

Clinical medication review by a pharmacist of patients on repeat prescriptions in general practice

Submission date 25/04/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 25/04/2003	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 27/08/2009	Condition category Other	<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

HTA 93/22/09

Study information

Scientific Title

Study objectives

This randomised controlled clinical trial will examine whether a suitably experienced pharmacist can conduct effective clinical medication review of patients on repeat prescriptions. The study will compare this approach with a control group who will be part of the normal review mechanisms for repeat prescriptions. The pharmacist will be an experienced clinical pharmacist with the appropriate clinical and consultation skills.

The objective is to assess whether clinical medication review by a pharmacist is a cost-effective method for improving the extent and quality of clinical control of repeat prescribing, compared with that achieved by the practices' normal procedures.

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)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Repeat prescriptions

Interventions

A total of 1200 patients from four practices will be randomised to the active or control group. Patients in the active group will be called for an appointment with the pharmacist at the surgery. The pharmacist will assess the continuing appropriateness of the medication regime and make appropriate recommendations (in conjunction with the GP). The control group will receive standard care.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The primary outcome was the number of repeat medication changes per patient over a 12-month period. The secondary outcome was the effect on the medication costs. The intervention group was compared with the control group to see whether a review had taken place, the numbers of medication changes, the numbers of repeat medications and the numbers of dosage times. The effects of the medication review clinics were considered in relation to practice consultations, outpatient consultations, hospital admissions and deaths from any cause. The number and nature of the pharmacist's interventions and recommendations were recorded, together with whether the recommendations were accepted by the GP.

Secondary outcome measures

Not provided at time of registration.

Overall study start date

01/03/1999

Completion date

31/03/2001

Eligibility**Key inclusion criteria**

Not provided at time of registration.

Participant type(s)

Patient

Age group

Not Specified

Sex

Both

Target number of participants

1200

Key exclusion criteria

Not provided at time of registration.

Date of first enrolment

01/03/1999

Date of final enrolment

31/03/2001

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

School of Healthcare Studies

Leeds

United Kingdom

LS2 9UT

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

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Sponsor type

Government

Website

<http://www.dh.gov.uk/en/index.htm>

ROR

<https://ror.org/03sbpja79>

Funder(s)

Funder type

Government

Funder Name

NIHR Health Technology Assessment Programme - HTA (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	08/12/2001		Yes	No