Adherence therapy for people with schizophrenia in Suanprung Psychiatric Hospital, Thailand: a randomised controlled trial

Submission date	Recruitment status	Prospectively registered	
24/10/2005	No longer recruiting	☐ Protocol	
Registration date	Overall study status	Statistical analysis plan	
26/04/2006	Completed	[X] Results	
Last Edited	Condition category	Individual participant data	
20/01/2020	Mental and Behavioural Disorders		

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

Nil known

IRAS number

ClinicalTrials.gov number

Nil known

Secondary identifying numbers

N/A

Study information

Scientific Title

Adherence therapy for people with schizophrenia in Suanprung Psychiatric Hospital, Thailand: a randomised controlled trial

Study objectives

Adherence therapy can reduce symptoms, increase attitude and satisfaction towards medication treatment

Ethics approval required

Old ethics approval format

Ethics approval(s)

Not provided at time of registration

Study design

Randomised controlled trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Hospital

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Schizophrenia

Interventions

Treatment group: adherence therapy Control group: treatment as usual such as medication treatment, group counselling, occupational and recreational therapy

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Symptoms of schizophrenia measured using the positive and negative syndrome scale

Secondary outcome measures

- 1. Symptoms of schizophrenia measured using the Global Assessment of Functioning Scale
- 2. Attitude towards medication measured using the Hogan Drug Attitude Inventory
- 3. Satisfaction with medication measured using the Satisfaction With Antipsychotic Medication Scale
- 4. Side effects of medication measured using the Liverpool University Neuroleptic Side Effect Rating Scale

Overall study start date

01/12/2004

Completion date

31/08/2005

Eligibility

Key inclusion criteria

- 1. Patients diagnosed by a psychiatrist with schizophrenia according to ICD-10 criteria (World Health Organisation ([WHO], 1992)
- 2. People with schizophrenia who are older than 20 years of age because they have reached the age of maturity legally and can provide consent independently
- 3. People with schizophrenia who have been assessed by a psychiatrist not involved in this study to determine that they can provide informed consent
- 4. People with schizophrenia who are residing in Muang Chiang Mai district

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

70

Total final enrolment

32

Key exclusion criteria

Patients suffering from organic brain diseases or learning disability

Date of first enrolment

01/12/2004

Date of final enrolment

Locations

Countries of recruitment

England

Thailand

United Kingdom

Study participating centre P O Box 30

London United Kingdom SE5 8AF

Sponsor information

Organisation

The Royal Thai Government

Sponsor details

Office of Educational Affairs Royal Thai Embassy 28 Prince's Gate London Thailand SW7 1QF +44 (0)207 584 4538 (5 lines) enquiries@oeathai.org

Sponsor type

Government

Website

http://www.oeathai.org

ROR

https://ror.org/05fp08r12

Funder(s)

Funder type

Funder Name

Praboromarajanok Institute of Workforce Development (Thailand)

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

Study outputs

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/07/2007	31/05/2019	Yes	No