

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

Protocol serial number
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

Study type(s)

Screening

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(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/04/2003

Eligibility

Key inclusion criteria

Recruiting 200 women in each arm

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Female

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

United Kingdom

England

Study participating centre

Department of Gynaecology

Birmingham

United Kingdom

B18 7QH

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type
Government

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

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary
Not provided at time of registration