

# Female fertility after laparoscopic pouch surgery

<b>Submission date</b> 13/04/2011	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
<b>Registration date</b> 01/06/2011	<b>Overall study status</b> Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
<b>Last Edited</b> 24/05/2012	<b>Condition category</b> Surgery	<input type="checkbox"/> Individual participant data

**Plain English summary of protocol**  
Not provided at time of registration

## Contact information

**Type(s)**  
Scientific

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## Additional identifiers

## Study information

### Scientific Title

Is postoperative first pregnancy rate higher after laparoscopic restorative proctocolectomy than after open restorative proctocolectomy?

### Study objectives

Adhesions to the fallopian tubes are thought to be the main cause of infertility after ileal pouch-anal anastomosis (IPAA) surgery, a so-called 'tubal factor infertility'. Recent studies demonstrated a significant decrease in adhesion formation after laparoscopic abdominal

surgery. Based on those study results, our hypothesis is that a higher pregnancy rate will be observed after laparoscopic IPAA compared to open IPAA, due to a reduction in adhesions of the Fallopian tubes

### **Ethics approval required**

Old ethics approval format

### **Ethics approval(s)**

The study protocol was approved by the ethics committee of the University Hospitals of the Catholic University in Leuven, Belgium on 30 November 2010, (B32220109939)

In the Netherlands ethics approval is not required for this study design.

### **Study design**

Observational multicentre cross-sectional design

### **Primary study design**

Observational

### **Study type(s)**

Screening

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

Restorative proctocolectomy and pregnancy

### **Interventions**

1. All potential participants will be contacted for consent and thereupon a questionnaire will be sent
2. The questionnaire is developed by the departments of Obstetrics and Gynaecology and the department of Surgery at the Academic Medical Centre and it addresses patients general medical history and their desire for children both before and after IPAA
3. If patients report that there ever was a desire for children, fertility and obstetric history will be inquired after in detail
4. All self-reported patient data will subsequently be verified and completed by checking the available medical records at the hospital of the IPAA
5. Pregnancy is defined as a clinical pregnancy i.e. the presence of a fetus with a heartbeat demonstrated by ultrasound
6. Time to pregnancy is defined as the number of cycles or months between stopping contraceptives until pregnancy occurred, defined as the first day of the last menstrual period
7. Most likely, several patients will report more than one pregnancy, these pregnancies however cannot be considered as independent and therefore only the time to first pregnancy will be used in this study
8. Kaplan-Meier survival curves will be plotted for time to first pregnancy after laparoscopic or open IPAA and will be compared with a log-rank test

### **Intervention Type**

Procedure/Surgery

### **Phase**

Not Applicable

**Primary outcome(s)**

Time (months) to first natural pregnancy after IPAA

**Key secondary outcome(s)**

1. Time (months) to any first pregnancy after IPAA, including pregnancies after fertility treatment
2. Subgroup analyses are planned for UC and FAP patients separately, since previous studies have shown different fertility outcomes after IPAA for these groups of patients

**Completion date**

01/04/2011

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

1. All living females who had IPAA between 1993 - 2009
2. IPAA in one of three tertiary referral centres in Belgium and the Netherlands
3. IPAA could be single- or two-stage and laparoscopic or open
4. At the time of IPAA patients had to be under 41 years of age
5. At the time of the study patients had to be 18 years or older

**Participant type(s)**

Patient

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Sex**

Female

**Key exclusion criteria**

1. Refusal to participate
2. Does not meet inclusion criteria

**Date of first enrolment**

01/02/2010

**Date of final enrolment**

01/04/2011

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

Belgium

Netherlands

**Study participating centre**  
**Academic Medical Centre**  
Amsterdam  
Netherlands  
1105 AZ

## Sponsor information

**Organisation**  
Academic Medical Centre (Netherlands)

**ROR**  
<https://ror.org/03t4gr691>

## Funder(s)

**Funder type**  
University/education

**Funder Name**  
Academic Medical Centre (Netherlands)

## Results and Publications

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

**IPD sharing plan summary**  
Not provided at time of registration

### Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#">Results article</a>	cross sectional study results	01/12/2012		Yes	No