

ENhancing DELivery And outcomes of VOcational Rehabilitation

Submission date 18/06/2010	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 18/06/2010	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 27/05/2014	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data

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
6104

Study information

Scientific Title
Enhancing delivery and outcomes of vocational rehabilitation (ENDEAVOR): improving outcomes for young people with a first episode psychosis

Acronym

ENDEAVOR

Study objectives

Primary hypothesis:

To compare the effectiveness of two vocational rehabilitation interventions in helping people attain and retain employment following a first episode of psychosis.

Secondary hypothesis:

To describe the pathways, barriers and facilitators to employment in a cohort of people suffering a first episode psychosis.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Charing Cross Research Ethics Committee approved on the 15th December 2008 (ref: 08/H0711 /136)

Study design

Multicentre randomised interventional treatment trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Topic: Mental Health Research Network; Subtopic: Psychosis; Disease: Schizophrenia

Interventions

1. Supported employment
2. Return to employment/education after first episode psychosis

Study entry: single randomisation only

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Employment status, measured at 12 months

Key secondary outcome(s)

Measured at 12 months:

1. Employment and education/training across follow up; qualifications obtained
2. Positive and Negative Symptom Scale (PANSS)

3. Global Assessment of Function
4. Time Budget
5. Client Service Receipt Inventory

Completion date

30/09/2011

Eligibility

Key inclusion criteria

1. First episode psychosis
2. Aged 18 - 35 years, either sex

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Upper age limit

35 years

Sex

All

Key exclusion criteria

1. Organic disease
2. Unable to comprehend written English
3. Unable to give informed consent

Date of first enrolment

01/06/2009

Date of final enrolment

30/09/2011

Locations

Countries of recruitment

United Kingdom

England

Study participating centre
Institute Of Psychiatry
London
United Kingdom
SE5 8AF

Sponsor information

Organisation

Kings College London (KCL) (UK)

ROR

<https://ror.org/0220mzb33>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research (NIHR) (UK) - Research for Patient Benefit (RfPB) programme

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	01/08/2014		Yes	No