ISRCTN67227760 https://doi.org/10.1186/ISRCTN67227760

# The role of follow-up after gynaecological cancer

| Submission date | Recruitment status | [ ] Prospectively registered |
|---|---|---|
| 29/09/2006 | No longer recruiting | [ ] Protocol |
| **Registration date** | **Overall study status** | [ ] Statistical analysis plan |
| 29/09/2006 | Completed | [ ] Results |
| **Last Edited** | **Condition category** | [ ] Individual participant data |
| 20/08/2012 | Cancer | [ ] 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

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**
N0128168765

## Study information

## Scientific Title

## Study objectives
In patients who have completed treatment for a gynaecological malignancy, this study will compare routine follow-up appointments with a system of patent-initiated follow-up.

## Ethics approval required
Old ethics approval format

## Ethics approval(s)
Prospective pragmatic randomised controlled trial

## Study design
Randomised controlled trial

## Primary study design
Interventional

## Secondary study design
Randomised controlled trial

## Study setting(s)
Not specified

## Study type(s)
Not Specified

## Participant information sheet
Not available in web format, please use the contact details below to request a patient information sheet

## Health condition(s) or problem(s) studied
Cancer: Gynaecological

## Interventions
Rutine follow-up appointments vs system of patent-initiated follow-up

## Intervention Type
Other

## Phase
Not Specified

## Primary outcome measure
Survival

## Secondary outcome measures
Quality of Life, cost effectiveness, rates of surgical and medical intervention in primary and secondary care and patient acceptability

**Overall study start date**
01/01/2004

**Completion date**
31/12/2009

# Eligibility

**Key inclusion criteria**
All women who have completed treatment and whose symptoms are controlled.

**Participant type(s)**
Patient

**Age group**
Adult

**Sex**
Female

**Target number of participants**
200

**Key exclusion criteria**
Patients who still require treatment of some kind.

**Date of first enrolment**
01/01/2004

**Date of final enrolment**
31/12/2009

# Locations

**Countries of recruitment**
England

United Kingdom

**Study participating centre**
**Gynaecology**
Liverpool
United Kingdom
L8 7SS

# Sponsor information

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

### Sponsor details
The Department of Health, Richmond House, 79 Whitehall
London
United Kingdom
SW1A 2NL
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dhmail@doh.gsi.org.uk

### Sponsor type
Government

### Website
http://www.dh.gov.uk/Home/fs/en

## Funder(s)

### Funder type
Government

### Funder Name
Liverpool Women's Hospital NHS Trust (UK), NHS R&D Support Funding

## Results and Publications

### Publication and dissemination plan
Not provided at time of registration

### Intention to publish date

### Individual participant data (IPD) sharing plan

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