

A randomized controlled trial of a web-based treatment for social phobia

Submission date 05/03/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered
		<input type="checkbox"/> Protocol
Registration date 26/09/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan
		<input checked="" type="checkbox"/> Results
Last Edited 23/05/2019	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Individual participant data

Plain English summary of protocol
Not provided at time of registration

Study website
<http://sa.online-therapy.ch>

Contact information

Type(s)
Scientific

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Additional identifiers

EudraCT/CTIS number
Nil known

IRAS number

ClinicalTrials.gov number
Nil known

Secondary identifying numbers

100011-112345

Study information

Scientific Title

A randomized controlled trial of a web-based treatment for social phobia

Acronym

OSP Trial (Online Social Phobia Trial)

Study objectives

To determine whether a new, web-based version of cognitive therapy is an effective treatment for social phobia.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approved by the Swiss National Science Foundation on 1 April 2006.

Study design

Randomised controlled trial.

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Not Specified

Participant information sheet

Health condition(s) or problem(s) studied

Social phobia

Interventions

1. Web-based cognitive-behavioral treatment (10 weeks)

2. 10 week wait-list control condition

Patients initially allocated to wait will subsequently receive the web-based cognitive-behavioral treatment.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

The following will be performed at pre-assessment (baseline), post-assessment (at 10 weeks) and follow-up (at 6 months):

1. Social Phobia Scale
2. Social Interaction Scale
3. Liebowitz Social Anxiety Scale

Secondary outcome measures

The following will be performed at pre-assessment (baseline), post-assessment (at 10 weeks) and follow-up (at 6 months):

1. Symptom Check List (SCL-90-R)
2. Beck Depression Inventory (BDI)
3. State-Trait Anxiety Inventory (STAI)
4. Inventory of Interpersonal Problems (IIP)
5. Goal Attainment Scaling (GAS)

Overall study start date

20/03/2007

Completion date

30/06/2008

Eligibility**Key inclusion criteria**

1. Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) (American Psychiatric Association, 1994) criteria for social phobia. Assessed with the Structured Clinical Interview for Diagnostic and Statistical Manual of Mental Disorders IV (SCID IV)
2. Patients are included if the cut off score is reached on the Social Phobia Scale and Social Interaction Scale
3. Age 18 to 35
4. Willing to accept random allocation

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

50

Total final enrolment

52

Key exclusion criteria

1. Current or past psychosis
2. Borderline personality disorder (other personality disorders are not a reason for exclusion)

Date of first enrolment

20/03/2007

Date of final enrolment

30/06/2008

Locations**Countries of recruitment**

Switzerland

Study participating centre

40 Blv. du Pont d'Arve

Geneva

Switzerland

1211

Sponsor information**Organisation**

The Swiss National Science Foundation (Switzerland)

Sponsor details

Division I: Humanities and Social Sciences

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Sponsor type

Government

Website

<http://www.snf.ch>

ROR

<https://ror.org/00yjd3n13>

Funder(s)

Funder type

Government

Funder Name

The Swiss National Science Foundation (Grant ref: 100011-112345)

Results and Publications

Publication and dissemination plan

Not provided at time of registration

Intention to publish date

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	results	01/10/2009	23/05/2019	Yes	No