

Clinical Trial Protocol

Iranian Registry of Clinical Trials

06 Jul 2026

Comparison between the effect of magnesium sulfate alone and magnesium sulfate with celecoxib to prevent spontaneous preterm labor in pregnant women

Protocol summary

Summary

In this randomized double-blind clinical trial, 164 singleton pregnant women with intact amniotic membranes, between 24-34 weeks gestational age, are enrolled with a diagnosis of preterm labor. They are assigned in two groups by colored cards. Magnesium sulfate is prescribed the first group at a dose of 4 grams and followed by two grams per hour intravenously for 24 hours, at the same time a 100-miligrams celecoxib capsule is administered orally. In the second group magnesium sulfate with the same dose is given and placebo is prescribed instead of celecoxib. Patients and the doctor who controls them are not aware of the kind of the consumed drug. Uterine contractions in both groups are recorded by tocometer every one hour. The first time that the uterine contraction decreases is considered as onset of drug action. If the uterine contractions do not remove after 8 hours, administering of celecoxib or placebo continues every 8 hours for 24 hours. And if this way also do not prevent the uterine contraction after 24 hours, the case is recorded as treatment failure and treatment is stopped.

General information

Acronym

-

IRCT registration information

IRCT registration number: **IRCT2015042621947N1**
Registration date: **2016-05-18, 1395/02/29**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2016-05-18, 1395/02/29

Registrant information

Name

Shahin Nemati

Name of organization / entity

Kosar Hospital

Country

Iran (Islamic Republic of)

Phone

+98 28 3223 6374

Email address

sh.nemati@qums.ac.ir

Recruitment status

Recruitment complete

Funding source

Qazvin University of Medical Science

Expected recruitment start date

2013-03-20, 1391/12/30

Expected recruitment end date

2015-03-20, 1393/12/29

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

Comparison between the effect of magnesium sulfate alone and magnesium sulfate with celecoxib to prevent spontaneous preterm labor in pregnant women

Public title

Effect of Celecoxib in order to prevent spontaneous preterm labor

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: Singleton pregnancy; Intact amniotic

membrane; Gestational age between 24-34 weeks; Positive tocometry. Exclusion criteria: PROM; Vaginal bleeding; Chorioamnionitis; Dilatation more than 2 cm and cervical effacement exceeding 80%; Polyhydramnios; Oligohydramnios; Intrauterine Fetal Demise; Intrauterine Growth Restriction; Fetal distress; Smoking and alcohol abuse; systemic disease; Congenital anomalies; uterine anomalies; Celecoxib intolerance.

Age

From **18 years** old to **46 years** old

Gender

Female

Phase

2

Groups that have been masked

No information

Sample size

Target sample size: **164**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Double blinded

Blinding description

Placebo

Used

Assignment

Parallel

Other design features

82 yellow cards and 82 green cards have put in the box, for each patient with study including criteria, have exit one card randomly: green card for first group (magnesium sulfate alone) and yellow card for second group (magnesium sulfate with celecoxib).

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Ethics committee of Qazvin University of Medical Sciences

Street address

Bahonar Blvd, Qazvin, Iran

City

Qazvin

Postal code

4321102589

Approval date

2010-08-20, 1389/05/29

Ethics committee reference number

28/20/8859

Health conditions studied

1

Description of health condition studied

Preterm Labor

ICD-10 code

O60.1

ICD-10 code description

Preterm spontaneous labour with preterm delivery

Primary outcomes

1

Description

Preterm Labor

Timepoint

One hour after the onset of drug use until 24 hours

Method of measurement

Tocometer

Secondary outcomes

empty

Intervention groups

1

Description

Intervention group: initial dose 4 g of magnesium sulfate intravenously and then continuation of magnesium sulfate administration at a dose of 2g per hour for 24 hours with a 100 mg celecoxib capsule and if needed its continuation every 8 hours for 24 hours

Category

Prevention

2

Description

Control group: initial dose 4g of magnesium sulfate intravenously and then continuation of magnesium sulfate administration at a dose of 2g per hour for 24 hours with a placebo capsule and if needed its continuation every 8 hours for 24 hours

Category

Placebo

Recruitment centers

1

Recruitment center

Name of recruitment center

Kosar Hospital

Full name of responsible person

Shahin Nemati

Street address

Kosar Hospital, Taleghani St, Qazvin, Iran

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

Shahin Nemati

Street address

Bahonar Blvd, Qazvin, Iran

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

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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