# Stop worrying: the effectivity of a self-help brochure for excessive worrying, with or without telephonic support

Submission date	Recruitment status	Prospectively registered
20/12/2005	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
20/12/2005	Completed	Results
Last Edited	Condition category	☐ Individual participant data
17/09/2008	Mental and Behavioural Disorders	Record updated in last year

## 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** 01; NTR177

# Study information

Scientific Title

#### **Acronym**

Stophetgetob

## **Study objectives**

A self help brochure will help people decrease their excessive worrying and telephonic support will increase the effect.

## Ethics approval required

Old ethics approval format

## Ethics approval(s)

Ethics approval received from the local medical ethics committee

## Study design

Randomised, active controlled, crossover trial

## Primary study design

Interventional

## Study type(s)

Treatment

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

Excessive worrying, anxiety

#### **Interventions**

A self help brochure, with or without telephonic support. These exercises take one months time. In the support condition participants are contacted by telephone on a weekly basis for (minimal) support.

#### Intervention Type

Other

#### Phase

**Not Specified** 

## Primary outcome(s)

Decrease in worrying intensity.

## Key secondary outcome(s))

Decrease in anxiety, depression, sleeplessness.

## Completion date

01/10/2005

# **Eligibility**

## Key inclusion criteria

- 1. Normal population 18+ years
- 2. Self-defined excessive worrying

## Participant type(s)

**Patient** 

## Healthy volunteers allowed

No

## Age group

Adult

## Lower age limit

18 years

#### Sex

All

## Key exclusion criteria

- 1. Under the age of 18 years
- 2. Undergoing psychiatric or psychological treatment
- 3. Using antidepressants
- 4. Clinical levels of anxiety and or depression

## Date of first enrolment

01/05/2005

## Date of final enrolment

01/10/2005

## Locations

## Countries of recruitment

Netherlands

# Study participating centre

Vrije Universiteit

Amsterdam Netherlands 1081 BT

# Sponsor information

## Organisation

Vrije University Medical Centre (VUMC) (The Netherlands)

#### **ROR**

# Funder(s)

## Funder type

Not defined

## Funder Name

Not provided at time of registration

# **Results and Publications**

Individual participant data (IPD) sharing plan

## IPD sharing plan summary

Not provided at time of registration