ISRCTN50023379 https://doi.org/10.1186/ISRCTN50023379

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

Registration date
12/09/2003

Overall study status
Completed

Last Edited
16/06/2014

Condition category
Urological and Genital Diseases

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

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

Study outputs

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