

Evaluating abdominal binding using kinesiotape to reduce infection after caesarean section

Submission date 11/03/2022	Recruitment status No longer recruiting	☒ Prospectively registered ☒ Protocol
Registration date 20/03/2022	Overall study status Completed	☐ Statistical analysis plan ☐ Results
Last Edited 08/09/2023	Condition category Pregnancy and Childbirth	☐ Individual participant data ☐ Record updated in last year

Plain English summary of protocol

Background and study aims

A caesarean section, or C-section, is an operation to deliver your baby through a cut made in your tummy and womb.

Caesarean delivery related superficial surgical site infection (SSI) rate ranges from 3% to 15% worldwide. SSI after caesarean section is associated with increased maternal illness, prolonged hospital stay, and increased medical costs. The aim is to evaluate abdominal binding using kinesiotape (elastic therapeutic tape) against micropore tape in reducing post Caesarean section SSI.

Who can participate?

Women scheduled for Caesarean section (planned and unplanned) in UMMC

What does the study involve?

The study involves using either kinesiotape or micropore tape (allocated randomly) as abdominal binding in patients post caesarean section and the wound will be assessed by clinical assessment (using the Center for Disease Control (USA) [CDC] SSI criteria) up to hospital discharge and further by phone-based questionnaire on day 15 and day 30.

What are the possible benefits and risks of participating?

Participants should not expect any benefit as it is not known whether the abdominal binding using kinesiotape is helpful in preventing infection of Caesarean wound. There are no expected serious drawbacks anticipated. If, at any time, participant finds it troublesome, she may request to change or remove the tape.

Where is the study run from?

Department of Obstetrics and Gynecology, UMMC (Malaysia)

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

January 2022 to March 2023

Who is funding the study?

Department of Obstetrics and Gynecology, UMMC (Malaysia)

Who is the main contact?

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Contact information

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

MREC ID 2021113-10747

Study information

Scientific Title

Effectiveness of abdominal binding using kinesiotape on caesarean surgical site infection

Study objectives

Abdominal binding will benefit obese women by lifting up the abdominal skin fold at the incision site and disrupting a favourable environment for SSI.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 07/02/2022, Medical Research Ethics Committee University of Malaya Medical Centre (Lembah Pantai, 59100 Kuala Lumpur, Malaysia; +60379493209/2251; iresearch@ummc.edu.my), ref: 2021113-10747

Study design

Single centre randomized controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Reducing rate of caesarean wound infection

Interventions

Participants will be randomised in the operating theatre during surgery after closure of the skin by opening the lowest number remaining, sealed and opaque envelope. The envelopes will be kept in a known location within obstetric operation theatres in a box with numbered envelopes arranged in sequence. Opened/unsealed envelopes will be discarded. The random allocation

sequence will be generated by an investigator with no clinical involvement in the study. Blinding is not possible due to the nature of the intervention.

After skin closure during Caesarean section and after vaginal cleansing, the abdominal binding will be applied according to randomization (either use Kinesiotape or sham (micropore) tape). Participants are to remove abdominal binding tape and stop using abdominal binding on day 3 after Caesarean delivery.

Intervention Type

Procedure/Surgery

Primary outcome(s)

Cumulative SSI rates to day 30 after Caesarean delivery, by clinical assessment (using the Center for Disease Control (USA) [CDC] SSI criteria) up to hospital discharge and further by phone-based questionnaire on day 15 and day 30.

Key secondary outcome(s)

1. Patient's satisfaction with abdominal binding using visual numerical rating scale (VNRS) at day 15 and day 30
2. Pain score on mobilisation using visual numerical rating scale (VNRS) at day 1, 15 and 30

Completion date

14/03/2023

Eligibility

Key inclusion criteria

1. Age 18 years and above
2. Scheduled for caesarean section (planned or unplanned)
3. BMI ≥ 27.5 kg/m²
4. Access to telephone

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Female

Key exclusion criteria

1. Preexisting abdominal skin infection
2. Hypersensitivity to abdominal kinesiotape or micropore
3. Midline skin incision
4. Category 1 (emergent) Caesarean section

Date of first enrolment

09/04/2022

Date of final enrolment

30/04/2024

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

Hospital/treatment centre

Funder Name

Department of Obstetrics and Gynecology, UMMC

Results and Publications

Individual participant data (IPD) sharing plan

The data-sharing plans for the current study are unknown and will be made available at a later date

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Participant information sheet	version 2	24/01/2022	16/03/2022	No	Yes
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Protocol file	version 1	03/11/2021	16/03/2022	No	No