

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

Submission date 05/08/2005	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 15/08/2005	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 15/09/2009	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> 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

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

Primary study design

Interventional

Study design

Randomised controlled trial

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