

Anger Management, a randomised, controlled trial

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 16/07/2009	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<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
N0230117542

Study information

Scientific Title

Study objectives

To assess the effectiveness of anger management by employing a cognitive behavioural framework.

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)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Anger management

Interventions

Study subjects will be selected from those referred to the Anger Management group run at the Department of Psychiatry, Royal South Hants Hospital. It will compare a cognitive behaviour therapy based anger management program with a control group of patients on the waiting list. The study will last for 9 months. <p>Questionnaires will be used at the start, end of group therapy (3 months after the start) and again at 6 months follow up.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Anger will be measured using the Navco Anger Assessment Scale (NAS) and State-Trait Anger Expression Inventory (STAXI). The Clinical Outcomes in Routine Elevations (CORE) scale will be used to measure the well-being and life functioning of the subjects. Depression and anxiety will

be measured using the Hospital Anxiety and Depression Scale. Evaluation belief scale will be used to measure the beliefs about self and others. Other variables to be collected include: age, gender, ethnicity, marital status and employment status.

Secondary outcome measures

Not provided at time of registration

Overall study start date

01/03/2003

Completion date

01/03/2004

Eligibility

Key inclusion criteria

30 Subjects within age range 16-65 from patients referred for Anger Management training.
<p>Inclusion criteria: a history of being unable to manage anger with or without a history of mental illness.

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

30

Key exclusion criteria

1. excessive use of drugs and/or alcohol
2. significant cognitive impairment
3. active psychosis.

Date of first enrolment

01/03/2003

Date of final enrolment

01/03/2004

Locations

Countries of recruitment

England

United Kingdom

Study participating centre
Dept of Psychiatry
Southampton
United Kingdom
S014 0YG

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
Research council

Funder Name
West Hampshire Consortium (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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/03/2009		Yes	No