

WELLFOCUS study: to investigate an intervention to improve well-being in people with psychosis

Submission date 12/11/2012	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 14/12/2012	Overall study status Completed	<input checked="" type="checkbox"/> Protocol
Last Edited 04/10/2018	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

Background and study aims

Well-being is important for everyone, including people with severe mental illness. Well-being can help to improve functioning, resilience, and life satisfaction and may protect against mental illness.

Positive Psychotherapy (PPT) was developed in the field of positive psychology. It aims to increase well-being by building positive emotions, character strengths, and meaning. So far, PPT has been shown to decrease symptoms and increase well-being in people with depression and other common mental disorders. There are no established interventions to increase well-being in people with severe mental illness, but PPT is a promising approach. In a previous project we adapted PPT to be suitable as a group therapy for people with psychosis. We have done so with the input of service users, professional carers, and experts in the field. In this study we will show how the adapted intervention can be best delivered to people with psychosis.

Who can participate?

Adults with an experience of psychosis from a London NHS Trust.

What does the study involve?

Participants will receive the WELLFOCUS programme, in group therapy format in addition to their usual care, or continue to receive their usual care as before. Participants will be assessed with a range of questionnaires before and after the study period and they will be asked to participate in personal interviews and focus groups.

What are the possible benefits and risks of participating?

Participants may experience increased personal well-being and there are no known risks for participants.

Where is the study run from?

Institute of Psychiatry at Kings College London, UK

When is study starting and how long is it expected to run for?
Recruitment will start in early 2013 the study is expected to run until 2015.

Who is funding the study?
Guys & St Thomas Charity, UK

Who is the main contact?
Dr Beate Schrank
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Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

Protocol serial number
G101016

Study information

Scientific Title
Pilot randomised controlled trial of a modified intervention to improve well-being in people with psychosis

Acronym
WELLFOCUS

Study objectives
No specific hypothesis is tested as this is a pilot randomised controlled study. This means that it will establish if the intervention works as it is expected to and if it can be delivered as planned in a research setting. The results will help to further adapt the intervention and the research process and help to plan a big study to investigate the intervention.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Pilot randomised controlled trial

Primary study design

Interventional

Study type(s)

Quality of life

Health condition(s) or problem(s) studied

Psychosis

Interventions

Participants will be randomised into either of the following groups:

1. The WELLFOCUS intervention: adapted PPT for people with psychosis in a group format, once a week over 12 weeks, in addition to treatment as usual.
2. Treatment as usual

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

Personal well-being assessed using the Warwick-Edinburgh Mental Well-Being Scale (WEMWBS)

Key secondary outcome(s)

1. Quality of life as assessed by the Manchester Short Assessment of Quality of Life (MANSA)
2. Happiness as assessed by the Short Depression-Happiness Scale (SDHS)
3. Hope as assessed by the Integrative Hope Scale (IHS)
4. Savouring as assessed by the Savouring Beliefs Inventory (SBI)
5. Symptoms and functioning as assessed by Brief Psychiatric Rating Scale (BPRS)
6. The Health of the Nation Outcome Scale (HoNOS)
7. The Global Assessment of Functioning (GAF)

Completion date

01/01/2015

Eligibility

Key inclusion criteria

Adults with a primary diagnosis of psychosis who are not currently receiving in-patient care or are in prison, speak and understand English and are sufficiently well to participate

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

All

Key exclusion criteria

1. Serious cognitive impairment
2. Unable to give informed consent

Date of first enrolment

01/01/2013

Date of final enrolment

01/01/2015

Locations**Countries of recruitment**

United Kingdom

England

Study participating centre

Kings College London

London

United Kingdom

SE5 8AF

Sponsor information**Organisation**

Kings College London (UK)

ROR

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

Funder(s)

Funder type

Charity

Funder Name

Guy's and St Thomas' Charity ref: G101016

Alternative Name(s)

Guy's and St Thomas' Charity, Guy's and St Thomas' Foundation, GSTTFoundation

Funding Body Type

Private sector organisation

Funding Body Subtype

Trusts, charities, foundations (both public and private)

Location

United Kingdom

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	01/01/2015		Yes	No
Results article	results	01/06/2016		Yes	No
Protocol article	protocol	03/06/2014		Yes	No
Participant information sheet	Participant information sheet	11/11/2025	11/11/2025	No	Yes
Study website	Study website	11/11/2025	11/11/2025	No	Yes