

# Gedragsactivatie door de POH-GGZ versus gebruikelijke behandeling voor depressie bij oudere volwassenen in de eerste lijn.

Gepubliceerd: 25-08-2016 Laatst bijgewerkt: 15-05-2024

The main hypothesis is that compared to TAU, BA will be more effective and less costly. A secondary goal is to explore several potential mechanisms of change, as well as predictors and moderators of treatment outcome of BA for late-life depression.

<b>Ethische beoordeling</b>	Positief advies
<b>Status</b>	Anders
<b>Type aandoening</b>	-
<b>Onderzoekstype</b>	Interventie onderzoek

## Samenvatting

### ID

NL-OMON20783

### Bron

NTR

### Verkorte titel

BeATDeP65

### Aandoening

Late-life depression, depressive symptoms, depressie, depressieve symptomen, depressieve klachten, elderly.

### Ondersteuning

**Primaire sponsor:** Pro Persona; RadboudUMC; Radboud Universiteit

**Overige ondersteuning:** ZonMw

### Onderzoeksproduct en/of interventie

## **Uitkomstmaten**

### **Primaire uitkomstmaten**

Depression severity as assessed with the Quick Inventory of Depressive Symptomatology (Q-IDS) during the 8-week treatment period and follow-up.

## **Toelichting onderzoek**

### **Achtergrond van het onderzoek**

With 12-25% prevalence, clinically significant depression is common in later life. However, the efficacy of current pharmacological and psychological treatments is limited. Behavioural programmes for late-life depression have recently received renewed attention with findings suggesting that Behavioural Activation (BA) may be effective. The primary objective of this study is to compare the effectiveness and cost-effectiveness of behavioural activation (BA) and treatment as usual (TAU) for late-life depression in primary care in the Netherlands. A cluster-randomised and controlled multicentre trial (RCT) is conducted, with two parallel groups: a) Behavioural activation, and b) Treatment as usual, conducted in primary care centres (PCC) with a follow-up of 52 weeks (FU).

### **Doele van het onderzoek**

The main hypothesis is that compared to TAU, BA will be more effective and less costly. A secondary goal is to explore several potential mechanisms of change, as well as predictors and moderators of treatment outcome of BA for late-life depression.

### **Onderzoeksopzet**

Participants in both the BA- and TAU-condition will complete these measures every two to three weeks during the 8 week therapy period, at post-treatment, and every three months during the 52-week follow-up.

### **Onderzoeksproduct en/of interventie**

In behavioural activation (BA) patients are encouraged to increase their activity levels, engage in more reinforcing and pleasurable activities, and modify avoidance and withdrawal patterns. BA is a component of cognitive-behavioural therapy (CBT), a more complex approach targeting both thoughts and behaviours.

# Contactpersonen

## Publiek

GGz Nijmegen, Outpatient Department for Anxiety Disorders, Nijmegen  
PO Box 7049,  
G.J. Hendriks  
Nijmegen 6503 GM  
The Netherlands  
+31-24-3837820

## Wetenschappelijk

GGz Nijmegen, Outpatient Department for Anxiety Disorders, Nijmegen  
PO Box 7049,  
G.J. Hendriks  
Nijmegen 6503 GM  
The Netherlands  
+31-24-3837820

## Deelname eisen

### Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

The main inclusion criterion is a PHQ-9 score >9.

### Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

Patients will be excluded from the trial in the case of I) severe mental illness in need of specialized treatment, including severe major depression, bipolar disorder, obsessive-compulsive disorder, (history of) psychosis; II) high risk of suicide, III) drug and/or alcohol abuse or dependence, IV) prior psychotherapy received in the previous 12 weeks V) current treatment by a mental health specialist. VI) moderate to severe cognitive impairment (MoCA <18).

# Onderzoeksopzet

## Opzet

Type:	Interventie onderzoek
Onderzoeksmodel:	Parallel
Toewijzing:	Gerandomiseerd
Blinding:	Open / niet geblindeerd
Controle:	Geneesmiddel

## Deelname

Nederland	
Status:	Anders
(Verwachte) startdatum:	01-07-2016
Aantal proefpersonen:	200
Type:	Onbekend

## Ethische beoordeling

Positief advies	
Datum:	25-08-2016
Soort:	Eerste indiening

## Registraties

### Opgevolgd door onderstaande (mogelijk meer actuele) registratie

ID: 47861  
Bron: ToetsingOnline  
Titel:

### Andere (mogelijk minder actuele) registraties in dit register

Geen registraties gevonden.

## In overige registers

Register	ID
NTR-new	NL5436
NTR-old	NTR6013
CCMO	NL54470.091.16
OMON	NL-OMON47861

## Resultaten

### Samenvatting resultaten

Hendriks, G. J., Oude Voshaar, R. C., Keijsers, G. P., Hoogduin, C. A. and van Balkom, A. J. (2008). Cognitive-behavioural therapy for late-life anxiety disorders: a systematic review and meta-analysis. *Acta Psychiatr Scand*, 117, 403-411.

Huibers, M. J., et al. (2014). Predicting response to cognitive therapy and interpersonal therapy, with or without antidepressant medication, for major depression: a pragmatic trial in routine practice. *J Affect Disord*, 152-154, 146-154.

Lemmens, L. H., Arntz, A., Peeters, F., Hollon, S. D., Roefs, A. and Huibers, M. J. (2015). Clinical effectiveness of cognitive therapy v. interpersonal psychotherapy for depression: results of a randomized controlled trial. *Psychol Med*, 1-16.

Licht-Strunk, E., Van Marwijk, H. W., Hoekstra, T., Twisk, J. W., De Haan, M. and Beekman, A. T. (2009). Outcome of depression in later life in primary care: longitudinal cohort study with three years' follow-up. *BMJ*, 338, a3079.