High blood pressure control and lipid lowering in patients at high cardiovascular risk: a cluster randomised factorial comparison of educational and protocol-based interventions to help primary care teams improve control of patients with poor risk factor control

Submission date	Recruitment status	 Prospectively registered
18/06/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
18/06/2010	Completed	Results
Last Edited	Condition category	[] Individual participant data
06/04/2017	Circulatory System	Record updated in last year

Plain English summary of protocolNot provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Mr Bob Scott

Contact details

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Additional identifiers

Study information

Scientific Title

High blood pressure control and lipid lowering in patients at high cardiovascular risk: a cluster randomised factorial comparison of educational and protocol-based interventions to help primary care teams improve control of patients with poor risk factor control

Acronym

HiLo

Study objectives

This is a 2 x 2 factorial design complex health services intervention research trial which examines the influence of none, either or both of two interventions designed to help primary care clinicians improve the blood pressure and cholesterol control of patients at high risk of cardiovascular events over 12 months. One intervention explores expertise sharing whilst the other explores protocol adherence by clinicians.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Single-centre randomised interventional prevention, process of care and treatment trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Cardiovascular, Diabetes Research Network, Primary Care Research Network for England; Subtopic: Both, Not Assigned, Cardiovascular (all Subtopics); Disease: Diabetic Control, Cardiovascular, All Diseases

Interventions

- 1. Case management
- 2. Protocol-based treatment suggestions
- 3. Clinical educational support
- 4. Expertise sharing

Follow-up length: 12 months

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Change in blood pressure (BP) treatment at 1 year

Key secondary outcome(s))

- 1. Change in lipid treatment at 1 year
- 2. Change in systolic blood pressure (BP) at 1 year
- 3. Change in total cholesterol at 1 year

Completion date

01/08/2011

Eligibility

Key inclusion criteria

- 1. Elevated systolic blood pressure greater than or equal to 160 mmHg within the last 12 months
- 2. Elevated total cholesterol greater than or equal to 5 mmol/l
- 3. Patients in secondary prevention, or risk equivalent, or hypertensives in primary prevention at more than 20% 10-year cardiovascular disease (CVD) risk by QRISK2 or Framingham equations
- 4. Male and female, lower age limit of 40 years

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

Compelling reasons to avoid increase in blood pressure or lipid-lowering medication, such as severe debilitating illness or terminal cancer

Date of first enrolment

01/06/2006

Date of final enrolment

01/08/2011

Locations

Countries of recruitment

United Kingdom

Study participating centre
Barts and The London NHS Trust
London
United Kingdom
EC1M 6BQ

Sponsor information

Organisation

William Harvey Research Institute (UK)

ROR

https://ror.org/0574dzy90

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

City and Hackney Teaching Primary Care Trust (UK)

Funder Name

Merck Schering-Plough (UK)

Funder Name

Pfizer (UK)

Alternative Name(s)

Pfizer Inc., Pfizer Consumer Healthcare, Davis, Charles Pfizer & Company, Warner-Lambert, King Pharmaceuticals, Wyeth Pharmaceuticals, Seagen, Pfizer Inc

Funding Body Type

Government organisation

Funding Body Subtype

For-profit companies (industry)

Location

United States of America

Funder Name

Tower Hamlets Primary Care Trust (UK)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Study outputs

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet
Participant information sheet
11/11/2025 No Yes