

A randomised controlled trial of automated electronic feedback to reduce the cardiovascular risk of individuals in general practice - the e-Nudge trial

Submission date 06/02/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 15/03/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 05/12/2017	Condition category Circulatory System	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Prof Margaret Thorogood

Contact details
Warwick Medical School
Gibbet Hill Road
Coventry
United Kingdom
CV4 7AL
+44 (0)2476 574509
margaret.thorogood@warwick.ac.uk

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

A randomised controlled trial of automated electronic feedback to reduce the cardiovascular risk of individuals in general practice - the e-Nudge trial

Study objectives

Primary research question: can an automated system of electronic feedback (e-Nudge) reduce the incidence of cardiovascular events in high-risk patients in general practice, compared to usual care?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved by the Warwickshire Local Research Ethics Committee, on 06/09/2005, reference number: 05/Q2803/85

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

GP practice

Study type(s)

Prevention

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular disease and diabetes

Interventions

A system of automatically generated alerts and electronic reminders versus usual care

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Cardiovascular event rates

Secondary outcome measures

Proportion of individuals at high cardiovascular risk

Overall study start date

01/03/2006

Completion date

28/02/2009

Eligibility

Key inclusion criteria

General practices in Warwickshire and Coventry using appropriate clinical software

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Up to 26 practices

Key exclusion criteria

Practices at which the principal investigator undertakes clinical practice

Date of first enrolment

01/03/2006

Date of final enrolment

28/02/2009

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Warwick Medical School

Coventry

United Kingdom
CV4 7AL

Sponsor information

Organisation

University of Warwick

Sponsor details

Gibbet Hill Road
Coventry
England
United Kingdom
CV4 7AL

Sponsor type

University/education

ROR

<https://ror.org/01a77tt86>

Funder(s)

Funder type

University/education

Funder Name

Department of Health PhD Studentship from Warwick Medical School

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Protocol article	protocol	28/04/2006		Yes	No
Results article	results	01/04/2010		Yes	No