

The investigation of functional impact of acute renal colic with assessment of relative renal function by multi-slice contrast enhanced computed tomography

Submission date 30/09/2004	Recruitment status No longer recruiting	☐ Prospectively registered
Registration date 30/09/2004	Overall study status Completed	☐ Protocol
Last Edited 18/10/2017	Condition category Urological and Genital Diseases	☐ Statistical analysis plan
		☐ Results
		☐ Individual participant data
		☐ Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

N0544139650

Study information

Scientific Title

The investigation of functional impact of acute renal colic with assessment of relative renal function by multi-slice contrast enhanced computed tomography

Study objectives

The aims of renal stone management are twofold: symptom control and maintenance of renal function. The trial is designed to evaluate the effects of management on the second of these two aims.

Does knowledge of relative renal function in acute renal colic patients improve outcome and can multi slice contrast enhanced computer tomography supply this information allowing a 'one stop' assessment?

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Primary study design

Interventional

Study design

Randomised controlled trial

Study type(s)

Diagnostic

Health condition(s) or problem(s) studied

Urological and Genital Diseases: Renal function

Interventions

Comparative imaging study with a randomised controlled arm. The interventions being compared are:

1. The availability of a functional image (not information that urologists currently routinely request or use in the management of renal colic patients)
2. No available functional image, as per current practice.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Not provided at time of registration

Key secondary outcome(s)

Not provided at time of registration

Completion date

26/01/2007

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

27/01/2004

Date of final enrolment

26/01/2007

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Addenbrooke's NHS Trust

Cambridge

United Kingdom

CB2 2QQ

Sponsor information

Organisation

Department of Health

Funder(s)**Funder type**

Government

Funder Name

Cambridge Consortium - Addenbrooke's (UK)

Results and Publications**Individual participant data (IPD) sharing plan****IPD sharing plan summary**

Not provided at time of registration