# Mechanical massage using an electric massage chair and the onset of coming in of breastmilk in first-time mothers

Submission date	<b>Recruitment status</b> No longer recruiting	[X] Prospectively registered		
16/02/2024		[X] Protocol		
Registration date	Overall study status	Statistical analysis plan		
19/02/2024	Completed	Results		
Last Edited	Condition category	Individual participant data		
27/02/2024	Pregnancy and Childbirth	Record updated in last year		

#### Plain English summary of protocol

Background and study aims

Hand (manual) massage especially along the back and foot after birth has been shown to speed up the first coming in of breast milk and of milk production. Hand massage is time consuming and a learnt skill. The electric mechanical massage chair is reusable, simple to operate and consistent in the massage it delivers but it has not been explored if it is effective. Delayed or perceived inadequate milk production can cause problems with the start or maintenance of breastfeeding of the baby.

WHO and UNICEF recommend that infants be exclusively breastfed for the first 6 months of life.

First time mothers take a longer time to first breastfeeding attempt, are more likely to have eight or fewer feeding attempts in the first 24 hours, to report early breastfeeding problems, are mixed feeding at hospital discharge and less likely to breastfeed through 6 months.

This study aims to compare the effect of two massage programs of an electric massage chair to evaluate if they work on to hasten the onset of lactation (lactogenesis stage II).

#### Who can participate?

First time mothers aged 18-45 years, without any serious illness or chronic disease, with a straightforward vaginal delivery, a healthy baby and planning to breastfeed.

#### What does the study involve?

Participants will be randomised into 2 groups:

(1) The intervention group will be given a back massage program using the electric massage chair twice daily, for 20 minutes each session, starting from 3-6 hours postdelivery, until the day of discharge.

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(2) The control group will be given a calf massage program using the electric massage chair twice

daily, for 20 minutes each session, starting from 3-6 hours postdelivery, until the day of discharge.

Follow up for 10 days post-delivery.

What are the possible benefits and risks of participating?

The allocated massage may hasten, have no effect or delay onset of lactation. The massages may increase relaxation or cause discomfort. Use of massage chair is not expected to cause major adverse consequences.

Where is the study run from? University of Malaya Medical Centre (UMMC) (Malaysia)

When is the study starting and how long is it expected to run for? January 2024 to December 2024

Who funds this study? The Department of Obstetrics & Gynaecology, UMMC (Malaysia)

Who is the main contact person?

Dr Nurul Atiqah Binti Radzali, atiqah.radzali@ummc.edu.my

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# Contact information

#### Type(s)

Public, Scientific, Principal investigator

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# Additional identifiers

## Clinical Trials Information System (CTIS)

Nil known

## ClinicalTrials.gov (NCT)

Nil known

#### Protocol serial number

MREC ID NO: 2023713-12663

# Study information

#### Scientific Title

Mechanical Massage By An Electric Massage Chair on Lactogenesis Stage II (Onset of Lactation) In First-Time Mothers: A Randomised Controlled Trial

## **Study objectives**

Mechanical back massage will hasten the onset of lactation (lactogenesis stage II) in first-time mothers after a vaginal delivery

## Ethics approval required

Ethics approval required

## Ethics approval(s)

approved 06/02/2024, Medical Research Ethics Committee University Of Malaya Medical Centre (Lembah Pantai, Kuala Lumpur, Kuala Lumpur, 59100, Malaysia; +60 03-79493209; ummc-mrec@ummc.edu.my), ref: MREC ID NO: 2023713-12663

# Study design

Interventional randomized controlled trial

# Primary study design

Interventional

# Study type(s)

Quality of life, Efficacy

# Health condition(s) or problem(s) studied

Onset of lactation in first time mothers

#### **Interventions**

Participants after written informed consenting will be randomised into two groups:

1. The intervention group will be given a back massage program using the electric massage chair twice daily, for 20 minutes each session, starting from 3-6 hours postdelivery, until the day of discharge.

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2. The control group will be given a calf massage program using the electric massage chair twice daily, for 20 minutes each session, starting from 3-6 hours postdelivery, until the day of discharge.

Randomisation sequence will be generated separately using an online generator (https://www.sealedenvelope.com/simple-randomiser/v1/lists), in 1 to 1 ratio and random blocks of 4 or 8, by a co-investigator who will not be involved in the recruitment process. Randomisation will be implemented using strict sequential assignment of the lowest-numbered remaining sealed envelope to the newest recruit.

#### Intervention Type

Device

#### Phase

Not Applicable

## Drug/device/biological/vaccine name(s)

Mechanical Electric Massage Chair

#### Primary outcome(s)

1. Time from birth to onset of lactation (lactogenesis stage II) [As reported by the mother on daily assessment]

Breast fullness is rated on a 5-point scale: from 1 (no change) to 3 (noticeably full) to 5 (uncomfortably full). Onset of lactation is defined as the first breast fullness score of 3 or greater as reported by the mother.

# Key secondary outcome(s))

- 1. Maternal satisfaction with their assigned mechanical massage chair experience using a 11 point (0-10) numerical rating scale prior to discharge. [As reported by the mother]
- 2. Baby's breastfeeding performance using Infant Breastfeeding Assessment Tool (IBFAT) prior to discharge. [As reported by the mother]
- 3. Baby's breastfeeding performance using Infant Breastfeeding Assessment Tool (IBFAT) at Day 10 after birth. [As reported by the mother]
- 4. Exclusive breastfeeding at Day 10 after birth. [As reported by the mother] IBFAT score of 10–12 indicates successful breastfeeding, a score of 7–9 relatively successful breastfeeding and a score of 0–6 unsuccessful breastfeeding.

#### Completion date

30/12/2024

# **Eligibility**

# Key inclusion criteria

- 1. Primiparous
- 2. Spontaneous vaginal deliveries mother without complications
- 3. 18 45 years
- 4. Singleton pregnancy
- 5. Term pregnancy > 37 weeks
- 6. Birth weight > 2500g
- 7. No serious illness/chronic disease
- 8. Ability to read and write, acceptable ability of listening and speaking to answer the questions

9. Intention to breastfeed10. Can communicate in Malay or English

## Participant type(s)

**Patient** 

## Healthy volunteers allowed

No

#### Age group

Adult

## Lower age limit

18 years

# Upper age limit

45 years

#### Sex

Female

#### Key exclusion criteria

- 1. Contraindication to breastfeeding
- 2. Postpartum complication e.g., haemorrhage, chorioamnionitis, retained placenta, any invasive procedures and blood transfusion
- 3. Newborns with complications

#### Date of first enrolment

01/04/2024

#### Date of final enrolment

31/10/2024

# **Locations**

#### Countries of recruitment

Malaysia

# Study participating centre University Of Malaya Medical Centre

Lembah Pantai Kuala Lumpur Malaysia 59100

# Sponsor information

#### Organisation

University Malaya Medical Centre

#### **ROR**

https://ror.org/00vkrxq08

# Funder(s)

#### Funder type

University/education

#### **Funder Name**

Universiti Malaya

#### Alternative Name(s)

University of Malaya, University Malaya, Malayan University, King Edward VII College of Medicine, Raffles College, University of Malaya in Singapore, , , , UM

#### **Funding Body Type**

Government organisation

# Funding Body Subtype

Universities (academic only)

#### Location

Malaysia

# **Results and Publications**

## Individual participant data (IPD) sharing plan

The datasets generated during and/or analysed during the current study are/will be available upon request from Dr Nurul Atiqah Radzali (atiqah.radzali@ummc.edu.my)

# IPD sharing plan summary

Available on request

# **Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	version 1	22/10/2023	19/02/2024	No	Yes
Participant information sheet	version 2.0	24/02/2024	27/02/2024	No	Yes
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes

Protocol file	version 1	22/10/2023	19/02/2024 No	No
Protocol file	version 2.0	24/02/2024	27/02/2024 No	No