

A randomised controlled trial to evaluate a novel CD-ROM learning package against the standard lecture - training system for research governance in the teaching of new researchers in an NHS Trust

Submission date	Recruitment status	<input type="checkbox"/> Prospectively registered
28/09/2007	No longer recruiting	<input type="checkbox"/> Protocol
Registration date	Overall study status	<input type="checkbox"/> Statistical analysis plan
28/09/2007	Completed	<input type="checkbox"/> Results
Last Edited	Condition category	<input type="checkbox"/> Individual participant data
12/10/2017	Other	<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

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Contact details

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The Cardiothoracic Centre
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Additional identifiers

Protocol serial number

N0054184076

Study information

Scientific Title

A randomised controlled trial to evaluate a novel CD-ROM learning package against the standard lecture - training system for research governance in the teaching of new researchers in an NHS Trust

Study objectives

Is the use of a CD-ROM teaching package more cost effective than traditional lecture methods of teaching Research Governance to new researchers?

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)

Treatment

Health condition(s) or problem(s) studied

Not Applicable: Education

Interventions

The purpose of this study is to determine whether a CD-ROM can provide education in research governance to new researchers as effectively but more cheaply and conveniently than routine lecture teaching. A CD-ROM teaching package has been produced that is identical to a Power point teaching presentation. Staff at the Cardiothoracic Centre - Liverpool NHS Trust who have research as part of their job description will be invited to attend a half day study day in the lecture theatre of the hospital and provided with an information sheet to decide if they wish to take part in the study or not.

On arrival in the lecture theatre subjects will be asked to consent to the study. Those who do will be randomised by pre numbered envelopes prepared by the clinical governance department.

1. Those subjects randomised to lecture will remain in the lecture theatre and receive 2 hours of the power point presentation with hand outs and web site addresses so that they have identical information to the CD-ROM group.

2. Those randomised to CD-ROM will be provided with a disc containing the PowerPoint presentation, web links and copies of important documents. They will be able to use this at home or with assistance from the librarian in the library.

All participants will be requested not to discuss source documents with other members of the research group. They will also be requested to return in 2 weeks time for a knowledge multichoice questionnaire and a satisfaction survey.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

1. Cost effectiveness will be measured by cost per hour per attendee taking into account only direct costs, with the assumption that the CD-ROM will be used for the same length of time as the lecture. Subjects who do not attend the exam will be considered exam failures and their data will be included in the cost effectiveness. Subjects who attend the exam but obtain less than the 50% pass mark will also be considered exam failures.
2. Satisfaction with the teaching method will be analysed via Likert scale questions from Price J 2005 and written responses to open ended questions.

Key secondary outcome(s)

Not provided at time of registration

Completion date

01/06/2006

Eligibility

Key inclusion criteria

Participants will be identified by obtaining lists of staff on Banding 6 & 7 with research as part of their job description. These staff will then be approached by letter also enclosing a study information sheet. Participants will be recruited and consented on attendance at the lecture theatre for the teaching session.

Subject inclusion criteria:

1. Age > 18 years
2. Able to give informed consent
3. Research active
4. Employee of the Cardiothoracic Centre - Liverpool NHS Trust

The sample size has been calculated to detect a £2.50 difference in cost using a 2 sample t test . at 90% power with a mean population wage of £ 20.46 and a standard deviation of 1.8. This will be calculated by taking the direct costs of subject wages per hour spent learning and the unit cost of the teaching method. This provides a sample size of 24 (12 in each arm of the study). To allow for drop outs and to obtain 24 evaluable questionnaires a total of 30 subjects will be enrolled.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

Not Specified

Key exclusion criteria

1. Unable to take ½ day for teaching and follow-up
2. No access to computer
3. Involved in another study within the last 30 days

Date of first enrolment

01/04/2006

Date of final enrolment

01/06/2006

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Clinical Trials Unit

Liverpool

United Kingdom

L14 3PE

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2007 Update - Department of Health

Funder(s)

Funder type

Government

Funder Name

The Cardiothoracic Centre Liverpool NHS Trust (UK), NHS R&D Support Funding

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