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Evaluation of relaxation techniques to reduce anxiety in patients undergoing MRI investigations

Submission date 30/09/2005	**Recruitment status** No longer recruiting	[ ] Prospectively registered [ ] Protocol [ ] Statistical analysis plan [ ] Results [ ] Individual participant data [ ] Record updated in last year
Registration date 30/09/2005	**Overall study status** Completed	
Last Edited 05/07/2016	**Condition category** Mental and Behavioural Disorders	

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

N0176148645; Protocol Dated 2003

Study information

Scientific Title
Evaluation of relaxation techniques to reduce anxiety in patients undergoing MRI investigations

Study objectives
This study is to test how effective relaxation techniques are for reducing anxiety in patients undergoing MRI (magnetic resonance imaging) investigations, and thus enable the patients to better tolerate the procedure.

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)
Quality of life

Participant information sheet
Not available in web format, please use contact details to request a participant information sheet

Health condition(s) or problem(s) studied
Anxiety disorders

Interventions
Randomised controlled trial with questionnaire component

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure

Writing up of study following data collection and analysis from 30 recruited patients for completion of course in May 04.

Secondary outcome measures
Not provided at time of registration

Overall study start date
16/03/2004

Completion date
31/12/2004

Eligibility

Key inclusion criteria
20 patients, 10 control patients

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
30

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
16/03/2004

Date of final enrolment
31/12/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
Radcliffe Infirmary
Oxford

United Kingdom
OX2 6HE

Sponsor information

Organisation
Department of Health

Sponsor details
Richmond House
79 Whitehall
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SW1A 2NL
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Sponsor type
Government

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

Funder(s)

Funder type
Government

Funder Name
Oxford Radcliffe Hospitals 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