A randomised trial comparing the effect of selfdirected online learning using a web-based resource with a classroom-based interactive workshop, on the knowledge and search skills of healthcare professionals

Submission date	Recruitment status	Prospectively registered
29/09/2006	No longer recruiting	Protocol
Registration date	Overall study status	Statistical analysis plan
29/09/2006	Completed	Results
Last Edited	Condition category	[] Individual participant data
12/05/2017	Other	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 N0643164123

Study information

Scientific Title

A randomised trial comparing the effect of self-directed online learning using a web-based resource with a classroom-based interactive workshop, on the knowledge and search skills of healthcare professionals

Study objectives

Is there a difference in terms of knowledge and skills, between self-directed learning using webbased resource directed by students, compared with a classroom based interactive workshop directed by tutors?

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)

Other

Health condition(s) or problem(s) studied

Knowledge and skill education

Interventions

At each location participants will be randomised into two groups:

- 1. Group one: access to a search-skills web resource (e-group)
- 2. Group two: receive a search workshop taught by a librarian (workshop group)

The e-learning resource and the workshop will have the same objectives and cover the same content. Each intervention group will attend one group session - the e-group will be shown a demonstration of the e-learning resource and asked to complete as much of the module as they can over the next week; the workshop group will receive a three-hour taught workshop with handouts and slides. At the beginning of these sessions, each participant will complete a pre-intervention search test (baseline) using a clinical scenario. One week after these initial sessions, a post-intervention search test should be completed by both groups. Pre- (baseline) and post-intervention search results will be compared. Follow-up tests will be sent at 4 weeks, 8 weeks (and 12 weeks if time allows).

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

30/04/2005

Eligibility

Key inclusion criteria

Health professionals

Participant type(s)

Health professional

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/03/2005

Date of final enrolment

30/04/2005

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Williams Room

Oxford United Kingdom OX3 7LF

Sponsor information

Organisation

Record Provided by the NHSTCT Register - 2006 Update - Department of Health

Funder(s)

Funder type

Government

Funder Name

Thames Valley Primary Care Research Partnership (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 No Yes