

Effectiveness of adding 'exposure in vivo' techniques to the return-to-work plan of workers with mental health problems: a cluster randomised controlled trial

Submission date 11/04/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
Registration date 11/04/2007	Overall study status Completed	<input type="checkbox"/> Protocol
Last Edited 13/09/2013	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

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

Acronym

Work up study

Study objectives

Occupational rehabilitation with a gradual return to work based on the principles of exposure in vivo will be more (cost)-effective in reducing absenteeism than usual occupational rehabilitation.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Study design

Randomised, active controlled, parallel group, multicentre trial

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Return to work of workers with common mental health complaints

Interventions

Level of occupational physician:

1. Two days of training followed by three intervention meetings

Level of worker:

1. Information folder with rationale
2. Homework assignments
3. Meeting with supervisor

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

1. Time to full return to work
2. Time to relapse
3. Percentage of contract hours worked
4. Work functioning

Secondary outcome measures

1. Psychological complaints
2. Work ability
3. Self efficacy in returning to work
4. Coping with work situations
5. Avoidance of work situations
6. Work adjustments
7. Satisfaction of worker with occupational physician

Overall study start date

01/01/2007

Completion date

01/01/2009

Eligibility**Key inclusion criteria**

Workers who:

1. Are two to six weeks absent from work
2. Have either:
 - a. a stress-related disorder (defined as having at least one psychological complaint with significant suffering or problems with functioning)
 - b. an anxiety disorder
 - c. a depressive disorder

Participant type(s)

Patient

Age group

Adult

Sex

Not Specified

Target number of participants

200

Key exclusion criteria

Workers with:

1. Severe mental illnesses (psychotic disorders, bipolar disorder)
2. Post Traumatic Stress Disorder (PTSD)
3. Addiction problems
4. A primary somatic disorder

Date of first enrolment

01/01/2007

Date of final enrolment

01/01/2009

Locations**Countries of recruitment**

Netherlands

Study participating centre

Academic Medical Centre

Amsterdam

Netherlands

1100 DE

Sponsor information**Organisation**

Academic Medical Centre (AMC) (The Netherlands)

Sponsor details

Coronel Institute for Occupational and Environmental Health

P.O. Box 22660

Amsterdam

Netherlands

1100 DD

Sponsor type

Hospital/treatment centre

Website

<http://www.amc.nl>

ROR

<https://ror.org/03t4gr691>

Funder(s)

Funder type

Research organisation

Funder Name

Knowledge centre on reintegration for professionals (Stichting Expertise Centrum Reintegratie [STECR]) (The Netherlands) - Aladdin Program

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	01/03/2013		Yes	No