# Consequences of chronic kidney disease (CKD) in older people

Submission date	Recruitment status	Prospectively registered
19/05/2010	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
19/05/2010	Completed	Results
Last Edited	Condition category	Individual participant data
13/12/2019	Urological and Genital Diseases	Record updated in last year

# Plain English summary of protocol

Not provided at time of registration

# Contact information

# Type(s)

Scientific

#### Contact name

Dr Stephen John

#### Contact details

Royal Derby Hospital Uttoxeter Road Derby United Kingdom DE22 3NE

# Additional identifiers

**EudraCT/CTIS** number

**IRAS** number

ClinicalTrials.gov number

Secondary identifying numbers 6258

# Study information

Scientific Title

Cardiovascular and functional consequences of chronic kidney disease in older people: a single centre non-randomised interventional treatment trial

#### **Study objectives**

The aim of this study is to investigate this area with an integrated range of non-invasive assessments, both without antihypertensive medication, and with BP control to published best-practice guidelines.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Trent MREC, 15/03/2007, ref: 06/MRE04/73

#### Study design

Single centre non-randomised interventional treatment trial

#### Primary study design

Interventional

#### Secondary study design

Non randomised controlled trial

## Study setting(s)

GP practice

# Study type(s)

Treatment

#### 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

Topic: Renal and Urogenital; Subtopic: Renal and Urogenital (all Subtopics); Disease: Renal

#### **Interventions**

Antihypertensive therapy using standard agents to target BP 130/80 mmHg. Assessments are performed after antihypertensive washout, reintroduction and 12 months follow up.

#### Intervention Type

Other

#### Phase

Phase IV

#### Primary outcome measure

Changes in Baroreflex sensitivity (a composite marker of autonomic function) in response to antihypertensive therapy, assessed at each assessment

#### Secondary outcome measures

Assessed at each assessment:

- 1. Cardiovascular predictors of Falls propensity
- 2. Cardiovascular consequences of antihypertensive therapy

## Overall study start date

01/08/2007

#### Completion date

08/02/2010

# **Eligibility**

#### Key inclusion criteria

- 1. Aged greater than 70 years
- 2. CKD 3, 4 or hypertensive (greater than 130/80) with no renal disease

#### Participant type(s)

Patient

#### Age group

Adult

#### Sex

Both

# Target number of participants

Planned sample size: 80

#### Key exclusion criteria

- 1. Use of more than 3 drugs for blood pressure control
- 2. Likely to develop CKD 5 within one year
- 3. Poor mobility precluding completion of assessment
- 4. Diabetes
- 5. Abbreviated mental test (AMT) score < 8
- 6. Recent acute illness (within 3 months)
- 7. Ischaemic heart disease requiring beta-blockade (nitrates/nicorandil are permitted)
- 8. Involvement in another clinical trial within 3 months
- 9. Attending day hospital or attending falls reduction services
- 10. Residents in nursing or residential homes
- 11. Renovascular disease precluding ACE-inhibitor (ACEi) or Angiotensin Receptor Blocker (ARB) usage
- 12. BP prior to antihypertensive withdrawal > 160/90mmHg
- 13. Malignant hypertension
- 14. Severe peripheral vascular disease or significant valvular heart disease
- 15. Heart failure (NYHA III/IV) or other cause to prevent diuretic withdrawal
- 16. Atrial fibrillation or other significant arrhythmia (precludes pulse wave velocity (PWV) measurement)

17. Currently taking antihypertensive medication for which MRHA approval (CTA) has not been granted

18. Active Obstructive uropathy

#### Date of first enrolment

01/08/2007

#### Date of final enrolment

08/02/2010

# Locations

# Countries of recruitment

England

**United Kingdom** 

# Study participating centre Royal Derby Hospital

Derby United Kingdom DE22 3NE

# Sponsor information

## Organisation

Derby Hospital NHS Foundation Trust (UK)

#### Sponsor details

Royal Derby Hospital
Uttoxeter Road
Derby
England
United Kingdom
DE22 3NE
+44 1332 340131
t.grieve@derbyhospitals.nhs.uk

#### Sponsor type

Hospital/treatment centre

#### Website

http://www.derbyhospitals.nhs.uk/

# Funder(s)

#### Funder type

Research organisation

#### Funder Name

British Renal Society (UK)

## Alternative Name(s)

**BRS** 

# **Funding Body Type**

Private sector organisation

## **Funding Body Subtype**

Associations and societies (private and public)

#### Location

**United Kingdom** 

# **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