

Clinical Trial Protocol

Iranian Registry of Clinical Trials

19 Jun 2026

The effect of motivational interviewing throughout pregnancy on birth preference: a randomized controlled trial

Protocol summary

Study aim

The present study aims to investigate the effect of motivational interviews conducted during pregnancy on birth preference.

Design

The design is of a randomized controlled experimental type. Obstetrics polyclinics of a university and a state hospital, Çanakkale, Turkey. A total of 100 pregnant women, 50 in the experimental group and 50 in the control group, were included in the study. Three motivational interviews, initiated during pregnancy (Week 26-28) and continued in the postnatal period, were conducted with the experimental group. Nutrition training was provided for the control group during pregnancy.

Settings and conduct

The study population consisted of all pregnant women who referred to obstetrics polyclinics of Çanakkale Onsekiz Mart University Healthcare, Practice and Research Hospital and Çanakkale State Hospital during 01 March 2019-02 March 2020.

Participants/Inclusion and exclusion criteria

Inclusion Criteria - Primigravid status (women experiencing their 1st pregnancy), - Married (civil marriage) pregnant women in gestational week 26-28, - 50% of the experimental group and the control group consisted of pregnant women from rural areas, - Pregnant women no physical disability, - No diagnosis of a psychiatric condition (verbally confirmed), - Subjects willing to participate in the study, - Those without pregnancy considered 'risky', - Pregnant women without planned cesarean section Exclusion criteria - know how to read and write - not be a foreign national

Intervention groups

Motivational interviews (face to face)

Main outcome variables

Mother's perception of self-efficacy regarding labor, fear of childbirth, intention, significance, confidence-efficacy parameters of motivational interviews, birth preference

General information

Reason for update

Acronym

IRCT registration information

IRCT registration number: **IRCT20220509054795N1**

Registration date: **2022-05-12, 1401/02/22**

Registration timing: **retrospective**

Last update: **2022-05-12, 1401/02/22**

Update count: **0**

Registration date

2022-05-12, 1401/02/22

Registrant information

Name

Eda CANGÖL

Name of organization / entity

Çanakkale Onsekiz Mart Üniversitesi

Country

Turkey

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+90 286 218 13 97

Email address

edacangol@comu.edu.tr

Recruitment status

Recruitment complete

Funding source

Expected recruitment start date

2019-03-01, 1397/12/10

Expected recruitment end date

2020-03-02, 1398/12/12

Actual recruitment start date

2019-03-01, 1397/12/10

Actual recruitment end date

2020-03-02, 1398/12/12

Trial completion date

empty

Scientific title

The effect of motivational interviewing throughout pregnancy on birth preference: a randomized controlled trial

Public title

The effect of motivational interviewing birth preference

Purpose

Education/Guidance

Inclusion/Exclusion criteria

Inclusion criteria:

- Primigravid status (women experiencing their 1st pregnancy) - Married (civil marriage) pregnant women in gestational week 26-28, - 50% of the experimental group and the control group consisted of pregnant women from rural areas, - Pregnant women no physical disability, - No diagnosis of a psychiatric condition (verbally confirmed), - Subjects willing to participate in the study, - Those without pregnancy considered 'risky', - Pregnant women without planned cesarean section

Exclusion criteria:

know how to read and write not be a foreign national

Age

No age limit

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **100**

Actual sample size reached: **63**

Randomization (investigator's opinion)

Randomized

Randomization description

A total of 100 pregnant women who met the inclusion criteria, i.e. those who referred to obstetrics polyclinics of Çanakkale Onsekiz Mart University Healthcare, Practice and Research Hospital and Çanakkale State Hospital during 01 March 2019-02 March 2020. Were assigned to the experimental group or the control group (50 women in each group) by randomization using the random sampling method. Randomization was performed by assigning numbers to the pregnant women according to their order of presentation using a table of random numbers to assign the first number in the experimental group and the second number in the control group.

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

Approval was obtained from Uşak University Ethics Committee

Street address

1 September Campus/Usak University

City

Uşak

Postal code

64000

Approval date

2016-04-06, 1395/01/18

Ethics committee reference number

local; ethics registration code: 97627247-050.99-11461

Health conditions studied

1

Description of health condition studied

None

ICD-10 code

ICD-10 code description

Primary outcomes

1

Description

Mother's perception of self-efficacy regarding labor

Timepoint

Pretest and posttest

Method of measurement

Self-Sufficiency in Labor Scale-Short Form

2

Description

The intent of motivational Interview, significance, confidence-efficacy parameters

Timepoint

Pretest and post-test

Method of measurement

MI Rating Scale

3

Description

fear of childbirth

Timepoint

Pretest and post-test

Method of measurement

Delivery Expectancy/Experience Questionnaire (W-DEQ), version A

Secondary outcomes

1

Description

Mother's perception of self-efficacy regarding labor

Timepoint

after 1 month

Method of measurement

Self-Sufficiency in Labor Scale-Short Form

2

Description

The intent of motivational Interview, significance, confidence-efficacy parameters

Timepoint

after 1 month

Method of measurement

MI Rating Scale

3

Description

fear of childbirth

Timepoint

after 1 month

Method of measurement

Wijma Delivery Expectancy/Experience Questionnaire (W-DEQ), version B

4

Description

Mode of Delivery

Timepoint

after 1 month

Method of measurement

Postnatal Maternal Information Form

Intervention groups

1

Description

To support vaginal delivery, three motivational interviews, initiated during pregnancy (Week 26-28) and continued in the postnatal period, were conducted with the experimental group. Motivational interviews in the third trimester were preferred as this is a period close to delivery. The interviews were conducted individually since the motivational interviewing technique is an individual-based method. Timeline of Motivational Interviews (MIs) was as follows; The first interview: Antenatal week 26-28 The second interview: Antenatal week 32-34 The third interview: Postnatal day 1

Category

Behavior

2

Description

Since it would not be ethically appropriate to apply no intervention in the control group, individual nutrition training was provided during pregnancy in this group.

The content of this training included managing nutrition during pregnancy, important nutrients during pregnancy, weight gain during pregnancy, and nutritional recommendations in physiological changes during pregnancy.

Category

Other

Recruitment centers

1

Recruitment center

Name of recruitment center

Faculty of Health Sciences, Çanakkale Onsekiz Mart University, Turkey

Full name of responsible person

Eda Cangöl

Street address

Çanakkale Onsekiz Mart University Terzioğlu Campus, Faculty of Health Sciences

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Sponsors / Funding sources

1

Sponsor

Name of organization / entity

Koç University

Full name of responsible person

Ayşe Ferda Ocakçı

Street address

Topkapı

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İstanbul

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34010

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Grant name

Grant code / Reference number

2019.1.6

Is the source of funding the same sponsor organization/entity?

Yes

Title of funding source

Koç University

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
Çanakkale Onsekiz Mart University
Full name of responsible person
Eda Cangöl
Position
Assistant Prof., PhD
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Person responsible for updating data

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Sharing plan

Deidentified Individual Participant Data Set (IPD)

Undecided - It is not yet known if there will be 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

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

Informed Consent Form

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

Clinical Study Report

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

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