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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
29/09/2006

Recruitment status
No longer recruiting

Registration date
29/09/2006

Overall study status
Completed

Last Edited
11/10/2017

Condition category
Cancer

[_] Prospectively registered
[_] Protocol
[_] Statistical analysis plan
[X] Results
[_] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Randomised controlled trial

Study setting(s)
Other

Study type(s)
Other

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

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 measure
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

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/11/2005

Completion date
30/04/2009

Eligibility

Key inclusion criteria
100 patients from Gynaecological Oncology

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
100

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
England

United Kingdom

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

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

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

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

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

Study outputs

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