

# Low rate versus high rate insufflation of the peritoneal cavity for laparoscopic surgery: a randomised comparative study

<b>Submission date</b> 29/09/2006	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
<b>Registration date</b> 29/09/2006	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
<b>Last Edited</b> 24/09/2013	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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**  
N0453168841

## Study information

**Scientific Title**

**Study objectives**

This study aims to find out whether the low rate and high rate of CO<sub>2</sub> insufflation at the commencement of laparoscopic surgery have different effects on cardiac function.

**Ethics approval required**

Old ethics approval format

**Ethics approval(s)**

Not provided at time of registration

**Study design**

Randomised comparative study

**Primary study design**

Interventional

**Study type(s)**

Treatment

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

Surgery: Laparoscopy

**Interventions**

The study involves insufflating the abdomen with gas at either a low rate or high rate and evaluating its impact on the efficacy of the heart pump and on the hormones released during surgery. It will be a randomised clinical study involving 2 groups (CO<sub>2</sub> insufflation at 1 l/1min & CO<sub>2</sub> insufflation of 18 l/min)

**Intervention Type**

Drug

**Phase**

Not Specified

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

CO<sub>2</sub>

**Primary outcome(s)**

Not provided at time of registration

**Key secondary outcome(s)**

Not provided at time of registration

**Completion date**

30/12/2005

**Eligibility****Key inclusion criteria**

Consenting adult patients aged 16-80 years

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Sex**

All

**Key exclusion criteria**

Not provided at time of registration

**Date of first enrolment**

19/07/2005

**Date of final enrolment**

30/12/2005

**Locations****Countries of recruitment**

United Kingdom

England

**Study participating centre**

**MRI Central Manchester & Manchester Children's University Hospitals**

Manchester

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**Sponsor information****Organisation**

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

**Funder(s)**

**Funder type**

Government

**Funder Name**

Central Manchester and Manchester Children's University Hospitals NHS Trust (UK) - Trust Endowment,

**Funder Name**

NHS R&D Support Funding

## **Results and Publications**

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

**IPD sharing plan summary**

Not provided at time of registration