

Clinical Trial Protocol

Iranian Registry of Clinical Trials

11 Jun 2026

The effect of aspirin administration for prevention of preeclampsia in pregnant women with abnormal findings in uterine artery doppler sonography

Protocol summary

Summary

In this double-blind randomized clinical trial, singleton pregnant women with gestational age between 11 to 15 weeks in prenatal clinic, Qazvin Kosar hospital are evaluated by uterine artery doppler sonography. Women with underlying disease, history of asthma, sensitivity to aspirin, termination of pregnancy before 20 weeks of gestation are excluded. One hundred pregnant women with abnormal findings in doppler sonography are randomly assigned to the intervention and control groups. Intervention group received aspirin 80 mg daily between 14 - 16 weeks of gestation until 37 weeks. The control group received a placebo pill daily. The two groups are followed until the end of pregnancy and pregnancy outcome in terms of preeclampsia, intrauterine growth restriction, preterm labor, mode of delivery, birth weight, first and fifth minutes apgar score are investigated.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT201505103025N5**
Registration date: **2015-06-08, 1394/03/18**
Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2015-06-08, 1394/03/18

Registrant information

Name

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Name of organization / entity

Qazvin University of Medical Sciences and Health

Services

Country

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Recruitment status

Recruitment complete

Funding source

Qazvin University of Medical Sciences

Expected recruitment start date

2014-09-23, 1393/07/01

Expected recruitment end date

2015-06-22, 1394/04/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of aspirin administration for prevention of preeclampsia in pregnant women with abnormal findings in uterine artery doppler sonography

Public title

Effect of aspirin for prevention of preeclampsia in women with abnormal ultrasonic findings in uterine artery

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: pregnant women; singletone; gestational age 11-15 weeks; abnormal finding in uterine artery doppler sonography exclusion criteria: underlying medical condition; history of asthma; allergic to aspirin; termination of pregnancy before 20 weeks of pregnancy

Age

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

Gender

Female

Phase

3

Groups that have been masked

No information

Sample size

Target sample size: **100**

Randomization (investigator's opinion)

Randomized

Randomization description**Blinding (investigator's opinion)**

Double blinded

Blinding description**Placebo**

Used

Assignment

Parallel

Other design features**Secondary Ids**

empty

Ethics committees**1****Ethics committee****Name of ethics committee**

Ethics Committee, Qazvin University of Medical Sciences

Street address

Iran, Qazvin, Bahonar Blvd

City

Qazvin

Postal code

4321102589

Approval date

2014-06-12, 1393/03/22

Ethics committee reference number

28/20/8853

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

Preeclampsia

ICD-10 code

O13

ICD-10 code description

Gestational [pregnancy-induced] hypertension without significant proteinuria

2**Description of health condition studied**

Preeclampsia

ICD-10 code

O14

ICD-10 code description

Gestational [pregnancy-induced] hypertension with significant proteinuria

Primary outcomes**1****Description**

Preeclampsia

Timepoint

Measurement of Blood pressure every two weeks until 28 w then every week until termination of pregnancy

Method of measurement

Barometer, mmHg

Secondary outcomes**1****Description**

Intrauterine Growth Retardation

Timepoint

Every two weeks until 28 w then every week until termination of pregnancy

Method of measurement

Doppler Ultrasound If fetal growth retardation is suspected

2**Description**

1 and 5 min apgar

Timepoint

1 and 5 min after birth

Method of measurement

Standard apgar score

3**Description**

Neonatal Weight

Timepoint

After Birth

Method of measurement

Digital Scale(gram)

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

Intervention Group: aspirine tablet 80 mili gram daily after 14-16 week of gestation until 37 week of pregnancy

Category

Prevention

2**Description**

Control Group: placebo tablet daily after 14-16 week of

gestation until 37 week of pregnancy

Category

Placebo

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

Kosar Hospital

Full name of responsible person

Farideh Movahed

Street address

Iran,Qazvin, Taleghani St, Kosar Hospital

City

Qazvin

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

Vice-Chancellor for Research of Qazvin University of Medical Sciences

Full name of responsible person

Farideh Movahed

Street address

Iran, Qazvin, Bahonar Blvd

City

Qazvin

Grant name**Grant code / Reference number****Is the source of funding the same sponsor organization/entity?**

Yes

Title of funding source

Vice-Chancellor for Research of Qazvin 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**

Qazvin University of Medical Sciences

Full name of responsible person

Farideh Movahed

Position

Associate Professor, Department of Obstetrics and Gynecology

Other areas of specialty/work**Street address**

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Full name of responsible person

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Web page address**Person responsible for updating data****Contact****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