

Clinical Trial Protocol

Iranian Registry of Clinical Trials

18 Jun 2026

The effect of multi-sensory stimulation on the development of neuromuscular, weight and physiological parameters in preterm infants

Protocol summary

Summary

Preterm birth is the most common cause of death in babies and as one of the risk factors is intended for developmental disabilities Which can lead to long-term effects on the Neuromuscular development. Since the original development is done during pregnancy, especially during the last months and weeks and the babies are born too early, so had ample opportunity for the development of critical systems developmental and physiological problems may have multiple. Multi-sensory stimulation is relatively new intervention to simulate the environment in the first weeks of intrauterine life; to maintain and facilitate the development in premature infants. This study aimed to effect of multi-sensory stimulation on the development of neuromuscular, weight and physiological parameters in preterm infants. In this clinical trial, 80 preterm infants with gestational age between 32 and 36 weeks were assigned randomly into two groups and controlled multisensory stimulation. Groups intervention, multisensory stimulation program for 12 minutes a day, 5 times a week at a time until discharge and the control group received only usual care. Physiological parameters (respiration, heart rate, blood pressure and oxygen saturation) using the monitoring and the development of neuromuscula using the Infant Neurological International Battery before and after the intervention, as well as daily weight gain the researcher measured.

General information

Acronym

IRCT registration information

IRCT registration number: **IRCT2016073114454N2**

Registration date: **2016-08-17, 1395/05/27**

Registration timing: **registered_while_recruiting**

Last update:

Update count: **0**

Registration date

2016-08-17, 1395/05/27

Registrant information

Name

Fatemeh Nasimi

Name of organization / entity

Mashhad University of Medical Sciences

Country

Iran (Islamic Republic of)

Phone

+98 72 1222 4259

Email address

nasimif911@mums.ac.ir

Recruitment status

Recruitment complete

Funding source

Vice Chancellor for Research, Jahrom University of Medical Sciences

Expected recruitment start date

2016-05-04, 1395/02/15

Expected recruitment end date

2016-09-02, 1395/06/12

Actual recruitment start date

empty

Actual recruitment end date

empty

Trial completion date

empty

Scientific title

The effect of multi-sensory stimulation on the development of neuromuscular, weight and physiological parameters in preterm infants

Public title

The effect of multi-sensory stimulation on the development of neuromuscular, weight and physiological parameters in preterm infants

Purpose

Prevention

Inclusion/Exclusion criteria

Inclusion criteria: parental consent; infants age between 37-30 weeks; hemodynamic stability (HR=120-160, RR=30-50, SPO2=85%-95% and keep axillary temperature range of 37-36 Centigrade degrees); Apgar scores of 7 or more on the first and fifth minutes of birth; spontaneous ventilation and no need for ventilation; lack of congenital malformations. Exclusion criteria: lack of consent to continue working for any reason; ventilation need during the study; infants with recurrent Apnea; seizures during the study.

Age

No age limit

Gender

Both

Phase

N/A

Groups that have been masked

No information

Sample size

Target sample size: 80

Randomization (investigator's opinion)

Randomized

Randomization description

Blinding (investigator's opinion)

Not blinded

Blinding description

Placebo

Not used

Assignment

Other

Other design features

Secondary Ids

empty

Ethics committees

1

Ethics committee

Name of ethics committee

Jahrom University of Medical Sciences

Street address

Jahrom University of Medical Sciences, Motahari Street, Jahrom

City

Jahrom

Postal code

Approval date

2016-03-09, 1394/12/19

Ethics committee reference number

IR.JUMS.REC.1394.207

Health conditions studied

1

Description of health condition studied

Preterm infants

ICD-10 code

P07.3

ICD-10 code description

Other preterm infants

Primary outcomes

1

Description

Heart Rate

Timepoint

Before and after the intervention.

Method of measurement

with monitoring

2

Description

Blood Pressure

Timepoint

Before and after the intervention.

Method of measurement

with monitoring

3

Description

Oxygen Saturation

Timepoint

Before and after the intervention.

Method of measurement

with monitoring

4

Description

Respiration Rate

Timepoint

Before and after the intervention.

Method of measurement

with monitoring

Secondary outcomes

1

Description

Neuromuscular development

Timepoint

Before and after the intervention.

Method of measurement

Using tool Infant Neurological International Battery

2

Description

Weight gain

Timepoint

Daily
Method of measurement
Using the Balance

Intervention groups

1

Description
The intervention group receive multi-sensory stimulation for 12 minutes every day, 5 times a week until discharge

Category
Prevention

2

Description
The control group received usual care

Category
N/A

Recruitment centers

1

Recruitment center
Name of recruitment center
Neonatal Intensive Care Unit, Motahari Hospital
Full name of responsible person
Fatemeh Nasimi
Street address
Neonatal Intensive Care Unit, Motahari Hospital,
Motahari Street, Jahrom
City
Jahrom

Sponsors / Funding sources

1

Sponsor
Name of organization / entity
Vice Chancellor for Research, Jahrom University of
Medical Sciences
Full name of responsible person
Dr. Kavus Solhjo
Street address
Jahrom University of Medical Sciences, Motahari
Street, Jahrom
City
Jahrom
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, Jahrom 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
Jahrom University of Medical Sciences
Full name of responsible person
Fatemeh Nasimi
Position
MSc in Nursing, Neonatal Intensive Care
Other areas of specialty/work
Street address
School of Nursing and Allied Health, Motahari Street,
Jahrom
City
Jahrom
Postal code
Phone
+98 71 5434 1501
Fax
Email
Nasimif@yahoo.com
Web page address

Person responsible for scientific inquiries

Contact
Name of organization / entity
Jahrom University of Medical Sciences
Full name of responsible person
Fatemeh Nasimi
Position
MSc in Nursing, Neonatal Intensive Care
Other areas of specialty/work
Street address
School of Nursing and Allied Health, Motahari Street,
Jahrom
City
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Phone
+98 71 5434 1501
Fax
Email
Nasimif@yahoo.com
Web page address

Person responsible for updating data

Contact
Name of organization / entity
Jahrom University of Medical Sciences
Full name of responsible person

Fatemeh Nasimi

Position

MSc in Nursing, Neonatal Intensive Care

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Street address

School of Nursing and Allied Health, Motahari Street,
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+98 71 5434 1501

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Email

Nasimif@yahoo.com

Web page address

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