ISRCTN91178137 https://doi.org/10.1186/ISRCTN91178137

A study to measure the health gain from providing GPs with training in the assessment and management of depression.

Submission date	Recruitment status	[_] Prospectively registered
23/01/2004	No longer recruiting	[_] Protocol
Registration date	**Overall study status**	[_] Statistical analysis plan
23/01/2004	Completed	[X] Results
Last Edited	**Condition category**	[_] Individual participant data
18/11/2010	Mental and Behavioural Disorders	

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
NMH11C

Study information

Scientific Title

Study objectives
Not provided at time of registration

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
GP practice

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Depression, anxiety, neuroses

Interventions
A randomised controlled trial involving 40 principals in general practice and a sample of their depressed patients. Half of the general practitioners will undergo training at the beginning of the study and the other half will receive this later. General practitioners will be randomly allocated to each group. Patients will not be randomised. Specific baseline and follow-up (3 months, 6 months and 12 months after recruitment) measures will be made on 400 patients being treated for depression by the general practitioners (i.e. approximately 10 per general practitioner).

Intervention Type
Other

Phase

Not Specified

Primary outcome measure
Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/04/2003

Completion date
30/04/2004

Eligibility

Key inclusion criteria
Consecutive GP surgery attenders fulfilling the following criteria: aged 16-65 years; GP intention to treat for depression or currently being treated for depression; duration of symptoms not more than six months.

Participant type(s)
Patient

Age group
Adult

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Those suffering from psychotic illness, and those in the recovery phase of an illness.

Date of first enrolment
01/04/2003

Date of final enrolment
30/04/2004

Locations

Countries of recruitment
England

United Kingdom

Study participating centre
National Primary Care Research and Development Centre (NPCRDC)
Manchester
United Kingdom
M13 9PL

Sponsor information

Organisation
Record Provided by the NHS R&D 'Time-Limited' National Programme Register - Department of Health (UK)

Sponsor details
The Department of Health
Richmond House
79 Whitehall
London
United Kingdom
SW1A 2NL

Sponsor type
Government

Website
http://www.doh.gov.uk

Funder(s)

Funder type
Government

Funder Name
NHS Mental Health National Research and Development Programme

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	01/01/2004		Yes	No