The efficacy of Attentional Training on Attentional Bias and Symptoms of Post-Traumatic Stress Disorder

Published: 27-12-2006 Last updated: 14-05-2024

To investigate the efficacy of attentional training

Ethical review Approved WMO

Status Recruitment stopped

Health condition type Anxiety disorders and symptoms

Study type Interventional

Summary

ID

NL-OMON29799

Source

ToetsingOnline

Brief title

Attentional training in PTSD

Condition

Anxiety disorders and symptoms

Synonym

psychotrauma, soldiers disease

Research involving

Human

Sponsors and support

Primary sponsor: Universiteit Leiden

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

Intervention

Keyword: attention, cognition, PTSD, training

Outcome measures

Primary outcome

Attentional Bias and Posttraumatic Symptoms

Secondary outcome

Depressive symptoms

Study description

Background summary

Efficacy of Attentional Training on Attentional Bias and Symptoms of Post-Traumatic Stress Disorder.

The symptoms of Posttraumatic Stress Disorder (PTSD) include many cognitive phenomena such as: re-experiencing the trauma, recurrent and intrusive recollections, nightmares, flashbacks, hyper vigilance and memory loss. Many studies have reported memory and attention problems in PTSD (Buckley, Blanchard, & Neill, 2000). Attentional bias (AB) is defined as the automatic and selective allocation of attentional resources to threatening and negative information, and is thought to be an important contributing factor in the development and maintenance of PTSD. AB is measured using reaction time tests. Recently, it has been demonstrated that AB can either be created or eliminated in the laboratory by using adapted versions of the attentional tests (*attentional training*). Preliminary evidence suggests that it may be effective as a treatment in anxiety disorders, e.g., social phobia. AB is larger in PTSD than in any other anxiety disorder.

Outcomes: Attentional bias, PTSD symptoms.

Investment: diagnostic interview, questionnaires, assessment of AB (2 hrs

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combined); Attentional training or placebo (8 times 20 minutes); Outcome assessment (30 minutes).

Risks: none. None of these procedures or questionnaires is any different from the procedures or questions that are asked in the context of their ongoing treatment. The attentional training task is considered boring by some participants (pilot data in students).

Study objective

To investigate the efficacy of attentional training

Study design

Randomized, double blind placebo-controlled trial.

Intervention

Attentional training or placebo training.

Study burden and risks

Participants will spend 8 hours in total in this researchproject. There are no risks to be expected.

Contacts

Public

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

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

Inclusion criteria

DSM-IV diagnosis of Post Traumatic Stress Disorder Score 24 points or higher on the Impact of Event Scale (IES) Attentional bias (Dot Probe test)

Exclusion criteria

psychotic disorder (lifetime) alcohol or drug abuse or dependence (current) high suicide risk (measured in a semi-structured interview)

Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Double blinded (masking used)

Control: Placebo

Primary purpose: Treatment

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 15-11-2007

Enrollment: 60

Type: Actual

Ethics review

Approved WMO

Application type: First submission

Review commission: METC Leiden-Den Haag-Delft (Leiden)

metc-ldd@lumc.nl

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 NL12042.058.06