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How can we reduce the frequency of unscheduled smear rates?

Submission date	Recruitment status	[ ] Prospectively registered
23/01/2004	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
23/01/2004	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
26/11/2015	Cancer	[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
RHC28303

Study information

Scientific Title
How can we reduce the frequency of unscheduled smear rates?

Study objectives
To improve the procedural effectiveness and efficiency of the NHS Cervical Screening Programme

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
Cervical neoplasia, cervical smear

Interventions
Not provided at time of registration

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
For each intervention, the rate of unscheduled smears will be compared between the intervention and control arms.

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/09/1999

Completion date
01/12/2002

Eligibility

Key inclusion criteria
Women aged 25-64 who fulfill study criteria. Eligible women identified from the Cytology database held by the Lancashire and South Cumbria Agency.

Participant type(s)
Patient

Age group
Adult

Sex
Female

Target number of participants
Not provided at time of registration

Key exclusion criteria
Does not match inclusion criteria

Date of first enrolment
01/09/1999

Date of final enrolment
01/12/2002

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
University of Manchester
Manchester
United Kingdom
M20 9QL

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

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.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive North West (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