Enhancing the quality of Life for people with severe mental health problems through supported employment

Submission date	Recruitment status No longer recruiting	Prospectively registered		
20/12/2005		☐ Protocol		
Registration date 20/12/2005	Overall study status Completed	Statistical analysis plan		
		[X] Results		
Last Edited 04/07/2019	Condition category Mental and Behavioural Disorders	Individual participant data		

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 NTR295

Study information

Scientific Title

A randomised controlled trial of the cost-effectiveness of individual placement and support in six European countries

Acronym

EQOLISE

Study objectives

Individual placement and supported (IPS) will be cost-effective in achieving regular (open, paid) employment. Differences between countries in the effectiveness of IPS can be explained by socio-economic variables like the employment rate and for instance the existence of a 'benefit trap'.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Received from the local medical ethics committee

Study design

Multicentre, randomised, active controlled, parallel group trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Psychiatric, mental disorders/illness

Interventions

Individual placement and support (EXP) will be compared to standard vocational rehabilitation (CC).

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

- 1. Achievement of open, paid employment
- 2. Hours of open employment
- 3. Time in open employment

Key secondary outcome(s))

- 1. Clinical improvement
- 2. Hospital admissions
- 3. Unmet needs for care
- 4. Quality of life

Completion date

01/01/2006

Eligibility

Key inclusion criteria

- 1. Diagnosis of a serious mental illness: schizophrenia, schizophrenia-like disorders, bipolar disorder with psychotic features
- 2. Age 18 65 years
- 3. Living in the community at baseline
- 4. Not employed for more then a month in the year before the start of the study
- 5. In mental health care for more then two years
- 6. Interest in competitive employment
- 7. Willing to give informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Total final enrolment

312

Key exclusion criteria

Use of the EXP or CC service in the preceding year

Date of first enrolment

01/01/2003

Date of final enrolment

01/01/2006

Locations

Countries of recruitment

Netherlands

Study participating centre

University Medical Centre Groningen

Groningen Netherlands 9700 RB

Sponsor information

Organisation

University Medical Centre Groningen (UMCG) (The Netherlands)

ROR

https://ror.org/03cv38k47

Funder(s)

Funder type

Government

Funder Name

European Community

Results and Publications

Individual participant data (IPD) sharing plan

IPD sharing plan summary

Not provided at time of registration

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/09/2009	04/07/2019	Yes	No