

Clinical Trial Protocol

Iranian Registry of Clinical Trials

28 Jun 2026

The effect of topical use of breast milk on episiotomy healing in Primiparous women at high risk for wound infection

Protocol summary

Study aim

Determining the effect of topical use of breast milk on episiotomy healing in Primiparous women at high risk for wound infection

Design

Clinical trial, superiority, with intervention and control groups, without blinding, simple random assignment, on 72 patients, www.randomization.com was used for randomization.

Settings and conduct

The present study will be performed in the maternity ward of Umm Al-Banin Hospital in Mashhad. In the intervention group, in addition to routine rinsing of the perineal episiotomy with 9% non-injectable normal saline, 2 ml is infused on the perineal wound site and 0.25 ml in the first three days. Gently squeeze with gas for 3 minutes and the control group will perform routine care. The episiotomy site will be examined in the first 24 hours, on the fifth and tenth days, and its improvement will be scored according to the REEDA scale. research units complete the follow-up checklist during the study period and deliver it to the researcher on days five and ten

Participants/Inclusion and exclusion criteria

The gestational age is between 38 to 42 weeks :Vaginal delivery! Be primiparous! Episiotomy incision should be of the mediolateral type! Have at least two conditions predisposing to infection at the same time! The episiotomy is first and second degree

Intervention groups

In the intervention group, topical application of episiotomy with breast milk, in addition to routine rinsing of the perineum with normal saline, is performed for ten days, and in the control group, only routine rinsing of the perineum with normal saline is performed during the same period

Main outcome variables

The interval between rupture of the amniotic sac and the time of delivery; BMI; Number of vaginal examinations;

The distance between the two cutting edges before skin repair; Number of skin sutures; Observance of hygienic principles; Use of antibiotics; Suture opening; Wound infection

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20210121050098N1**

Registration date: **2021-03-24, 1400/01/04**

Registration timing: **registered_while_recruiting**

Last update: **2021-03-24, 1400/01/04**

Update count: **0**

Registration date

2021-03-24, 1400/01/04

Registrant information

Name

Kobra Sadat Hosseini

Name of organization / entity

Country

Iran (Islamic Republic of)

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+98 51 3859 1511

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Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2021-02-19, 1399/12/01

Expected recruitment end date

2021-05-20, 1400/02/30

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of topical use of breast milk on episiotomy healing in Primiparous women at high risk for wound infection

Public title

The effect of topical use of breast milk on episiotomy healing in Primiparous women at high risk for wound infection

Purpose

Prevention

Inclusion/Exclusion criteria**Inclusion criteria:**

Iranian nationality and resident of Mashhad Have at least literacy Have informed consent to participate in the research The gestational age is between 38 - 42 weeks The fetus is single and alive Show the fetus at the top of the head Vaginal and spontaneous delivery has been performed Be primiparous Episiotomy incision should be of the mediolateral type Have at least two conditions predisposing to infection at the same time The episiotomy is first and second degree

Exclusion criteria:

Do follow a special diet . such as a diet of vegetables or just meat Do use tobacco or drugs The fetus does suffer from severe and persistent respiratory distress Do use certain medications (including anticoagulants, antidepressants, antiepileptics, alcohol and benzodiazepines) Use of assistive devices in natural childbirth The death of a baby or infant with major abnormalities Have a history of reconstructive surgery and vaginal and perineum lesions

Age

From **15 years** old to **49 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **72**

Randomization (investigator's opinion)

Randomized

Randomization description

Sampling is available by simple random assignment. After determining the sample size, through a table of random numbers using the site www.randomization.com, the numbers are divided into two groups of sequences marked with the letters A and B. The sequence stored in sealed envelopes that the researcher uses when selecting a research unit to send individuals to two groups that meet the inclusion criteria.

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 Mashhad University of Medical Sciences, School of Nursing and Midwifery

Street address

School of Nursing and Midwifery, Doktora Crossroads, Daneshgah St., Mashhad

City

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Province

Razavi Khorasan

Postal code

9137913199

Approval date

2020-12-29, 1399/10/09

Ethics committee reference number

IR.MUMS.NURSE.REC.1399.071

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

episiotomy wound healing

ICD-10 code

O86.0

ICD-10 code description

Infection of obstetric surgical wound

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

Redness

Timepoint

The first 24 hours, the fifth and tenth days after delivery

Method of measurement

REEDA Scale

2**Description**

Edema

Timepoint

The first 24 hours, the fifth and tenth days after delivery

Method of measurement

REEDA Scale

3

Description

Ecchymosis

Timepoint

The first 24 hours, the fifth and tenth days after delivery

Method of measurement

REEDA Scale

4

Description

Discharge

Timepoint

The first 24 hours, the fifth and tenth days after delivery

Method of measurement

REEDA Scale

5

Description

Approximation

Timepoint

The first 24 hours, the fifth and tenth days after delivery

Method of measurement

REEDA Scale

Secondary outcomes

1

Description

The rate of episiotomy healing in women at high risk for wound infection

Timepoint

The first 24 hours, days five and ten after delivery

Method of measurement

REEDA Scale

Intervention groups

1

Description

Intervention group: In addition to routine rinsing of the perineal episiotomy with normal saline 9% non-injectable (twice a day every 12 hours - routine washing of Mashhad training hospitals with normal saline for ten days), the perineal wound will be soaked in breast milk. This is done for ten days after delivery twice a day (every 12 hours) each time with a new 5 ml syringe at the rate of 2 ml of breast milk and with a 2 ml syringe without a needle from the first day on the perineal wound. It is instilled (in the first three days, this amount is 0.25 ml, which is instilled into the perineal wound) and then gently squeezed with gas for 3 minutes.

Category

Treatment - Other

2

Description

Control group: Recommended to wash the perineum with

9% non-injectable normal saline twice a day for ten days (according to the routine of Umm Al-Banin Hospital in Mashhad)

Category

Treatment - Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Ommolbanin hospital

Full name of responsible person

Faride Akhlaghi

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Mashhad University of Medical Sciences

Full name of responsible person

Dr. Mohsen Tafaghodi

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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
Mashhad 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

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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

Documents are shared after being unidentified

When the data will become available and for how long

8 months

To whom data/document is available

All researchers

Under which criteria data/document could be used

All women with natural childbirth who have had an episiotomy and Midwifery service personnel

From where data/document is obtainable

To email address hoseinik973@mums.ac.ir

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

First, send an e-mail and if you do not respond within a week, you can refer to the library of Mashhad School of Nursing and Midwifery at Ibn Sina St.

Comments