

Pilot trial to assess the feasibility and response rates for an RCT evaluating the effectiveness of a computer tailored intervention for smoking cessation in general practice

Submission date 30/09/2004	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 30/09/2004	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 01/04/2008	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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
N0530137082

Study information

Scientific Title

Study objectives

The aim of the proposed research is to carry out a pilot study to assess the feasibility of, and response rates to, an intervention for smoking cessation. We have already developed a computer-based system for generating individually tailored feedback reports designed to encourage and help smokers to quit. These self-help materials tailored for each individual mimic the care used in clinical settings, but make this available to the general public. We plan to adapt this system for use in general practice, modifying the questionnaire and the feedback reports to different educational levels. The pilot work will lead to a large RCT evaluating the effectiveness of the intervention. The hypothesis to be tested on smokers identified from general practitioners lists is that: Personalised feedback reports tailored to levels of reading ability and sent to smokers in varying stages of readiness to quit, will increase quitting activity and modify the attitudes of the smokers over and above that found with usual standard care received from the practice.

Please note that this trial was completed on the 30th June 2005. The previous end date for this pilot study was 31/05/2009. The main RCT of this pilot study has been registered under ISRCTN05385712 (see <http://www.controlled-trials.com/ISRCTN05385712>).

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial - pilot study

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Smoking addiction

Interventions

Questionnaires; Before-After-Study, Randomised Controlled Trial (RCT) comparison between:

1. Normal treatment
2. Normal treatment plus tailored feedback reports group

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Computer tailored feedback, adapted to reading levels and readiness to quit, is a simple intervention which could be widely replicated and delivered cost effectively to a large proportion of the smoking population. A modest success rate could have a large effect on public health given its recruitment potential and make a valuable contribution to lowering smoking prevalence. The method mimics the guidelines of asking, advising, assessing, assisting and arranging and could be incorporated into GPs standard treatment at less time and cost. These materials, tailored to the requirements of each individual, would offer GPs and practice nurses an efficient way of integrating smoking cessation counselling into a busy primary care practice. The computer programme could also be made available on CD Rom to produce immediate printed feedback for use by primary care professionals.

Key secondary outcome(s)

Not provided at time of registration

Completion date

30/06/2005

Eligibility

Key inclusion criteria

1. Smokers
2. Aged between 18 and 65 years, either sex
3. Identified from records in four practices

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

65 years

Sex

All

Key exclusion criteria

Does not comply with above inclusion criteria

Date of first enrolment

01/01/2004

Date of final enrolment

30/06/2005

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

University College London

London

United Kingdom

WC1E 6BT

Sponsor information

Organisation

Department of Health

Funder(s)

Funder type

Government

Funder Name

North Central London Research Consortium (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?
Results article	results	01/09/2007		Yes	No