

# Clinical Trial Protocol

## Iranian Registry of Clinical Trials

29 Jun 2026

### The Effect of Massage on Neonatal Hyperbilirubinemia in Neonatal Intensive Care Unit

#### Protocol summary

##### Study aim

Determining the effect of massage on Hyperbilirubinemia in infants admitted to the neonatal intensive care unit

##### Design

Randomised, superiority, parallel group trial with blinded outcome assessment. Randomisation was centralised and computerised with concealed randomisation sequence carried out at an external site.

##### Settings and conduct

This clinical trial study was conducted in two intervention and control groups on 70 admitted infants in the intensive care unit. Eligible babies will be assigned to intervention and control groups using the random block. Both mothers and infants mothers will be given individual education about breastfeeding and jaundice. Control neonates will receive only routine care. However, the newborns in the intervention group will be massaged in addition to receiving usual care for 5 days, 3 times a day, and 15 minutes each, based on the technique of Fidel's mother (after training by the researcher). The amount of skin infant bilirubin (Cutaneous Billirubin Trans = TCB) is measured daily for five days (although the measurement of bilirubin is not due to research, but according to the routine of the hospital for newborns with jaundice on a daily basis), the first time the meconium is excreted and The number of meconium in 24 hours and in both groups is recorded for five days. During the study, the person receiving the bilirubin level is blind to the groups.

##### Participants/Inclusion and exclusion criteria

Infants with fetal age 37 to 42 weeks without congenital anomalies. Lack of knowledge of known gastrointestinal and central nervous system problems. Babies weighing over 2500 to 4000 grams. Babies with a one-day age who are not hospitalized at least until the fifth day of birth, and do not have a massage ban.

##### Intervention groups

Neonates with Hyperbilirubinemia who are admitted to the intensive care unit

#### Main outcome variables

Evaluation of blood bilirubin reduction in patients in control and intervention groups.

#### General information

##### Reason for update

##### Acronym

##### IRCT registration information

IRCT registration number: **IRCT20150621022852N2**

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

Registration timing: **retrospective**

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

Update count: **0**

##### Registration date

2018-02-20, 1396/12/01

##### Registrant information

##### Name

##### Name of organization / entity

##### Country

Iran (Islamic Republic of)

##### Phone

+98 44429613

##### Email address

dr.s.asaadi@sbm.ac.ir

##### Recruitment status

**Recruitment complete**

##### Funding source

##### Expected recruitment start date

2018-01-21, 1396/11/01

##### Expected recruitment end date

2018-02-20, 1396/12/01

##### Actual recruitment start date

empty

##### Actual recruitment end date

empty

**Trial completion date**

empty

**Scientific title**

The Effect of Massage on Neonatal Hyperbilirubinemia in Neonatal Intensive Care Unit

**Public title**

The Effect of Massage on Neonatal Jaundice in Neonatal Intensive Care Unit

**Purpose**

Treatment

**Inclusion/Exclusion criteria****Inclusion criteria:**

Normal Birth Weight:2500 to 4000 grams No history of any acute congenital disorder Negative symptom or sign of Sepsis or infection

**Exclusion criteria:****Age**From **1 day** old to **5 days** old**Gender**

Both

**Phase**

N/A

**Groups that have been masked**

- Outcome assessor

**Sample size**Target sample size: **64****Randomization (investigator's opinion)**

Randomized

**Randomization description**

The method of random sampling is blocking. To determine whether each child is in the control group or experience the use of the blue-ring method, it is determined by the throwing of the six-sided dice that controls or experiences from the first to the last one. To implement the blocking method, two modes A and B are considered so that the six blocks are as follows from the different types of placement of these two modes together: 1)ABAB 2)AABB 3) ABBA 4)BBAA 5)BABA 6) BAAB Then, with the throwing of six-sided dice, ten blocks of four (ten modes for the direction of 70 children) are obtained randomly according to the above conditions; each of the selected letters will be considered for children 1-42. A and B modes are also determined randomly (lottery) as controls or as experiences (eg, mode A as control and B as experience). In addition, with the throw of dice, the order of the block is specified. Draw and throw dice in the presence of a supervisor or consultant professor. Examples of dice throwing scenarios, for example, are achieved, and the following results are obtained and the assignment of samples to the experiment group and control will be based on it.

**Blinding (investigator's opinion)**

Single blinded

**Blinding description**

A person who performs the Bilirubin test will be Blind to the groups.

**Placebo**

Not used

**Assignment**

Parallel

**Other design features****Secondary Ids**

empty

**Ethics committees****1****Ethics committee****Name of ethics committee**

Ethics committee of Tehran University of Medical Sciences

**Street address**

Poursina st,Enghelab Ave,Tehran,Iran

**City**

Tehran

**Province**

Tehran

**Postal code**

34252523525

**Approval date**

2017-06-19, 1396/03/29

**Ethics committee reference number**

IR.TUMS.FMN.REC.1396.2684

**Health conditions studied****1****Description of health condition studied**

Hyper bilirubinemia

**ICD-10 code**

P59.9

**ICD-10 code description**

Neonatal jaundice, unspecified Physiological jaundice (intense)(prolonged) NOS

**Primary outcomes****1****Description**

level of bilirubin in serum

**Timepoint**

in first five days of intervention

**Method of measurement**

Measurement of serum bilirubin levels in blood samples of patients.

**Secondary outcomes**

empty

**Intervention groups****1****Description**

Intervention group:The intervention group of the

newborns in the intervention group, in addition to receiving usual care, will be given massage for a period of five days, 3 times a day, and 15 minutes each time, according to the technique of the Fidel Massage. The practice of massage is taught to mothers by the trained researcher in accordance with the guidelines of the International Association of Infant Massage (IAIM). Babies will be massaged during the first day by a researcher or mother's participation. After assessing and assuring the mother's learning and mastery of massage; Mothers from the first to fifth days do their massage therapy on their babies, and their performance is monitored by the researcher in a timely and correct manner. Massage after washing and Warming the hands is done in a warm environment. Each massage is composed of 3 phases of 5 minutes and a total of 15 minutes. In the first and third phases, the baby is placed in the abdomen and the massage is done as follows: 1) from the top to the neck and vice versa, 2 ) From the neck to the shoulders and vice versa, 3) from the back to the waist and vice versa, 4) from the shoulder to the hand, then in the direction of return in both hands, 5) from the thigh to the ankle and then in the direction of return in both Leg. In the second stage, motor stimulation involves extensibility and flexion of the organs in the supine position.

**Category**

Treatment - Other

**2**

**Description**

Control group: In the control group after birth and transfer to the neonatal ward, they will be evaluated and qualified to join the study by researchers. Then, the goal of the study will explained to mothers and after obtaining consent and completing the informed consent form they will be enrolled in study. All mother will be inform about breastfeeding and jaundice. The control group will receive only routine care, such as umbilical cord care, audiometry, and vaccination. During study days control group will be assess about same study variables that will be check in intervention group.

**Category**

N/A

**Recruitment centers**

**1**

**Recruitment center**

**Name of recruitment center**

Arash Hospital

**Full name of responsible person**

Zeinab Shokati mir

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Eastern 162th St.,3rd Sq.,Tehran Pars District 4, Tehran

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sina.asadi90@yahoo.com

**Sponsors / Funding sources**

**1**

**Sponsor**

**Name of organization / entity**

Tehran University of Medical Sciences

**Full name of responsible person**

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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**

Tehran 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**

Tehran University of Medical Sciences

**Full name of responsible person**

Zeinab Shokati Mir

**Position**

Nurse

**Latest degree**

Bachelor

**Other areas of specialty/work**

Nursery

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Nursing And Midwifery School, Tehran University Of Medical Science, Tehran Province, Tehran, District 6,

Nosrat St  
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## Person responsible for scientific inquiries

### Contact

**Name of organization / entity**  
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**Position**  
Nurse  
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Bachelor  
**Other areas of specialty/work**  
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## Person responsible for updating data

### Contact

**Name of organization / entity**  
Tehran University of Medical Sciences  
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## Sharing plan

### Deidentified Individual Participant Data Set (IPD)

Yes - There is a plan to make this available

### Study Protocol

No - There is not a plan to make this available

### Statistical Analysis Plan

No - There is not a plan to make this available

### Informed Consent Form

Yes - There is a plan to make this available

### Clinical Study Report

No - There is not a plan to make this available

### Analytic Code

No - There is not a plan to make this available

### Data Dictionary

No - There is not a plan to make this available

### Title and more details about the data/document

Data from the study under the supervision of the scientific and practical scientist will be the study of collecting and sharing the results and raw data with full supervision.

### When the data will become available and for how long

From the beginning of the study to the publication of the results.

### To whom data/document is available

Researcher and supervisor of the plan and the officials of the Ethics Committee of the University

### Under which criteria data/document could be used

To verify the results and verify the accuracy of the documentation.

### From where data/document is obtainable

Must be referred to the scientific and practical planner.

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

Call the scientific and practical scientist through the phone or email.

### Comments