

Diffusion and perfusion weighted magnetic resonance imaging (MRI) following acute aneurysmal subarachnoid haemorrhage; early detection of vasospasm-induced ischaemia

Submission date 12/09/2003	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 12/09/2003	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 22/05/2017	Condition category Circulatory System	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N0016111404

Study information

Scientific Title

Diffusion and perfusion weighted magnetic resonance imaging (MRI) following acute aneurysmal subarachnoid haemorrhage; early detection of vasospasm-induced ischaemia

Study objectives

To assess the utility of diffusion and perfusion MRI in predicting and assessing ischaemia related to vasospasm following acute aneurysmal subarachnoid haemorrhage.

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)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Cardiovascular: Ischaemia

Interventions

Prospective imaging study (analysis blinded to clinical outcome).

Intervention Type

Other

Phase

Not Applicable

Primary outcome measure

Not provided at time of registration

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/05/2002

Completion date

01/05/2003

Eligibility

Key inclusion criteria

Not provided at time of registration

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

Initially 10 patients and 10 controls, total 20 participants

Key exclusion criteria

Not provided at time of registration

Date of first enrolment

01/05/2002

Date of final enrolment

01/05/2003

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

Department of Imaging

London

United Kingdom

W6 8RF

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

Hammersmith Hospital NHS 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 summary**

Not provided at time of registration