Changing catastrophic scenarios in imagery in people with severe social anxiety

Submission date	Recruitment status	Prospectively registered
14/02/2020	No longer recruiting	☐ Protocol
Registration date	Overall study status	Statistical analysis plan
17/02/2020	Completed	Results
Last Edited	Condition category	Individual participant data
16/03/2021	Mental and Behavioural Disorders	Record updated in last year

Plain English summary of protocol

Background and study aims

Social anxiety disorder, also called social phobia, is a long-lasting and overwhelming fear of social situations. It's a common problem that usually starts during the teenage years. Social anxiety is more than shyness. It's an intense fear that does not go away and affects everyday activities, self-confidence, relationships and work or school life.

Patients with social anxiety disorder (SAD) do not only experience dysfunctional negative thoughts in anxiety-provoking social interactions but recent studies show that they also suffer from recurrent, intrusive, negative images. A study showed in most patients with social anxiety disorder these images were linked to past social traumatic memories. Recently a treatment technique 'imagery rescripting' is applied to update these traumatic memories. This resulted in a strong decrease in social anxiety complaints. However, rescripting of the intrusive, negative images itself, the future-oriented catastrophic scenarios or 'flashforwards' has not been studied yet. It is possible this is an even more direct way to treat anxiety in social situations.

The research question is whether imagery rescripting focused on flashforwards can reduce social anxiety symptoms in patients with social anxiety disorder.

Who can participate?

Patients with a diagnosis of social anxiety disorder, IQ above 80, and able to read and write in Dutch or English.

What does the study involve?

Patients with social anxiety disorder follow treatment with imagery rescripting focused on flashforwards: images of future

catastrophic scenarios of social interaction. The treatment consists of two sessions of 60 minutes per week with a total of 12 sessions. At waitlist, pre-treatment, post-treatment, 1-month follow-up and 6-month follow-up patients fill out various questionnaires to assess their symptom level. Prior to each session and twice a week during waiting-list patients fill out 2 short questionnaires on social anxiety complaints and 1 that comprised idiosyncratic social situations. Sessions were audio-recorded.

What are the possible benefits and risks of participating?

The possible benefits of participating of this study are that the treatment will reduce social

anxiety disorder complaints. There are no risks involved in participating in this study. The disadvantage of participating in this study is the time it requires to participate in the interviews and fill out the questionnaires.

Where is the study run from?

- 1. Virenze-Riagg Maastricht (Netherlands)
- 2. Vincent van Gogh Instituut (Netherlands)

When is the study starting and how long is it expected to run for? December 2013 to September 2018

Who is funding the study?
Maastricht University (Netherlands)

Who is the main contact?
Prof Marleen Rijkeboer
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Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

Clinical Trials Information System (CTIS)

Nil known

ClinicalTrials.gov (NCT)

Nil known

Protocol serial number

ECP-134; 04 11 2013

Study information

Scientific Title

Flashforward Rescripting in Social Anxiety Disorder

Acronym

FFinSAD

Study objectives

A treatment regimen focused on flashforward rescripting can substantially change social anxiety symptoms in patients with Social Anxiety Disorder (SAD)

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved 15/09/2014, Maastricht University Ethics Committee Psychology (ECP) (Ethics Review Committee Psychology and Neuroscience (ERCPN), Maastricht University, Department of the Faculty of Psychology and Neuroscience, Postbox 616, 6200 MD Maastricht, Netherlands; +31 43 3884008; no email provided), ref: ECP-134; 04 11 2013

Study design

Interventional case series

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Social anxiety disorder

Interventions

This study has a case-study design, with a randomized waitlist between 2 and 7 weeks. Patients were randomised over the number of weeks of their waitlist period. The randomization list was provided by a computerized randomization program.

Patients with social anxiety disorder (n=6) followed treatment with imagery rescripting focused on flashforwards: images of future catastrophic scenarios of social interaction. The treatment consists of two sessions of 60 minutes per week with a total of 12 sessions. At waitlist, pretreatment, post-treatment, 1-month follow-up and 6-month follow-up patients fill out various questionnaires to assess their symptom level. Prior to each session and twice a week during waiting-list patients fill out 2 short questionnaires on social anxiety complaints and 1 that comprised idiosyncratic social situations. Sessions were audio-recorded.

Flashforward Rescripting: 60 minutes sessions twice a week, with a total of 12 sessions depending on the progress of treatment. Three types of imagery exercises are provided, all focus on future catastrophic scenarios.

Intervention Type

Behavioural

Primary outcome(s)

At waitlist, pre-treatment, post-treatment, 1-month follow-up and 6-month follow-up:

- 1. The Social Interaction Anxiety Scale (SIAS)
- 2. The Social Performance Scale (SPS)
- 3. Situational anxiety severity and avoidance measured using the idiosyncratic rating scale

Key secondary outcome(s))

At waitlist, pre-treatment, post-treatment, 1-month follow-up and 6-month follow-up:

- 1. The shortened Social Phobia and Anxiety Inventory (SPAI-18)
- 2. Center for Epidemiologic Studies Depression scale (CES-D)
- 3. Self-focused Attention scale (SFA)
- 4. Social Phobic Beliefs (SPB)
- 5. Subtle Avoidance Frequency Examination (SAFE)
- 6. Remoralisation measured using the Remoralisation Questionnaire
- 7. Subjective Happiness Scale (SHS)
- 8. The avoidant personality items of the Personality Disorders Questionnaire (ADP-IV)

Completion date

01/09/2018

Eligibility

Key inclusion criteria

- 1. Diagnosis of Social Anxiety Disorder
- 2. IQ above 80
- 3. Able to read and write in Dutch or English

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Sex

Αll

Total final enrolment

6

Key exclusion criteria

- 1. Diagnosis of substance abuse disorder
- 2. Diagnosis of Psychotic Disorder
- 3. Diagnosis of Bipolar Disorder
- 4. Acute Suicidality

Date of first enrolment

16/12/2013

Date of final enrolment 01/09/2018

Locations

Countries of recruitment

Netherlands

Study participating centre Virenze-Riagg Maastricht

Parallelweg 45-47 Maastricht Netherlands 6221 BD

Study participating centre Vincent van Gogh Instituut

Stationsweg 46 Venray Netherlands 5803 AC

Sponsor information

Organisation

Maastricht University

ROR

https://ror.org/02jz4aj89

Funder(s)

Funder type

University/education

Funder Name

Universiteit Maastricht

Alternative Name(s)

Maastricht University, UM

Funding Body Type

Private sector organisation

Funding Body Subtype

Universities (academic only)

Location

Netherlands

Results and Publications

Individual participant data (IPD) sharing plan

The current data sharing plans for this study are unknown and will be available at a later date.

IPD sharing plan summary

Data sharing statement to be made available at a later date

Study outputs

Output type Details Date created Date added Peer reviewed? Patient-facing?

Participant information sheet Participant information sheet 11/11/2025 11/11/2025 No Yes