Guided self-help course on the Internet for Turkish migrants with depression: a randomised controlled trial

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This study will investigate the effectiveness of a culturally adapted and internet-based guided self-help intervention among Turkish migrants, in terms of reduction of depressive symptoms (CES-D). As a secondary objective, the underlying mechanisms...

Ethical review Approved WMO

Status Recruitment stopped

Health condition type Mood disorders and disturbances NEC

Study type Interventional

Summary

ID

NL-OMON34779

Source

ToetsingOnline

Brief title

AOC-TR for Turkish migrants with depression

Condition

Mood disorders and disturbances NEC

Synonym

down, well-being

Research involving

Human

Sponsors and support

Primary sponsor: Vrije Universiteit

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

Intervention

Keyword: depression, e-health, ethnic groups, mental health

Outcome measures

Primary outcome

Depressive symptoms

Secondary outcome

Somatic symptoms, symptoms of anxiety, acculturation, satisfaction and the quality of life.

Study description

Background summary

The Turkish population living in the Netherlands has a high prevalence of psychological complaints and experiences a high threshold for seeking professional help for these problems. Seeking help through the Internet can bridge these barriers. This project aims to evaluate the effectiveness of "Alles Onder Controle TR" (in Dutch and Turkish), a web-based guided self-help intervention for depressed Turkish migrants.

Study objective

This study will investigate the effectiveness of a culturally adapted and internet-based guided self-help intervention among Turkish migrants, in terms of reduction of depressive symptoms (CES-D). As a secondary objective, the underlying mechanisms of change will be measured in terms of somatic symptoms, anxiety, acculturation, satisfaction and the quality of life.

Study design

This study is a randomized controlled trial with two conditions 1) web-based guided self-help intervention (experimental group) and 2) the wait-list control group (access to the intervention after 4 months).

Intervention

The intervention we will evaluate is the adapted version of the original website Alles Onder Controle (AOC; Allesondercontrole.nu), a brief problem solving intervention based on *self-examination* therapy (Bowman et al, 1995): Alles Onder Controle TR (in Dutch) and Her *ey Kontrol Alt*nda (in Turkish).

The course is available in two languages and both are adapted by a Turkish psychologist in collaboration with the VU University, by:

- cultural sensitivity in the languages and presentation concerning psychological problems
- use of cultural specific cases and problems that are recognizable for the target group concerned
- cultural specific examples of persons with similar problems

The intervention consists of 5 sessions and takes 5 weeks in total. During that period the respondents indicate what they think is important in their lives, they make a list of their *problems and worries* and they categorize their problems into three groups: unimportant (not related to what they think is important in their lives), important and solvable (these problems are solved by a systematic problem-solving approach consisting of 6 steps), or important but unsolvable (for example having lost someone by death, having a chronic general medical disease; for these problems they make a plan how to live with it). At the end of the course, the participant will receive a certificate for successfully completing the course.

Study burden and risks

For the study, participants will be asked to complete 3 questionnaires during 4 months. One questionnaire takes 30 minutes. The total amount for following the intervention varies per participant. Ideally, it will take on average 20 minutes each day. By participating to the study, participants don*t run any known or notable risks. Participants have always the opportunity to withdraw from this study without obligation to state their reasons and can decide by their own the amount of spent time during the study. We expect that participants with depressive complaints will benefit from the intervention if it works as expected. Overall, all participants are free to take medicine, therapy by a professional or seek for help for their complaints.

Contacts

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Trial sites

Listed location countries

Netherlands

Eligibility criteria

Age

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

Inclusion criteria

- 1) aged 18 years or older
- 2) depressive symptoms (CES-D score * 16)
- 3) Turkish ethnicity (which will be based if the participant or at least on of his/her parents is born in Turkey)
- 4) having access to a PC and the Internet and an e-mail address
- 5) provide informed consent

Exclusion criteria

Suicidal ideations or plans (M.I.N.I. score > low risk)

Study design

Design

Study type: Interventional

Intervention model: Parallel

Allocation: Randomized controlled trial

Masking: Open (masking not used)

Control: Active

Primary purpose: Prevention

Recruitment

NL

Recruitment status: Recruitment stopped

Start date (anticipated): 16-06-2010

Enrollment: 200

Type: Actual

Ethics review

Approved WMO

Date: 10-06-2010

Application type: First submission

Review commission: METC Amsterdam UMC

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 NL31548.029.10