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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

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
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

Secondary study design
Single-centre

Study setting(s)
Hospital

Study type(s)
Treatment

Participant information sheet

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 measure
Independent rating of treatment outcome (psychological impairment)

Secondary outcome measures
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

Overall study start date
01/09/2005

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

Age group
Adult

Lower age limit
18 Years

Upper age limit
65 Years

Sex
Both

Target number of participants
294

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
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Sponsor information

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

Sponsor details
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Sponsor type
Government

Website
http://www.bmbf.de

ROR
https://ror.org/04bwf3e34

Funder(s)

Funder type
Government

Funder Name
German Federal Ministry of Education and Research (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