# Collaborative Care: Depression in Occupational Care

Submission date	Recruitment status	[X] Prospectively registered		
01/12/2006	No longer recruiting	☐ Protocol		
Registration date	Overall study status	Statistical analysis plan		
01/12/2006	Completed	[X] Results		
Last Edited	Condition category	Individual participant data		
23/08/2012	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

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

# Study information

#### Scientific Title

#### Acronym

CC:DOC

#### Study objectives

A collaborative care intervention for patients with major depressive disorder who are on sick leave will result in more reduction in depressive symptoms and a faster return to work than usual care.

#### Ethics approval required

Old ethics approval format

#### Ethics approval(s)

Ethics approval received from the Ethics Committee of the VUmc (METc, VUmc, Amsterdam) on the 3rd August 2007 (ref: 2006/246).

#### Study design

Randomised controlled parallel armed trial

#### Primary study design

Interventional

# Secondary study design

Randomised controlled trial

## Study setting(s)

Not specified

# Study type(s)

Treatment

#### Participant information sheet

# Health condition(s) or problem(s) studied

Depressive disorders

#### **Interventions**

The intervention follows a collaborative care model with adherence and compliance enhancing techniques, contracting, Problem Solving Treatment (PST), antidepressant medication, a workplace intervention, and manual guided self help aimed at return to work and healthy lifestyle.

#### Intervention Type

Other

#### **Phase**

#### Primary outcome measure

The primary outcome measure is the extent of reduction in depressive symptoms, as measured by the PHO-9.

#### Secondary outcome measures

The secondary outcome measure is time to return to work, which will be acquired upon inquiry with company doctor and patient and which refers to the duration of absence through illness until work is resumed.

## Overall study start date

01/03/2007

#### Completion date

01/06/2010

# **Eligibility**

#### Key inclusion criteria

Employees on sick leave lasting between 4 to 12 weeks with major depressive disorder and who do not have the prospect of full return to work yet.

#### Participant type(s)

Patient

## Age group

Adult

#### Sex

Not Specified

#### Target number of participants

126

#### Key exclusion criteria

- 1. Patients who are suicidal, psychotic or have dementia, as noticed by the company doctor
- 2. Addicted to drugs or alcohol, as assessed by the Mini International Neuropsychiatric Interview (MINI)
- 3. Patients who do not speak Dutch sufficiently to fill in the questionnaires

Patients who are already receiving psychiatric treatment can be included in the study, in case of mutual agreement with the current care giver.

#### Date of first enrolment

01/03/2007

#### Date of final enrolment

01/06/2010

# Locations

#### Countries of recruitment

Netherlands

### Study participating centre

Trimbos-instituut/Netherlands institute of Mental Health and Addiction

Utrecht Netherlands 3500 AS

# Sponsor information

#### Organisation

Trimbos-institute/Netherlands Institute of Mental Health and Addiction (The Netherlands)

# Sponsor details

P.O. Box 725 Utrecht Netherlands 3500 AS +31 (0)30 2971100 info@trimbos.nl

#### Sponsor type

Hospital/treatment centre

#### Website

http://www.trimbos.nl/default37.html

#### **ROR**

https://ror.org/02amggm23

# Funder(s)

# Funder type

Research organisation

#### **Funder Name**

Foundation Reserves Voormalige Vrijwillige Ziekenfondsverzekering (RVVZ) (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

**Study outputs** 

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
Results article	results	01/06/2012		Yes	No