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Patient centred work in psychiatric care: the Weddinger Modell

Submission date	Recruitment status	[ ] Prospectively registered
03/02/2011	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
22/09/2011	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
22/09/2011	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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
Version I 31.05.2010

Study information

Scientific Title

Patient centered work-satisfaction and resilience in the Trialogue - an observational study (Weddinger Modell) (Patientenzentriertes Arbeiten- Zufriedenheit und Resilienz im Trialog - eine Beobachtungsstudie (Das "Weddinger Modell")

Acronym
Weddinger Modell

Study objectives
The aim of the study is a novel patient centred approach (Weddinger Modell) in psychiatric care for problematic urban districts. Improve measures of patient contentedness, resilience, goal-attainment and therapeutic relationship evaluated by patients, relatives and staff.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Ethics Committee of the Charité-University Medicine Berlin (Ethikkommission der Charité - Universitätsmedizin) approved on 29th June 2010, ref: EA4/074/10

Study design
Controlled parallel group observational study

Primary study design
Observational

Secondary study design
Non randomised controlled trial

Study setting(s)
GP practice

Study type(s)
Quality of life

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

Health condition(s) or problem(s) studied
Psychiatric disorders leading to hospital admission

Interventions
A controlled, clinical, longitudinal trial using trialogical questionnaires including self-rating as well as ratings by relatives and by professionals.

Intervention Type
Other

Phase
Not Applicable

Primary outcome measure
1. WPAZ
2. Resilence-Questionnaire
3. Work Ability Index (WAI)
4. Goal-Attainment-Scale, Global Assessment of Functional Scale (GAF)
5. Staff contentedness

Measured before and after the intervention.

Secondary outcome measures
Need for hospital admission and extent of comunity-based care one year prior and one year after inclusion

Overall study start date
01/09/2010

Completion date
31/12/2013

Eligibility

Key inclusion criteria
1. Age more than 18 years
2. All admitted patients -patients treated in three psychiatric hospitals in Berlin, who give informed consent
3. Relatives of admitted patients
4. Staff

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Sex
Both

Target number of participants
1200

Key exclusion criteria
1. Duration of stay in hospital less than 24 hours
2. Suicidal indication and severe aggression
3. Serious organic brain disease

Date of first enrolment
01/09/2010

Date of final enrolment
31/12/2013

Locations

Countries of recruitment
Germany

Study participating centre
Große Hamburger Strasse 5-11
Berlin
Germany
10115

Sponsor information

Organisation
Charité - University Medicine Berlin (Charité - Universitätsmedizin Berlin) (Germany)

Sponsor details
c/o Dr. Lieselotte Mahler
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Sponsor type
Hospital/treatment centre

ROR
https://ror.org/001w7jn25

Funder(s)

Funder type
Hospital/treatment centre

Funder Name
Charité University Medicine (Germany)

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