ISRCTN63839707 https://doi.org/10.1186/ISRCTN63839707

Post-Laparoscopic Cholecystectomy Pain: Benefit of Intraperitoneal Saline

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol
Registration date 30/09/2005	**Overall study status** Completed	[_] Statistical analysis plan [_] Results
Last Edited 09/09/2016	**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

Contact name
Mr Daniel Thomas

Contact details
Queen Mary's Sidcup NHS Trust
Sidcup
United Kingdom
DA14 6LT
+44 20 8302 2678
daniel.thomas@qms.nhs.uk

Additional identifiers

Protocol serial number
N0533152392

Study information

Scientific Title
Post-Laparoscopic Cholecystectomy Pain: Benefit of Intraperitoneal Saline

Study objectives

Study aims:
1. To assess a simple way of pain reduction post-operatively
2. To evaluate if this intervention influences the timing of patient discharge and the rate of day case Laparoscopic Cholecystectomy
3. To examine individual differences and identify patient characteristics that may predict likely success and failure from this intervention

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)
Not Specified

Health condition(s) or problem(s) studied
Signs and Symptoms: Post operative pain

Interventions
1. Normal saline instilled intraperitoneally at end of operation
2. Control

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
1. Pain Score
2. Nausea
3. Vomiting
4. Duration of hospitalisation
5. Re-admission within 10 days

Key secondary outcome(s)
Not provided at time of registration

Completion date
30/09/2005

Eligibility

Key inclusion criteria
80 patients at Queen Mary's

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Not Specified

Sex
Not Specified

Key exclusion criteria
1. Any procedure which had to be converted to laparotomy
2. Any patient where the operating surgeon did not adhere to the study protocol
3. Any patient whose cognitive function does not allow them to understand consent to this study

Date of first enrolment
01/10/2004

Date of final enrolment
30/09/2005

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Queen Mary's Sidcup NHS Trust
Sidcup
United Kingdom
DA14 6LT

Sponsor information

Organisation
Department of Health

Funder(s)

Funder type
Government

Funder Name
Queen Mary's Sidcup NHS Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration