Study on Cost-Effectiveness of Personality disorder TREatment

Submission date	Recruitment status	Prospectively registered
28/12/2006	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
28/12/2006	Completed	☐ Results
Last Edited	Condition category	Individual participant data
16/01/2007	Mental and Behavioural Disorders	Record updated in last year

Plain English summary of protocol

Not provided at time of registration

Study website

http://www.vispd.nl/projects.htm

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Acronym

SCEPTRE

Study objectives

- 1. Effect sizes of the psychotherapeutic treatments vary between 0.5 and 2.0.
- 2. A lower dosage of psychotherapeutic treatment is mainly related to symptomatic improvement, while a higher dosage is also related to structural and functional improvement.
- 3. The dosage for an optimal cost-effectiveness is lower than the maximum dosage.
- 4. When compared to cluster B personality disorders, for cluster C personality disorders a lower dosage is required to reach optimal cost-effectiveness.
- 5. A lower level of psychological mindedness and/or ability for a positive working alliance with the therapist is related to a lower level of (cost)-effectiveness.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approval received from the Research Ethics Committee (REC) of Erasmus Medical Centre, Rotterdam, on the 7th January 2004 (ref: MEC 229.834/2003/122).

Study design

Non-randomised clinical trial

Primary study design

Interventional

Secondary study design

Multi-centre

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Personality Disorder

Interventions

Psychotherapeutic treatment programs for patients with personality pathology that will be distinguished from one another according to the following dosages:

- 1. Short (maximum of six months) outpatient
- 2. Short day hospital
- 3. Short inpatient
- 4. Long (at least six months) outpatient
- 5. Long day hospital
- 6. Long inpatient

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Structural improvement of personality pathology (assessed with the Outcome Questionnaire [OQ45], and the Severity Indices of Personality Problems [SIPP])
- 2. Quality Adjusted Life Years (QALYs)

Secondary outcome measures

- 1. Symptomatic improvement (assessed with the Symptoms CheckList [SCL-90])
- 2. Functional improvement (assessed with, amongst others, questions regarding work situation and care consumption)
- 3. Quality of life

Overall study start date

01/09/2003

Completion date

31/12/2008

Eligibility

Key inclusion criteria

- 1. At least one Diagnostic and Statistical Manual of Mental Disorders Fourth Edition (DSM-IV) axis-II diagnosis
- 2. An indication for psychotherapeutic treatment aiming at structural improvement of the personality pathology (besides symptomatic and/or functional improvement)

Participant type(s)

Patient

Age group

Not Specified

Sex

Not Specified

Target number of participants

821

Key exclusion criteria

- 1. Insufficient command of Dutch language
- 2. Severe cognitive impairments
- 3. Mental retardation
- 4. Axis-I schizophrenia

Date of first enrolment

01/09/2003

Date of final enrolment

31/12/2008

Locations

Countries of recruitment

Netherlands

Study participating centre

Viersprong Institute for Studies on Personality Disorders (VISPD)

Halsteren Netherlands 4660 AA

Sponsor information

Organisation

Viersprong Institute for Studies on Personality Disorders (VISPD) (The Netherlands)

Sponsor details

P.O. Box 7 Halsteren Netherlands 4660 AA

Sponsor type

Research organisation

Website

http://www.vispd.nl/

ROR

https://ror.org/048jnwk41

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Psychotherapeutic Center De Viersprong (The Netherlands)

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