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How effective is hand massage in stress relief for patients waiting for coronary angiography?

Submission date 30/09/2004	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol
Registration date 30/09/2004	**Overall study status** Completed	[ ] Statistical analysis plan [ ] Results
Last Edited 04/09/2015	**Condition category** Signs and Symptoms	[ ] 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

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
N0205139800

Study information

Scientific Title
How effective is hand massage in stress relief for patients waiting for coronary angiography?

Study objectives
The project aims to identify the stress levels for patients waiting for coronary angiography, and whether these can be modified with the use of massage therapy.

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
Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied
Signs and Symptoms: Stress

Interventions
The study takes the form of a randomised control trial.

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
A recognised tool for measurement of stress will be used called the state-trait anxiety score. Clients will also be asked to complete questionnaires regarding massage.

Secondary outcome measures
Not provided at time of registration

Overall study start date
20/01/2004

Completion date
30/09/2004

Eligibility

Key inclusion criteria
The study aims to identify 60 patients to recruit.

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
60

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
20/01/2004

Date of final enrolment
30/09/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
The Royal London Hospital
London
United Kingdom
E1 1BB

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.dh.gov.uk/Home/fs/en

Funder(s)

Funder type
Government

Funder Name
Barts and The London NHS Trust

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