

64 slice computed tomography angiography (CTA) and rotational digital subtraction angiography (DSA) in the assessment of patients with subarachnoid haemorrhage: a study to improve effectiveness and reduce patient radiation dose

Submission date 26/05/2005	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered <input type="checkbox"/> Protocol
Registration date 13/07/2005	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input checked="" type="checkbox"/> Results
Last Edited 30/07/2012	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

05/Q0108/194

Study information

Scientific Title

Study objectives

64 slice CTA is just as good as DSA for the detection of intra cerebral aneurysms and is superior to 16 slice CTA in resolution and dose to the patient.

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)

Hospital

Study type(s)

Diagnostic

Participant information sheet

Health condition(s) or problem(s) studied

Subarachnoid haemorrhage

Interventions

64 slice computed tomography angiography (CTA) versus 16 slice CTA versus rotational digital subtraction angiography (DSA)

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Ability to detect intracranial aneurysms

Secondary outcome measures

Radiation dose administered

Overall study start date

01/10/2005

Completion date

31/12/2006

Eligibility

Key inclusion criteria

All patients admitted with sub-arachnoid haemorrhage to the Neurosurgery department.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

100

Key exclusion criteria

Does not match inclusion criteria

Date of first enrolment

01/10/2005

Date of final enrolment

31/12/2006

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University Department of Radiology
Cambridge

United Kingdom
CB2 2QQ

Sponsor information

Organisation

Addenbrookes NHS Trust (UK)

Sponsor details

Addenbrooke's Hospital
Hills Road
Cambridge
England
United Kingdom
CB2 2QQ

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/055vbx86>

Funder(s)

Funder type

University/education

Funder Name

University of Cambridge (UK) - Internal University departmental funding

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	01/05/2005		Yes	No