ISRCTN19638644 https://doi.org/10.1186/ISRCTN19638644

How can rehospitalisations of patients with schizophrenia be avoided? A comparison between different compliance programs

Submission date	Recruitment status	[ ] Prospectively registered
02/11/2006	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
14/02/2007	Completed	[ ] Results
Last Edited	**Condition category**	[ ] Individual participant data
01/02/2019	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
NCT00514423

Secondary identifying numbers

01GL0509

Study information

Scientific Title
How can rehospitalisations of patients with schizophrenia be avoided? A comparison between different compliance programs

Study objectives
Participation in one of the three interventions (psychoeducation by professionals, psychoeducation by peer-moderators or video-education) can reduce the rehospitalisation rate of schizophrenic patients compared to a control group.

Ethics approval required
Old ethics approval format

Ethics approval(s)
Approval received from the local ethics committee on the 10th February 2006 (ref: 1468/06).

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Treatment

Participant information sheet

Health condition(s) or problem(s) studied
Schizophrenia or schizoaffective disorder

Interventions
1. Psychoeducation by professionals
2. Psychoeducation by peer-moderators
3. Video-education

Intervention Type
Other

Phase
Not Specified

Primary outcome measure
Rehospitalisation rate

Secondary outcome measures
1. Rehospitalisation days
2. Resulting costs per patient
3. Cost-benefit-analysis of the different interventions
4. Compliance
5. Compliance self-assesment (Medication Adherence Rating Scale [MARS]/Drug Attitude Inventory [DAI])
6. Knowledge of illness
7. Attitude towards illness
8. Quality of life (World Health Organisation Quality Of Life [WHOQOL]-BREF)
9. Satisfaction questionnaire (ZUF-8)
10. Duration of consultation time
11. Dealing with mental illness (Mantonakis, Family Member Questionnaire [FMQ], Beck Depression Inventory [BDI])

Overall study start date
01/09/2006

Completion date
31/12/2007

Eligibility

Key inclusion criteria
1. Schizophrenia or schizoaffective disorder (International Classification of Diseases [ICD-10])
2. Age 18 to 67 years
3. Hospitalised or treated in a day care clinic

Participant type(s)
Patient

Age group
Adult

Lower age limit
18 Years

Upper age limit
67 Years

Sex
Not Specified

Target number of participants
896 patients, family members as available

Key exclusion criteria

1. More than 12 months of hospitalisation within the last two years
2. Substance-dependency (principal diagnosis)
3. Mental retardation ICD-10 Chapter F70-79
4. Fluency of German language not given

Date of first enrolment
01/09/2006

Date of final enrolment
31/12/2007

Locations

Countries of recruitment
Germany

Study participating centre
Technical University Munich
Muenchen
Germany
81675

Sponsor information

Organisation
Technical University Munich (Germany)

Sponsor details
Klinikum rechts der Isar
Department of Psychiatry and Psychotherapy
Moehlstrasse 26
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81675

Sponsor type
University/education

Website
http://www.tum.de/

ROR
https://ror.org/02kkvpp62

Funder(s)

Funder type
Government

Funder Name
Federal Ministry of Education and Research (Bundesministerium für Bildung und Forschung [BMBF]) (Germany) (grant ref: 01GL0509)

Alternative Name(s)
Federal Ministry of Education and Research, BMBF

Funding Body Type
Government organisation

Funding Body Subtype
National government

Location
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