

Effectiveness of an implementation strategy of Clinical Practice Guideline for patients with anxiety disorders in Primary Care

Submission date 18/06/2010	Recruitment status No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
Registration date 04/08/2010	Overall study status Completed	<input checked="" type="checkbox"/> Protocol
Last Edited 29/12/2020	Condition category Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<input type="checkbox"/> Individual participant data
		<input type="checkbox"/> Record updated in last year

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

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Additional identifiers

Protocol serial number
PI09/90304

Study information

Scientific Title
Cluster randomized trial for evaluate the effectiveness of an implementation strategy of a clinical practice guideline for patients with anxiety disorders

Acronym

GRITA

Study objectives

The percentage of patients with a decrease of 2 or more points in the Goldberg Anxiety Scale after 6 months, will be 20% higher in those that were involved in the focused organisational strategy over those that weren't involved in it.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The Committee of Ethical Investigation of the 9th Madrid Health Area approved on the 29th of April 2010

Study design

Multicentre cluster randomised 2 arm open label controlled trial

Primary study design

Interventional

Study type(s)

Treatment

Health condition(s) or problem(s) studied

Generalised anxiety disorder, panic disorder and panic attacks

Interventions

1. Control group:

- 1.1. The Clinical Practice Guideline (CPG) will be included on the official working place website
- 1.2. The CPG will be presented in a formative session of 2 hours of length to which will be invited to assist 1 or 2 members of each one of the 22 Health Centres involved in the study

2. Intervention group:

- 2.1. They will also receive an special formative session of 2 hours of length focused on the professionals with the protocol of intervention
- 2.2. They will receive special educational material as information sheets for patients, quick guides, etc.
- 2.3. A local opinion leader will be chosen on each team as the responsible person for the CPG
- 2.4. A defined system for reminding, audit and feedback will be developed to give back information about the intermediate results of the study at 3 and 6 months of the beginning.

Intervention Type

Other

Phase

Not Applicable

Primary outcome(s)

1. Variation of 2 or more points in the Goldberg Anxiety Scale
2. Evaluative criteria
 - 2.1. 4 or more affirmative responses in the Anxiety subscale

2.2. 2 or more affirmative responses in the Depression subscale
Outcomes will be measured at baseline, 6 and 12 months.

Key secondary outcome(s)

1. Compliance with the Guideline recommendations on treatment
2. Information and referrals to Mental Health Services

Outcomes will be measured at baseline, 6 and 12 months in data collecting notebooks

Completion date

15/09/2012

Eligibility

Key inclusion criteria

1. Patients older than 18
2. Diagnosed as generalised anxiety disorder, panic disorder and panic attacks by Diagnostic and Statistical Manual of Mental Disorders 4th edition (text revision) (DSM-IV-TR)
3. Signed informed consent

Participant type(s)

Patient

Healthy volunteers allowed

No

Age group

Adult

Lower age limit

18 years

Sex

All

Key exclusion criteria

1. Subjects that have been diagnosed of one of the following types of anxiety:
 - 1.1. Post-traumatic stress disorders
 - 1.2. Acute stress disorders
 - 1.3. drug-related anxiety disorders
 - 1.4. Anxiety disorders due to a medical or a mental illness
2. Subjects not able to read or understand Spanish language
3. Subjects that will not stay living on the Health Area in the next year after their inclusion on the study
4. Institutionalised or immobilised patients

Date of first enrolment

15/09/2010

Date of final enrolment

15/09/2012

Locations

Countries of recruitment

Spain

Study participating centre

Avda. de los Pinos, 30.

Leganés

Spain

28914

Sponsor information

Organisation

Carlos III Institute of Health (Instituto de Salud Carlos III) (Spain)

ROR

<https://ror.org/00ca2c886>

Funder(s)

Funder type

Hospital/treatment centre

Funder Name

Carlos III Health Institute (Instituto de Salud Carlos III) (Spain) - Spanish Ministry of Health Supporting Investigation Department (ref: PI09/90304)

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?
Protocol article	protocol	01/12/2011	29/12/2020	Yes	No

