Clinical Trial Protocol
Iranian Registry of Clinical Trials

07 Nov 2026

The effect of Hypericum Perforatum (Perforan) on wound healing episiotomy in women

Protocol summary

Summary
The objective of this study was the effect of Hypericum Perforatum (Perforan) on wound healing of episiotomy. 120 primiparous mothers with singleton pregnancy for whom episiotomy had been done after uncomplicated normal vaginal delivery were randomly allocated into either intervention or control group. Subjects with any acute or chronic systemic diseases, hypersensitivity, severe wound infection at fifth day after delivery or allergy to Hypericum Perforatum derivatives were excluded. Intervention under study was using Hypericum Perforatum ointment (three times a day for to 10 days) and Patients in control group received placebo ointment. Main outcome measures were pain on wound site, wound site edema, wound infection, suppurative secretion, number of remained sutures at fifth and tenth day after delivery, and wound dehiscence.

General information

Acronym
-

IRCT registration information
IRCT registration number: **IRCT201505061557N5**
Registration date: **2015-06-27, 1394/04/06**
Registration timing: **retrospective**

Last update:
Update count: **0**

Registration date
2015-06-27, 1394/04/06

Registrant information
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Katayon Vakilian
Name of organization / entity
Arak University of Medical Sciences and Health Services
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Recruitment status
Recruitment complete

Funding source
Vice chancellor for research, Arak University of Medical Sciences

Expected recruitment start date
2014-10-07, 1393/07/15
Expected recruitment end date
2015-03-06, 1393/12/15
Actual recruitment start date
empty
Actual recruitment end date
empty
Trial completion date
empty

Scientific title
The effect of Hypericum Perforatum (Perforan) on wound healing episiotomy in women

Public title
Clinical trial the effect of Hypericum Perforatum (Perforan) on wound healing episiotomy in women
Purpose
Prevention
Inclusion/Exclusion criteria
Inclusion criteria: singleton pregnancy;primiparous; episiotomy Exclusion criteria: severe infection in 5th day after episiotomy; any acute or chronic diseases or allergy to Hypericum Perforatum
Age
From **16 years** old to **45 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
Blinding (investigator's opinion)
Double blinded
Blinding description
Placebo
Used
Assignment
Parallel
Other design features
random allocation

Secondary Ids

empty

Ethics committees

1

Ethics committee
Name of ethics committee
Ethics committee of Arak University of Medical Sciences
Street address
Educational Complex campus, Basij Square, Sardasht, arak, Markazi, Iran, Islamic Republic Of
City
Arak
Postal code
3848176941
Approval date
2014-02-03, 1392/11/14
Ethics committee reference number
92-158-8

Health conditions studied

1

Description of health condition studied
Episiotomy during vaginal delivery
ICD-10 code
o90.1
ICD-10 code description
Disruption of wound of; episiotomy

Primary outcomes

1

Description
pain
Timepoint
Before intervention, 5 and 10 day after intervention
Method of measurement
visual analogue scale

2

Description
Discharge
Timepoint
Before the intervention, 5 and 10 days after the intervention
Method of measurement
Presence of suppurative vaginal discharge

3

Description
remained sutures
Timepoint
Before the intervention, 5 and 10 days after the intervention
Method of measurement
number of remained sutures

4

Description
Dehiscence
Timepoint
10th day after intervention
Method of measurement
Incidence of wound dehiscence

5

Description
Edema
Timepoint
Before the intervention, 5 and 10 days after the intervention
Method of measurement
Diameter of edema on wound site in Centimiter

6

Description
Redness
Timepoint
Before the intervention, 5 and 10 days after the intervention
Method of measurement
Diameter of redness in millimeter

Secondary outcomes

1

Description
Infection
Timepoint
Before the intervention, 5 and 10 days after the intervention
Method of measurement
Presence of infective discharge and dehiscence of wound

Intervention groups

1

Description
intervention group:Hypericum Perforatum (Perforan) ointment, Combined Vaseline with Hypericum Perforatum extracts, 25% concentration, three times daily for 10 days, POURSINA PHARMACEUTICAL COMPANY
Category
Prevention

2

Description
Control group: Vaseline ointment, three times daily for 10 days, POURSINA PHARMACEUTICAL COMPANY
Category
Placebo

Recruitment centers

1

Recruitment center
Name of recruitment center
Ayatollah Taleghani Hospital
Full name of responsible person
Kobra Abbasinia
Street address
Imam Khomeini Street, Ayatollah Taleghani hospital, Arak.
City
Arak

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Vice Chancellor for research of Arak University of Medical Sciences
Full name of responsible person
Kobra Abbasinia
Street address
Educational Complex campus, Basij Square, Sardasht, arak.
City
Arak
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 Arak 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
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Arak University of Medical Sciences
Full name of responsible person
Kobra Abbasinia
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Person responsible for updating data

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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