

Systematic Intervention to Transform Environment: a randomised controlled trial of nidoththerapy in an assertive outreach team

Submission date
15/10/2006

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
06/02/2007

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
08/06/2011

Condition category
Mental and Behavioural Disorders

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

Prof Peter Tyrer

Contact details

Department of Psychological Medicine
Imperial College
St Dunstan's Road
London
United Kingdom
W8 8RP
+44 (0)20 7386 1237
p.tyrer@imperial.ac.uk

Additional identifiers

Protocol serial number

site/DNMM06

Study information

Scientific Title

Acronym

SITE

Study objectives

To test the feasibility and likely effectiveness of nidothrapy, a new treatment for personality disorder and chronic mental illness, in a group that is most difficult to treat - those cared for by an assertive outreach team, in an exploratory randomised controlled trial.

Ethics approval required

Old ethics approval format

Ethics approval(s)

St Marys Hospital Research Ethics Committee on 16/07/2003 (ref: R&D03/X008E).

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Co-morbid personality disorder and severe mental illness (schizophrenia, bipolar disorder or severe depressive illness)

Interventions

Nidothrapy (a new treatment designed to change the environment to suit the needs of the peson with the chronic disorder) versus assertive treatment as usual in the team.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Reduction of usage of psychiatric beds one year after treatment

Key secondary outcome(s)

1. Reduction in number of admissions
2. Social function (using the Social Functioning Questionnaire)
3. Psychiatric symptomatology (using Brief Psychiatric Rating Scale [BPRS])
4. Patient satisfaction (Consultation Satisfaction Questionnaire [CSQ])
5. Engagement with services (Engagement and Acceptance Scale [EAS])

Completion date

31/12/2006

Eligibility

Key inclusion criteria

Randomisation of all eligible patients who give consent currently under the care of an assertive outreach team.

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Not Specified

Sex

Not Specified

Key exclusion criteria

Those who refuse to take part.

Date of first enrolment

01/10/2003

Date of final enrolment

31/12/2006

Locations

Countries of recruitment

United Kingdom

England

Study participating centre

Department of Psychological Medicine

London

United Kingdom

W8 8RP

Sponsor information

Organisation

Funder(s)

Funder type

Charity

Funder Name

Nicola Pigott Memorial Fund (UK) (ref: CNWL019070)

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	01/04/2009		Yes	No