Can a review of the medication of elderly nursing and residential home patients improve the quality of prescribing and residents' outcomes?

Recruitment status No longer recruiting	Prospectively registered	
	☐ Protocol	
Overall study status	Statistical analysis plan	
Completed	[X] Results	
Condition category	[] Individual participant data	
	No longer recruiting Overall study status Completed	

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

N0436118166

Study information

Scientific Title

Study objectives

- 1. Improve quality of therapeutics care for elderly residents of homes
- 2. Reduce risk of adverse drug reactions and interactions
- 3. Improve economy of prescribing
- 4. Mental and physical well-being as measured by mini-mental state and Barthel scores respectively

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Medication review of elderly patients

Interventions

Random allocation to:

- 1. Standard treatment
- 2. Medication review

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Compare the number of changes made to repeat medication schedules between baseline and end data collection
- 2. The cost of repeat medications and the number per patient will be compared in this order of priority

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/01/2002

Completion date

01/01/2004

Eligibility

Key inclusion criteria

Patients aged 65 or over and on at least one repeat medication at the time of identification and in a nursing or residential home.

Participant type(s)

Patient

Age group

Senior

Sex

Both

Target number of participants

Not provided at time of registration

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/01/2002

Date of final enrolment

01/01/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Division of Academic Pharmacy Practice
Leeds
United Kingdom
LS2 9LN

Sponsor information

Organisation

Department of Health (UK)

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL

Sponsor type

Government

Website

http://www.doh.gov.uk

Funder(s)

Funder type

Government

Funder Name

Leeds Teaching Hospitals NHS Foundation Trust (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 summaryNot provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	Results	01/11/2006		Yes	No