

Clinical Trial Protocol

Iranian Registry of Clinical Trials

27 Jun 2026

The effect of healthy nutrition training and physical activity with pedometer on weight changes and weight-efficacy of life style woman with high body mass index in postpartum period

Protocol summary

Summary

In this study, two groups, randomized clinical trial that aim of determining the effect of healthy nutrition training and physical activity with Pedometer on weight changes and weight-efficacy of life style woman with high body mass index in postpartum period. Sample size includes 64 obese and overweight women who go to the health center at 6-16 weeks postpartum. For sampling, Mashhad Center No. 2 and No. 3 were selected. The among their subsidiary centers, randomly, four centers are selected as interventions and four centers are selected as controls. The training is carried out by a researcher in groups of 5 to 7. Teaching is a lecture method, a question and answer that takes place in two sessions (90-60) minutes a week, once a week. Session One: Measurement of anthropometric index and body composition, completion of questionnaires including Clark's Weight Lifestyle Self-Efficacy Questionnaire, Physical Activity Indicator Questionnaire, an explanation on how to fill out a checklist of food and daily physical activity, then training Regarding physical activity and its benefits during postpartum period, and the method of using the pedometer is given. Second session: Training on healthy nutrition counseling and people will be taught to choose their food according to the food substitution list At the end of the session, a healthy lifestyle education booklet (with content of physical activity and healthy nutrition advice) will be presented. In the intervention group, after the completion of the training in the 2nd and 6th weeks, a telephone call is made to follow up the research units. The control group will receive routine care. Maternal weight at the end of weeks 4 and 8 is monitored in both groups, then the mean weight changes and lifestyle self-efficacy related to weight at the end of week 8 will be compared between the two groups.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2017062834339N1**
Registration date: **2017-08-31, 1396/06/09**
Registration timing: **retrospective**

Last update:

Update count: **0**

Registration date

2017-08-31, 1396/06/09

Registrant information

Name

Hoda Naderi

Name of organization / entity

Mashhad University of Medical Sciences

Country

Iran (Islamic Republic of)

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+98 76 3537 7297

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naderih931@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

Mashhad University of Medical Sciences

Expected recruitment start date

2017-01-14, 1395/10/25

Expected recruitment end date

2017-08-16, 1396/05/25

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of healthy nutrition training and physical activity with pedometer on weight changes and weight-efficacy of life style woman with high body mass index in postpartum period

Public title

Effect of training on weight changes and self-efficacy of women's lifestyle with high body mass index in postpartum period

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: have written consent to participate in research; to visit the health center at 6-16 weeks after delivery; have physical activity level poor (score between 20-39) or inactive (score Less than 20); the pre-pregnancy body mass index is between (25-34/9); after delivery body mass index is between (25-34/9). Exclusion criteria: do not participate in one of the training sessions; walking has not been done in 3 continuous Sessions or 5 discontinuous sessions; in more than 5 sessions, the number of steps recorded is less than 100-120 steps per minute.

Age

From **18 years** old to **45 years** old

Gender

Female

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: **64**

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Single blinded

Blinding description

Placebo

Not used

Assignment

Parallel

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Mashhad University of Medical Sciences

Street address

Qaryashi Building, University Street, Mashhad Holy

City

Mashhad

Postal code

Approval date

2016-12-10, 1395/09/20

Ethics committee reference number

IR.MUMS.REC.1395.429

Health conditions studied

1

Description of health condition studied

Overweight and Obesity after childbirth

ICD-10 code

E66.8

ICD-10 code description

Other obesity

Primary outcomes

1

Description

Weight changes

Timepoint

Before the intervention, 4 weeks after the intervention, 8 weeks after the intervention

Method of measurement

Scales

2

Description

Weight-Efficacy of Life style

Timepoint

Before intervention, 8 weeks after intervention

Method of measurement

Questionnaire

Secondary outcomes

1

Description

Physical Activity Index

Timepoint

Before the intervention, 8 weeks after the intervention

Method of measurement

Questionnaire

2

Description

Waist to Hip Ratio

Timepoint

Before the intervention, 4 weeks after the intervention, 8 weeks after the intervention

Method of measurement

Meter

3

Description

Hip Circumference

Timepoint

Before the intervention, 4 weeks after the intervention, 8 weeks after the intervention

Method of measurement

Meter

4

Description

Waist Circumference

Timepoint

Before the intervention, 4 weeks after the intervention, 8 weeks after the intervention

Method of measurement

Meter

5

Description

Body Composition

Timepoint

Before the intervention, 8 weeks after intervention

Method of measurement

Electrical Impedance Analysis Device

6

Description

Body Mass Index

Timepoint

Before the intervention, 4 weeks after the intervention, 8 weeks after the intervention

Method of measurement

Scales and Meters

Intervention groups

1

Description

In the intervention group, in addition to the standard care provided in the Health centers, a healthy nutrition education and physical activity with a pedometer will be held by the researcher in groups of 5 to 7. Teaching is a lecture method, a question and answer that takes place in two sessions (90-60) minutes, once a week. At the end of the training sessions, a healthy lifestyle booklet will be given to the mothers with the content of healthy nutrition and physical activity after childbirth.

Category

Lifestyle

2

Description

The control group will receive standard care in Health centers.

Category

Lifestyle

Recruitment centers

1

Recruitment center**Name of recruitment center**

Health Center No. 2

Full name of responsible person

Hoda Naderi

Street address

Beginning of Boulevard of Unity, next to Wahdat Park, Health Center No. 2

City

Mashhad

2

Recruitment center**Name of recruitment center**

Health Center No. 3

Full name of responsible person

Hoda naderi

Street address

Akhund Khorasani Ave. 24

City

Mashhad

Sponsors / Funding sources

1

Sponsor**Name of organization / entity**

Mashhad University of Medical Sciences

Full name of responsible person

Research Mashhad University of Medical Sciences

Street address

Mashhad, Daneshgah street, Mashhad University of Medical Science

City

Mashhad

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

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

Faculty of nursing and midwifery of Mashhad

Full name of responsible person

Masoumeh Kordi

Position

Master of midwifery

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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