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

Submission date	Recruitment status	Prospectively registered
30/09/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
30/09/2005	Completed	Results
Last Edited	Condition category	☐ Individual participant data
05/07/2016	Mental and Behavioural Disorders	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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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

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

Sponsor information

Organisation

Department of Health

Sponsor details

Richmond House 79 Whitehall London United Kingdom SW1A 2NL +44 (0)20 7307 2622 dhmail@doh.gsi.org.uk

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