

Internet-based relapse prevention for Eating Disorders following inpatient treatment: randomised controlled trial for Anorexia Nervosa

Submission date

08/11/2006

Recruitment status

No longer recruiting

Registration date

04/01/2007

Overall study status

Completed

Last Edited

22/05/2019

Condition category

Mental and Behavioural Disorders

☐ Prospectively registered

☐ Protocol

☐ Statistical analysis plan

☒ Results

☐ Individual participant data

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Contact details

Klinik Roseneck
Am Roseneck 6
Priem am Chiemsee
Germany
83209

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

01GV0604

Study information

Scientific Title

Internet-based relapse prevention for Eating Disorders following inpatient treatment: randomised controlled trial for Anorexia Nervosa

Acronym

EDNET 3 AN

Study objectives

Evaluation of the efficacy of an internet based relapse prevention program in anorexia nervosa after inpatient treatment.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Approval was obtained from the ethics board of the "Bayerische Landsaerztekammer" (Bavarian General Medical Council) on August 2, 2006 (reference number: 06034).

Study design

Multi-centre, prospective, randomised superiority trial with two parallel arms each

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Not specified

Study type(s)

Treatment

Participant information sheet

Health condition(s) or problem(s) studied

Anorexia Nervosa

Interventions

Experimental intervention:

Application of a behaviourally oriented Relapse Prevention Program (RPP) via internet with chat room, SMS and e-mail support and feed-back.

Control intervention:

Treatment As Usual (TAU) and internet newsletter.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Difference between body mass index at the end of the treatment (nine months) and at randomisation

Secondary outcome measures

Explorative measures of psychopathology to investigate the time course and to support the clinical claim

Overall study start date

01/01/2007

Completion date

31/12/2009

Eligibility**Key inclusion criteria**

1. Female
2. More than 16 years
3. Diagnosis anorexia nervosa
4. Successful inpatient therapy

Participant type(s)

Patient

Age group

Adult

Sex

Female

Target number of participants

Included: 258, completers: 180

Total final enrolment

258

Key exclusion criteria

1. Mental or somatic disability prohibiting participation in study
2. Acute psychosis
3. Chronic organic or schizophrenic psychosis
4. Severe suicidal ideation/behaviour

Date of first enrolment

01/01/2007

Date of final enrolment

31/12/2009

Locations

Countries of recruitment

Germany

Study participating centre

Klinik Roseneck

Prien am Chiemsee

Germany

83209

Sponsor information

Organisation

Medical-Psychosomatic Hospital Klinik Roseneck (Medizinisch-Psychosomatische Klinik Roseneck) (Germany)

Sponsor details

Affiliated with the Ludwig-Maximilians-University (LMU) Munich

Am Roseneck 6

Prien am Chiemsee

Germany

83209

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/007ztdc30>

Funder(s)

Funder type

Government

Funder Name

German Federal Ministry for Education and Research (Bundesministerium für Bildung und Forschung) (Germany) (ref: 01 GV 0604)

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	01/03/2012	22/05/2019	Yes	No
Results article	results	30/07/2013	22/05/2019	Yes	No
Results article	results	08/02/2015	22/05/2019	Yes	No