mm in diameter. The final oocyte maturation will be achieved by the administration of HCG 10000 IU when 3 follicles become 17mm in diameter. Oocyte retrieval will be carried out 36 hours after HCG administrations. Embryo transfer will be performed on day 3 after oocyte retrieval and two grade 1 embryos will be transferred. Luteal phase supplementation consist of 3 suppository progesterone will be started the day after oocyte retrieval and will be continued to 7 weeks of gestation. For patients in the intervention group uterine cavity irrigation with normal saline in an outpatient setting with or without mild sedation will be done by doing hysteroscopy in the days 5-7 of stimulation cycle . After 2 and 4 weeks of embryo transfer chemical and clinical pregnancy will be evaluated by BHCG test and TVS respectively.

#### General information

##### Acronym

cycle hysteroscopy

##### IRCT registration information

IRCT registration number: **IRCT2016011022795N2**

Registration date: **2016-03-04, 1394/12/14**

Registration timing: **registered\_while\_recruiting**

Last update:

Update count: **0**

##### Registration date

2016-03-04, 1394/12/14

##### Registrant information

###### Name

Marzieh Ghasemi

###### Name of organization / entity

Zahedan University of Medical Sciences

###### Country

Iran (Islamic Republic of)

###### Phone

+98 54 3329 5581

###### Email address

drghasemi@zaums.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

Tehran University of Medical Sciences

##### Expected recruitment start date

2015-04-21, 1394/02/01

##### Expected recruitment end date

2016-03-05, 1394/12/15

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

##### Trial completion date

empty

##### Scientific title

Randomised clinical trial of comparison between with and without atraumatic hysteroscopy on the

improvement of endometrial receptivity on the shariati hospitals patient in 1394

## Public title

evaluation of hysteroscopic effect on endometrial receptivity of the patients in the IVF cycle

## Purpose

Treatment

## Inclusion/Exclusion criteria

Inclusion criteria:regular mense;age less than 40;BMI=19-30;without any prior hysteroscopy examination or previous IVF or ICSI; with a normal Transvaginal sonography (TVS) (in the day 2-3 same cycle);normal hystrosalpingo graphy (HSG) (between 6to24 mounths) . Exclusion criteria : recurrent miscamiage (3 or more miscarriage); inter menstrual bleeding; any doubt about uterine cavity abnormality; server male factor request to PESA or TESE; patients with poly cystic ovary syndrome or endometriosis ;hydrosalpinx ;ovarian cyst ; cancellation of the same cycle for any reason

## Age

From **18 years** old to **40 years** old

## Gender

Female

## Phase

N/A

## Groups that have been masked

*No information*

## Sample size

Target sample size: **250**

## Randomization (investigator's opinion)

Randomized

## Randomization description

## 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

Tehran University of Medical Sciences

##### Street address

central building of Tehran University of Medical Sciences ,keshavarz BLV,Tehran,Iran

##### City

Tehran

##### Postal code

1417614411

#### Approval date

2010-09-23, 1389/07/01

#### Ethics committee reference number

IR.TUMS.REC.1394.1598

## Health conditions studied

### 1

#### Description of health condition studied

Infertility

#### ICD-10 code

N97.0,N97.

#### ICD-10 code description

Female infertility associated with anovulation, Female infertility of tubal origin, Female infertility of other origin, Female infertility, unspecified,

## Primary outcomes

### 1

#### Description

clinical pregnancy rate in two groups

#### Timepoint

4 weeks after embryo transfer transvaginal sonography will be done and gestational sac will be evaluated

#### Method of measurement

Find gestational sac on transvaginal sonography

## Secondary outcomes

### 1

#### Description

chemical pregnancy rate in two group ,abortion , implantation rate in two groups

#### Timepoint

two weeks after transfer for biochemical pregnancy,4weeks for clinical pregnancy,12 weeks after transfer for abortion,two weeks after transfer for implantation rate

#### Method of measurement

BHCG for biochemical ,trans vaginal sonography for clinical,physical examination and trans vaginal sonography for abortion ,trans vaginal sonography for implantation

## Intervention groups

### 1

#### Description

Performing hysteroscopy in the 5-7 days of the IVF cycle treatment and uterine irrigation with normal saline in the patients in intervention group . For patients in the control group did not add any intervention.

#### Category

Treatment - Other

## 2

### Description

In the control group the traditional IVF cycle will be done without any additional intervention

### Category

Treatment - Drugs

## Recruitment centers

### 1

#### Recruitment center

##### Name of recruitment center

Shriati hospital,Infertility ward

##### Full name of responsible person

Dr Marzieh Ghasemi,fellowship of infertility

##### Street address

Tehran,Jalal-Al -Ahmad BLV crossed north kargar,Shariati hospital

##### City

Tehran

## Sponsors / Funding sources

### 1

#### Sponsor

##### Name of organization / entity

Tehran University of Medical Sciences

##### Full name of responsible person

Dr Ashraf Alyacin

##### Street address

Tehran, Keshavarz Blvd, Tehran University of Medical Sciences

##### 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

Tehran University of Medical Sciences,infertility ward

#### Full name of responsible person

Marzieh Ghasemi

#### Position

fellowship in infertility

#### Other areas of specialty/work

#### Street address

Shariati Hospital , Jalal-Al-Ahmad BLV intersection Kargar shomali,Tehran, Iran

#### City

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ma@razi.tums.ac.irghasemim841@yahoo.com

#### Web page address

## Person responsible for scientific inquiries

### Contact

#### Name of organization / entity

Tehran University of Medical Sciences,Obestetrics and Gynecology group ,infertility ward

#### Full name of responsible person

Dr Ashraf Alyacin

#### Position

professor of Obestetrics and Gynecology

#### Other areas of specialty/work

#### Street address

,Infertility ward ,Shariati Hospital ,Tehran University of Medical Sciences

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## Person responsible for updating data

### Contact

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#### Full name of responsible person

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#### Position

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#### Other areas of specialty/work

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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*