

Gepubliceerd: 18-10-2010 Laatste bijgewerkt: 13-12-2022

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Ethische beoordeling	Niet van toepassing
Status	Werving nog niet gestart
Type aandoening	-
Onderzoekstype	Interventie onderzoek

Samenvatting

Health condition

anxiety;depression;self concept;self-help

angst; depressie; zelfbeeld; zelfhulp

Ondersteuning

Primaire sponsor : prof. dr. P. Cuijpers
head of department of Clinical Psychology at the Vrije
Universiteit Amsterdam

Overige ondersteuning : The faculty of Psychology and Education, department of
Clinical psychology at the Vrije Universiteit Amsterdam.

Onderzoeksproduct en/of interventie

Uitkomstmaten

Primaire uitkomstmaten

The intervention's primary focus is enhancing self-esteem. The primary outcome measure used in this study is the Rosenberg self-esteem scale (Rosenberg, 1965). This scale includes 10 statements (such as 'generally speaking, I am content with myself'). The respondent then checks a score on a 4-point scale which indicates the extent to which the respondent identifies with the statement. Total scores vary from 0 to 30, higher scores indicate higher self esteem. In a

sample of the Dutch population, a mean score of 20.9 was reported with a standard deviation of 4.4. Both international and Dutch studies report high reliability, validity and internal homogeneity (Cronbach alpha .86; Franck et al., 2008).

Toelichting onderzoek

Achtergrond van het onderzoek

Recently, a self-help book based on cognitive behavioural techniques targeted at improving one's self image has been published (de Neef, 2010). In the current study, the effectiveness of the use of this self-help book is studied and compared to a wait list control group of people with low self esteem and subclinical anxiety and/or depressive symptoms. Guidance provided to all participants receiving the self-help book entails feedback and answers to questions regarding the assignments described in the book. Recruitment takes place among the general population in the Netherlands.

Doel van het onderzoek

To study's primary objective is to establish the effectiveness of a low-threshold guided self-help intervention targeted at improving one's self image in people with anxiety and/or depressive symptoms and low self esteem. We expect to find both a significant improvement of self-esteem (primary outcome measure) and a significant diminshment of anxiety and depression.

Onderzoeksopzet

1. Pretreatment (T0);
2. Posttreatment 10 weeks after T0(T1);
3. 12 weeks after T1 (T2).

Onderzoeksproduct en/of interventie

Respondents in the experimental condition are offered the self-help book. The intervention or

course entails 6 different techniques, based on elements from cognitive behavioural therapy. Duration of the intervention may vary from 6 to 10 weeks, depending on the pace of the individual respondent. Guidance by specially trained students and research assistants is mainly focused on motivating the respondent to work through the entire book, whilst providing feedback and information in case of any misunderstanding of the assignments described in the book.

Respondents in the control condition are placed on a waiting-list. Throughout the waiting period they do not receive the self-help book, but are free to seek help through the common channels. After the follow-up period of 3 months, they are offered the self-help book, with the same guidance provided by the university.

Contactpersonen

Publiek

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Deelname eisen

Belangrijkste voorwaarden om deel te mogen nemen (Inclusiecriteria)

1. The presence of clinically relevant anxiety and/or depressive symptoms as established by a score of 7 or

higher on the HADS-A and/or a score of 16 or higher on the CES-D;

2. The participant is suffering from low self-esteem as established by administration of the Rosenberg self-esteem

scale (Rosenberg, 1965);

3. The participant is troubled and/or limited by the aforementioned symptoms.

Belangrijkste redenen om niet deel te kunnen nemen (Exclusiecriteria)

Insufficient understanding of the Dutch language.

Onderzoeksopzet

Opzet

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

Deelname

Nederland	
Status :	Werving nog niet gestart
(Verwachte) startdatum :	15-11-2010
Aantal proefpersonen :	128
Type :	Verwachte startdatum

Ethische beoordeling

Niet van toepassing

Soort : Niet van toepassing

Registraties

In dit register bekende (historische) registraties

Geen registraties gevonden

In overige registers

Source : NTR

Register	ID
NTR-new	NL2457
NTR-old	NTR2573
CCMO	NL33798.029.10
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Resultaten

Samenvatting resultaten

N/A