Depression and anxiety: Indicated Prevention in homes for the elderly

Submission date Recruitment status [X] Prospectively registered 12/10/2006 No longer recruiting [X] Protocol [] Statistical analysis plan Registration date Overall study status 12/10/2006 Completed [X] Results [] Individual participant data **Last Edited** Condition category Mental and Behavioural Disorders 28/07/2014

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

Protocol serial number N/A

Study information

Scientific Title

Acronym

DIP

Study objectives

An indicated, stepped care prevention programme is feasible in homes for the elderly in Amsterdam and reduces the incidence of major disorders of depression and anxiety with 15% compared to care as usual. Total costs of health care use will be lower in the intervention group as compared to care as usual.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the local medical ethics committee

Study design

Randomised controlled trial

Primary study design

Interventional

Study type(s)

Prevention

Health condition(s) or problem(s) studied

Depressive disorders, Anxiety disorders

Interventions

A stepped care programme consisting watchful waiting, biblio-therapy, life review/Problem Solving Treatment (PST) and consultation of a mental health specialist as compared to usual care.

Intervention Type

Other

Phase

Not Specified

Primary outcome(s)

Incidence of depressive and/or anxiety disorders in residents after one year as compared to usual care.

Key secondary outcome(s))

- 1. Symptoms of depression and/or anxiety
- 2. Psychological measures such as life satisfaction and wellbeing
- 3. Satisfaction with care and treatment costs

Completion date

01/11/2010

Eligibility

Key inclusion criteria

- 1. Residents of homes for the elderly in Amsterdam
- 2. Aged 75 year and older
- 3. Scoring for symptoms of depression and anxiety on the screening measurement (Center for Epidemiologic Studies Depression scale [CES-D])
- 4. Capable of informed consent
- 5. Sufficient understanding of the Dutch language

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Senior

Sex

All

Key exclusion criteria

- 1. Residents meeting criteria for major depression and/or clinical anxiety according to the Mini-International Neuropsychiatric Interview (MINI)
- 2. Residents suffering cognitive impairment according to the Mini Mental State Examination (MMSE) are also excluded

Date of first enrolment

01/11/2006

Date of final enrolment

01/11/2010

Locations

Countries of recruitment

Netherlands

Study participating centre EMGO-Institute/VU medical centre

Amsterdam Netherlands 1000 SN

Sponsor information

Organisation

VU University Medical Center/EMGO-Institute (The Netherlands)

ROR

https://ror.org/00q6h8f30

Funder(s)

Funder type

Research organisation

Funder Name

Netherlands Organisation for Health Research and Development (ZonMw) (The Netherlands)

Alternative Name(s)

Netherlands Organisation for Health Research and Development

Funding Body Type

Private sector organisation

Funding Body Subtype

Other non-profit organizations

Location

Netherlands

Results and Publications

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	cost-effectiveness results	01/02/2014		Yes	No
Results article	results	01/02/2014		Yes	No
<u>Protocol article</u>	protocol	08/03/2007		Yes	No