Outcome monitoring and outcome management in inpatient psychiatric care (Ergebnis-Monitoring und Ergebnis-Management in der stationären psychiatrischen Versorgung)

Submission date	Recruitment status	[X] Prospectively registered
05/08/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
15/08/2005	Completed	Results
Last Edited	Condition category	Individual participant data
15/09/2009	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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Additional identifiers

Protocol serial number

01 GL 0504

Study information

Scientific Title

Acronym

EMM

Study objectives

To examine whether continuous monitoring and management of outcome (EMM) improves the quality of inpatient psychiatric care and contributes to an adaptive allocation of treatment resources. Hypotheses are that EMM:

- 1. Leads to a short and medium-term improvement of the clinical treatment outcome
- 2. Is cost effective because of a reduction of length of stay (LOS) and decrease of subsequent treatment costs
- 3. Is feasible i.e. can also be used with severely impaired patients
- 4. Causes higher clinician and patient satisfaction with treatment

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)

Treatment

Health condition(s) or problem(s) studied

Severe mental disorder

Interventions

Outcome monitoring and management (intervention group) versus outcome monitoring (control group)

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Independent rating of treatment outcome (psychological impairment)

Key secondary outcome(s))

- a. Patient and clinician rating of outcome
- b. Costs for utilisation of health care services (from health insurance files and patient information)
- c. Patient and clinician satisfaction

Completion date

01/07/2007

Eligibility

Key inclusion criteria

Patients at the age of 18 to 65 years admitted to a large psychiatric hospital in rural Bavaria ('Bezirkskrankenhaus Guenzburg')

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

65 years

Sex

All

Key exclusion criteria

Primary diagnosis of substance disorder, organic brain disorder, or dementia; insufficient knowledge of the German language, limited reading and/or writing abilities.

Date of first enrolment

01/09/2005

Date of final enrolment

01/07/2007

Locations

Countries of recruitment

Germany

Study participating centre

University of Ulm

Guenzburg Germany D-89312

Sponsor information

Organisation

German Federal Ministry of Education and Research, Project Management by the German Aerospace Center (DLR)

ROR

https://ror.org/04bwf3e34

Funder(s)

Funder type

Government

Funder Name

German Federal Ministry of Education and Research (Germany)

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration