

Clinical Trial Protocol

Iranian Registry of Clinical Trials

07 Jul 2026

The effect of expressive writing on postpartum depression and stress of mothers with preterm infants hospitalized in the neonatal intensive care unite

Protocol summary

Study aim

Determination and comparison of average score of postpartum depression of mothers with preterm infants hospitalized in NICU between two intervention and control group before and after doing intervention
Determination and comparison of average score of postpartum stress of mothers with preterm infants hospitalized in NICU between two intervention and control group before and after doing intervention

Design

In this study, 96 mothers with the preterm infants less than 37 weeks which have the conditions of the study and hospitalized for more than three days in the neonatal intensive care unit (NICU) of Uromieh Motahary Educational Center, are chosen. Mothers randomly assigned into two intervention and control groups and one code is allocated to each participant.

Settings and conduct

In this study, the effect of the expressive writing on postpartum depression and stress of mothers with preterm infants hospitalized in the neonatal intensive care unit (NICU) of Uromieh Motahary Educational Center will be evaluated. Postpartum depression in the control and intervention group will be measured by using Edinburgh's questionnaire on the third and 10th day, one and three months after the admission of the infant to the NICU and mothers' stress in both groups will be measured by using Parents' stress in the NICU questionnaire On the third and 10th day after the admission and Cohen' perceived stress- 14 items questionnaire in one and three months after the admission of the infant in the NICU.

Participants/Inclusion and exclusion criteria

Inclusion criteria: attending the NICU at least three days a week.

Intervention groups

Mothers in the intervention group will be asked to write

their deepest emotions and thoughts about their infants' birth and hospitalized in a private place, regardless of grammar and sentence structure, three times for 15 minutes each time. The duration of the intervention will be once a day from third to 10th day of hospitalized to the NICU. Mothers in the control group will receive the routine care.

Main outcome variables

postpartum depression; postpartum stress

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20170726035328N2**

Registration date: **2018-02-20, 1396/12/01**

Registration timing: **registered_while_recruiting**

Last update: **2018-02-20, 1396/12/01**

Update count: **0**

Registration date

2018-02-20, 1396/12/01

Registrant information

Name

Naemeh VatankhahAlamdary

Name of organization / entity

Uromieh University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 44 3223 4897

Email address

vatankhah.na@umsu.ac.ir

Recruitment status

Recruitment complete

Funding source

Investigator

Expected recruitment start date

2017-07-25, 1396/05/03

Expected recruitment end date

2018-03-21, 1397/01/01

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of expressive writing on postpartum depression and stress of mothers with preterm infants hospitalized in the neonatal intensive care unite

Public title

Postpartum depression and stress of mothers with preterm infants

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria:

Having at least a low secondary education Attendance at the NICU for at least three days a week

Exclusion criteria:

previous record of newborns admitted to the neonatal intensive care unite (NICU) acute stressful conditions, such as death of loved ones within the past six months history of depression and hospitalization and antidepressant use within the past six month history of psychoactive use and lack of drug addiction during the past six months genetic disease in the newborn

Age

No age limit

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 96

Randomization (investigator's opinion)

Randomized

Randomization description

A total number of 96 envelopes containing number one or two was designed for mothers to choose from. mothers who received number one were assigned to intervention group (48 mothers) and the others were assigned to control group (48 mothers).

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 Uromieh University of Medical Sciences

Street address

Jahad Street, Urmia University of Medical Sciences, Uromieh

City

Urmia

Province

West Azarbaijan

Postal code

57147-83734

Approval date

2017-06-07, 1396/03/17

Ethics committee reference number

IR.UMSU.rec.1396.103

Health conditions studied

1

Description of health condition studied

postpartum depression and stress

ICD-10 code

F53- F43.1

ICD-10 code description

Mental and behavioural disorders associated with the puerperium, not elsewhere classified - Post-traumatic stress disorder

Primary outcomes

1

Description

postpartum depression

Timepoint

The third day, 10th day, one and three months after the admission of the baby to the neonatal intensive care unite

Method of measurement

Edinburgh questionnaire

2

Description

Stress of mothers' with preterm infants

Timepoint

The third day, 10th day, one and three months after the admission of the baby to the neonatal intensive care unite

Method of measurement

Parental stressor scale for neonatal intensive care units

and Cohen' perceived Stress - 14 items questionnaire

Secondary outcomes

empty

Intervention groups

1

Description

intervention group: Mothers will be asked to write in their own private place without attention to grammar about their deepest emotions and their work on the inconvenient event. Writes three times a week and each time for 15 minutes. The period from the third day until the tenth day will be admitted to NICU.

Category

N/A

2

Description

Control group: The control group will receive the routine care.

Category

Behavior

Recruitment centers

1

Recruitment center

Name of recruitment center

Urmia Motahary Educational Center

Full name of responsible person

Naemeh VatankhahAlamdary

Street address

Ayatollah Kashani Ave: Motahari Educational Center

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Web page address

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Oroumia University of Medical Sciences

Full name of responsible person

Dr. Iraj Mohebbi

Street address

Jahad Street, Urmia University of Medical Sciences

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Web page address

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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

Oroumia 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

Person responsible for general inquiries

Contact

Name of organization / entity

Assistance for Research and Technology of Uromieh University of Medical Sciences

Full name of responsible person

Dr. Iraj Mohebbi

Position

Assistant for Research and Technology

Latest degree

Specialist

Other areas of specialty/work

Occupational Medicine

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Person responsible for scientific inquiries

Contact

Name of organization / entity

Urmia University of Medical Sciences

Full name of responsible person

Soheila Rabeepoor

Position

Faculty member

Latest degree

Ph.D.

Other areas of specialty/work

Midwifery

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Web page address

Sharing plan

Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

Study Protocol

Undecided - It is not yet known if there will be a plan to make this available

Statistical Analysis Plan

Not applicable

Informed Consent Form

Undecided - It is not yet known if there will be a plan to make this available

Clinical Study Report

Not applicable

Analytic Code

Undecided - It is not yet known if there will be a plan to make this available

Data Dictionary

Undecided - It is not yet known if there will be a plan to make this available

Title and more details about the data/document

Because the paper out from the thesis will probably be published until July 2018

When the data will become available and for how long

I have not decided yet - its release plan is still unclear

To whom data/document is available

I have not decided yet - its release plan is still unclear

Under which criteria data/document could be used

I have not decided yet - its release plan is still unclear

From where data/document is obtainable

I have not decided yet - its release plan is still unclear

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

I have not decided yet - its release plan is still unclear

Comments