A randomised study comparing satisfaction with follow up led by a trained cancer nurse versus conventional medical follow-up after primary treatment for ovarian cancer

Submission date	Recruitment status No longer recruiting	Prospectively registered	
29/09/2006		☐ Protocol	
Registration date	Overall study status Completed	Statistical analysis plan	
29/09/2006		[X] Results	
Last Edited	Condition category	[] Individual participant data	
11/10/2017	Cancer		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Ms A Lanceley

Contact details

Gynaecological Oncology
Gynaecological Cancer Research Centre
Institute for Women's Health
University College London, Maple House, 149 Tottenham Court Road
London
United Kingdom
W1P 9LL
+44 (0)20 7380 6906
a.lanceley@ucl.ac.uk

Additional identifiers

Protocol serial number

N0263172828

Study information

Scientific Title

A randomised study comparing satisfaction with follow up led by a trained cancer nurse versus conventional medical follow-up after primary treatment for ovarian cancer

Study objectives

To elicit and compare women's satisfaction with nurse-led follow up after treatment for ovarian cancer with routine follow up, in order to generate information, which enhances the ability of service providers to meet their needs.

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)

Other

Health condition(s) or problem(s) studied

Cancer: Ovarian

Interventions

- 1. Conventional medical follow up
- 2. Follow up care led by a trained cancer nurse

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Patient satisfaction and quality of life (QOL). QOL will be measured using the European Organisation for research and treatment of cancer (EORTC) core QOL questionnaire and supplementary ovarian cancer module

Key secondary outcome(s))

Not provided at time of registration

Completion date

30/04/2009

Eligibility

Key inclusion criteria

100 patients from Gynaecological Oncology

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Female

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/11/2005

Date of final enrolment

30/04/2009

Locations

Countries of recruitment

United Kingdom

England

Study participating centre University College London

London United Kingdom W1P 9LL

Sponsor information

Organisation

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

Funder(s)

Funder type

Government

Funder Name

University College London Hospitals NHS Foundation Trust

Alternative Name(s)

University College London Hospitals, UCLH

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

United Kingdom

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

Study outputs

Output type	Details	Date created Date added	Peer reviewed?	Patient-facing?
Results article	results	01/01/2017	Yes	No
Participant information sheet	Participant information sheet	11/11/2025 11/11/2025	No	Yes