# Post-Laparoscopic Cholecystectomy Pain: Benefit of Intraperitoneal Saline

Submission date	Recruitment status	<ul><li>Prospectively registered</li></ul>
30/09/2005	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	Individual participant data
09/09/2016	Signs and Symptoms	[] 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

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# 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

#### Study design

Randomised controlled trial

#### Primary study design

Interventional

#### 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