Anger Management, a randomised, controlled trial

| Recruitment status No longer recruiting | Prospectively registered | |
|--|---|--|
| | ☐ Protocol | |
| Overall study status | Statistical analysis plan | |
| Completed | [X] Results | |
| Condition category Montal and Robavioural Disorders | Individual participant data | |
| | No longer recruiting Overall study status Completed | |

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Dr Faroog Naeem

Contact details

Dept of Psychiatry Royal South Hants Hospital Brintons Terrace Southampton United Kingdom S014 0YG

Additional identifiers

Protocol serial number 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

Study type(s)

Not Specified

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. 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(s)

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.

Key secondary outcome(s))

Not provided at time of registration

Completion date

01/03/2004

Eligibility

Key inclusion criteria

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

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Αll

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

United Kingdom

England

Study participating centre Dept of Psychiatry

Southampton United Kingdom S014 0YG

Sponsor information

Organisation

Department of Health (UK)

Funder(s)

Funder type

Research council

Funder Name

West Hampshire Consortium (UK)

Results and Publications

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 |