Psychotherapy for residual depression following initial treatment: effectiveness, relapse prevention and mechanisms of change

Published: 05-12-2006 Last updated: 09-05-2024

To compare the effectiveness of CBT versus IPT in the prevention of future relapse among residually depressed patients who received initial treatment. Our research questions: Are CBT and IPT following initial treatment effective interventions that...

Ethical review Approved WMO **Status** Recruiting

Health condition type Mood disorders and disturbances NEC

Study type Interventional

Summary

ID

NL-OMON30225

Source

ToetsingOnline

Brief title

Psychotherapy for residual depression following initial treatment

Condition

Mood disorders and disturbances NEC

Synonym

depression, mood disorder

Research involving

Human

Sponsors and support

Primary sponsor: Universiteit Maastricht

1 - Psychotherapy for residual depression following initial treatment: effectiveness ... 29-05-2025

Source(s) of monetary or material Support: Ministerie van OC&W

Intervention

Keyword: cognitive behaviour therapy, depression, mechanisms of change, relapse prevention

Outcome measures

Primary outcome

Depressive relapse/recurrence over the course of two years.

Secondary outcome

Severity of depressive symptoms; episodes of major depression according to the DSM-IV; other Axis I and Axis II disorders according to the DSM-IV; psychological problems; health care consumption; social functioning; patient perspectives; mechanism of change variables (explicit and implicit); genetic material (buccal mucous membrane).

Study description

Background summary

Although psychotherapy and antidepressants seem to help initially, many depressed patients suffer from relapse and recurrence. Recent findings suggest cognitive behaviour therapy (CBT) may reduce that risk in the long-term, but the mechanisms of change that prevent relapse and recurrence are still unknown. We will be the first to study the effectiveness of CBT compared to interpersonal therapy (IPT) for residual depression after initial treatment (reduction of symptoms; prevention of relapse and recurrence) and the underlying mechanisms of change (explicit and implicit measures).

Study objective

To compare the effectiveness of CBT versus IPT in the prevention of future relapse among residually depressed patients who received initial treatment. Our research questions: Are CBT and IPT following initial treatment effective interventions that prevent relapse or recurrence of depression in the

long-term? What are the underlying psychological mechanisms of change in CBT, compared to IPT? Do genetic variations in polymorphisms modify the short and long-term efficacy of CBT and IPT?

Study design

Randomized trial with three groups: CBT (n=75), IPT (n=75) and an 8-week waitinglist condition (n=30)

Intervention

Cognitive behaviour therapy (CBT) and interpersonal therapy (IPT). CBT and IPT will be offered at our clinical site (Riagg Maastricht). Both interventions are written out in a treatment manual, will be delivered by qualified therapists and will both contain 18 sessions and two booster sessions.

Study burden and risks

Participants will have to visit the research centre twice, and will spend approximately 20 hours filling out questionnaires or being interviewed over the course of 24 months. Part of the assessments is a clinical interview at the beginning of the study and at the end of follow-up, and a computer task in an observation laboratory. No risks are involved in participation.

Contacts

Public

Universiteit Maastricht

Postbus 616 6200 MD Maastricht NL

Scientific

Universiteit Maastricht

Postbus 616 6200 MD Maastricht NL

Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

Adults (18-64 years) Elderly (65 years and older)

Inclusion criteria

depressive symptoms (BDI > 10)

Exclusion criteria

concurrent treatment for depressive symptoms

Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

Control: Active

Primary purpose: Basic science

Recruitment

NL

Recruitment status: Recruiting
Start date (anticipated): 01-02-2007

Enrollment: 180

Type: Actual

Medical products/devices used

Registration: No

Ethics review

Approved WMO

Date: 05-12-2006

Application type: First submission

Review commission: METC academisch ziekenhuis Maastricht/Universiteit

Maastricht, METC azM/UM (Maastricht)

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register ID

CCMO NL14107.068.06