

A randomised trial of telecolposcopy screening in the management of women with minor cervical smear abnormalities

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 16/06/2014	Condition category Urological and Genital Diseases	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

Contact name

Mr Ian Etherington

Contact details

Department of Gynaecology
City Hospital
Dudley Road
Birmingham
United Kingdom
B18 7QH
+44 (0)121 507 4377

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0064069074

Study information

Scientific Title

Study objectives

The primary objective of the study is to evaluate the impact of telecolposcopy screening on the number of visits women have to the colposcopy clinic before discharge to community cytology.

Secondary objectives are:

1. To evaluate the anxiety levels experienced by women in the study and control groups.
2. To evaluate the cost impact to the Screening Programme of introducing telecolposcopy
3. To evaluate the impact of such screening on the time interval between the first abnormal smear and the detection of high-grade cervical intraepithelial neoplasia (CIN)
4. To evaluate the ability of telecolposcopy screening to differentiate between grades of CIN

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)

Screening

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

Urological and Genital Diseases: Telecolposcopy

Interventions

Impact of telecolposcopy screening

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/08/2000

Completion date

01/04/2003

Eligibility

Key inclusion criteria

Recruiting 200 women in each arm

Participant type(s)

Patient

Age group

Not Specified

Sex

Female

Target number of participants

400

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/08/2000

Date of final enrolment

01/04/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Department of Gynaecology
Birmingham
United Kingdom
B18 7QH

Sponsor information

Organisation
Department of Health (UK)

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
<http://www.doh.gov.uk>

Funder(s)

Funder type
Government

Funder Name
Sandwell and West Birmingham Hospitals NHS Trust - City Hospital (UK)

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