

# An exploratory randomised trial of the effectiveness of a training intervention delivered to staff of a district nursing service in improving the knowledge and detection of common mental disorders in a mixed adult and older adult caseload

<b>Submission date</b> 03/02/2003	<b>Recruitment status</b> No longer recruiting	<input type="checkbox"/> Prospectively registered
<b>Registration date</b> 03/02/2003	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 24/05/2016	<b>Condition category</b> 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**

G84/6075

## **Study information**

**Scientific Title**

An exploratory randomised trial of the effectiveness of a training intervention delivered to staff of a district nursing service in improving the knowledge and detection of common mental disorders in a mixed adult and older adult caseload

**Study objectives**

To ascertain whether a training programme delivered to district nurses and associated health workers can produce significant and sustained improvements in their abilities to detect cases of clinically significant psychological disorder in their patients.

**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)**

Screening

**Health condition(s) or problem(s) studied**

Mental and Behavioural Disorders

**Interventions**

A training package comprising seven sessions will be developed based upon evidence-based clinical practice guidelines and existing teaching resources. This will be delivered utilising varied learning approaches, predominantly interactive educational workshops such as those involving small group discussion or case studies, to a randomly allocated group of staff.

In a second phase one year later, the training intervention will be delivered to the prior control group.

**Intervention Type**

Other

**Phase**

Not Specified

**Primary outcome(s)**

Changes in measures of case recognition utilising standard properties of diagnostic instruments will be compared between intervention and control groups pre- and post-intervention, and at 12 months. Included patients will complete Hospital Anxiety Depression Scale (Zigmond & Snaith, 1983)

those over 65 will additionally complete the Short Geriatric Depression Scale (15) (Sheik & Yesavage, 1986). Staff will concurrently complete a five-point measure of presence and severity of mental disorder for their patients. Staff knowledge and attitudes will be measured at baseline and at three and 12 month intervals using an adapted version of the Depression Attitude Questionnaire (Botega, Blizard, Wilkinson, Mann, 1992), and multiple choice questions based upon World Health Organisation (WHO) primary care mental health literature.

### **Key secondary outcome(s)**

Not provided at time of registration

### **Completion date**

29/09/2005

## **Eligibility**

### **Key inclusion criteria**

1. All consenting state qualified nursing staff will be included, home care staff (levels I & II) will be randomly selected
2. Patients on their caseloads aged over 18, who provide consent, will be included

### **Participant type(s)**

Patient

### **Healthy volunteers allowed**

No

### **Age group**

Adult

### **Lower age limit**

18 years

### **Sex**

All

### **Key exclusion criteria**

For patients, diagnoses of: severe mental illness; current substance misuse disorder; moderate or severe organic brain disorder.

### **Date of first enrolment**

30/09/2002

### **Date of final enrolment**

29/09/2005

# Locations

## Countries of recruitment

United Kingdom

England

## Study participating centre

**Health Services Research Department**

London

United Kingdom

SE5 8AF

# Sponsor information

## Organisation

Medical Research Council (MRC) (UK)

# Funder(s)

## Funder type

Research council

## Funder Name

Medical Research Council (MRC) (UK)

## Alternative Name(s)

Medical Research Council (United Kingdom), UK Medical Research Council, Medical Research Committee and Advisory Council, MRC

## Funding Body Type

Government organisation

## Funding Body Subtype

National government

## Location

United Kingdom

# Results and Publications

## **Individual participant data (IPD) sharing plan**

### **IPD sharing plan summary**

Not provided at time of registration