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Methods of implementing guidelines for hypertension in primary care: acceptability, effectiveness and cost-effectiveness

Submission date 23/01/2004	**Recruitment status** No longer recruiting	[_] Prospectively registered [_] Protocol [_] Statistical analysis plan [_] Results [_] Individual participant data [_] Record updated in last year
Registration date 23/01/2004	**Overall study status** Completed	
Last Edited 24/10/2019	**Condition category** Circulatory System	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
C/COM/04/16-04-94/O'BRIEN/F

Study information

Scientific Title
Methods of implementing guidelines for hypertension in primary care: acceptability, effectiveness and cost-effectiveness

Study objectives
This project examines the acceptability, effectiveness and cost-effectiveness of different methods of implementing clinical guidelines for hypertension in primary care, by means of a randomised controlled trial of financial incentive and /or behaviour modification. Both quantitative and qualitative methods will be used to explore fully issues surrounding the management of hypertension and the implementation of guidelines in primary care.

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

Secondary study design
Randomised controlled trial

Study setting(s)
GP practice

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Cardiovascular diseases: Hypertension

Interventions
Not provided at time of registration

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/08/1994

Completion date
01/12/1996

Eligibility

Key inclusion criteria
Not provided at time of registration

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/08/1994

Date of final enrolment
01/12/1996

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
South and West Devon HA
Dartington
United Kingdom
TQ9 6JE

Sponsor information

Organisation
NHS R&D Regional Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL
+44 (0)20 7307 2622
dhmail@doh.gsi.org.uk

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Executive South West (UK)

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