

Preoperative fasting and postoperative nausea and vomiting after elective caesarean section under spinal anaesthesia: a randomised controlled study

Submission date 30/09/2004	Recruitment status No longer recruiting	☐ Prospectively registered
		☐ Protocol
Registration date 30/09/2004	Overall study status Completed	☐ Statistical analysis plan
		☐ Results
Last Edited 14/11/2014	Condition category Signs and Symptoms	☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

N0084125057

Study information

Scientific Title

Study objectives

To compare the effect of two fasting regimens on the incidence of postoperative nausea and vomiting (PONV) after elective lower segment caesarean sections (LSCS) under spinal anaesthesia.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Signs and Symptoms: Post-operative nausea and vomiting (PONV)

Interventions

Patients undergoing elective caesarean section will be randomly assigned to either of the following two groups:

Group A: Fasted overnight from 12 midnight

Group B: A light breakfast of two slices of toasts and a cup of tea/coffee will be allowed 4 hours preoperatively

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/06/2004

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

08/08/2001

Date of final enrolment

01/06/2004

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Anaesthetics

Hull

United Kingdom

HU3 2JZ

Sponsor information**Organisation**

Department of Health

Funder(s)

Funder type

Government

Funder Name

The North and South Bank Research and Development Consortium (UK)

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary**

Not provided at time of registration