

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

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

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### Contact details

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## 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?
<a href="#">Protocol article</a>	protocol	28/04/2006		Yes	No
<a href="#">Results article</a>	results	01/04/2010		Yes	No