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PRIDE Trial: PRimary care Intervention for Depression in the Elderly

Submission date	Recruitment status	[ ] Prospectively registered
09/03/2005	No longer recruiting	[ ] Protocol
Registration date	**Overall study status**	[ ] Statistical analysis plan
26/05/2005	Completed	[X] Results
Last Edited	**Condition category**	[ ] Individual participant data
16/09/2009	Mental and Behavioural Disorders	

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
0250024

Study information

Scientific Title
A controlled trial to investigate the feasibility of a new model of intervention for the treatment of late-life depression in primary care

Acronym
PRIDE

Study objectives
The aim of the trial is to test whether the new intervention, delivered in primary care, leads to improved outcomes for elderly depressed people.

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

Study type(s)
Treatment

Health condition(s) or problem(s) studied
Late-life depression

Interventions
The intervention comprises: antidepressants (where appropriate) prescribed by the general practitioner, a brief psychological intervention delivered by a community psychiatric nurse (CPN) employed by the study, and care management according to a defined algorithm. The CPN is working closely with the patient's general practitioner but the model demands close support from and liaison with the specialist community mental health team for older people.

Control: usual care

Intervention Type
Other

Phase
Not Specified

Primary outcome(s)
SCID scores

Key secondary outcome(s))
Montgomery-asberg scores

Completion date
30/09/2005

Eligibility

Key inclusion criteria
Elderly people (>60) with depression as defined by the GP or community or practice nurse are referred into the study.
If depressed (Geriatric Depression Scale [GDS] %) they are randomised to intervention or usual care group.

Participant type(s)
Patient

Healthy volunteers allowed
No

Age group
Senior

Sex
All

Key exclusion criteria
Does not meet inclusion criteria

Date of first enrolment
01/02/2004

Date of final enrolment
30/09/2005

Locations

Countries of recruitment
United Kingdom

England

Study participating centre
Rusholme Academic Unit
Manchester
United Kingdom
M14 5NP

Sponsor information

Organisation
University of Manchester (UK)

ROR
https://ror.org/027m9bs27

Funder(s)

Funder type
Government

Funder Name
Department of Health (UK) (ref: 0250024) - the call for bids was 2002, National Service Framework for Older People

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?
Results article	results	01/06/2006		Yes	No
Results article	results	01/05/2007		Yes	No
Results article	results	04/07/2007		Yes	No