

Preventing hospitalisation and early death from chronic heart failure in primary care: a randomised controlled trial of pharmacist led medication review

Submission date 15/06/2006	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 06/07/2006	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 10/04/2012	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

04/SO701/8

Study information

Scientific Title

Study objectives

Pharmacists in primary care can prevent hospitalisation and early death through medication review

Ethics approval required

Old ethics approval format

Ethics approval(s)

2004; Greater Glasgow Primary Care NHS Trust Local Research Ethics Committee

Study design

Randomised controlled trial with cost effectiveness analysis

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Prevention

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Chronic heart failure

Interventions

Practices in the control group, in common with intervention group, receive a complete disease register for patients with a diagnosis of chronic heart failure due to left ventricular systolic dysfunction. Control group practices do not receive any further pharmacist led support. Intervention practices receive pharmacist medication review for their patients with heart failure.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Death or heart failure hospitalisations

Secondary outcome measures

Survival times until occurrence of first heart failure hospitalisation or death

Overall study start date

01/05/2004

Completion date

01/06/2009

Eligibility**Key inclusion criteria**

1. Consenting practices
2. Patients with confirmed left ventricular systolic dysfunction (LSVD) providing written consent

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

2400

Key exclusion criteria

1. Non LVSD
2. Not chronic heart failure
3. Renal failure requiring dialysis
4. Patient under enhanced form of Primary Care

Date of first enrolment

01/05/2004

Date of final enrolment

01/06/2009

Locations**Countries of recruitment**

United Kingdom

Study participating centre
Primary Care Research and Development Directorate
Scotland
United Kingdom
G12 OXH

Sponsor information

Organisation

Greater Glasgow Primary Care NHS Trust (UK)

Sponsor details

Primary Care Research and Development Directorate
Gartnavel Royal Hospital
1055 Great Western Road
Glasgow
Scotland
Scotland
United Kingdom
G12 OXH

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/05kdz4d87>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

NHS Glasgow Research and Development

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?
Results article	1. results	01/02/2012		Yes	No