

Improving care of people with dementia in Black African and Caribbean groups

Submission date
19/11/2015

Recruitment status
No longer recruiting

☐ Prospectively registered

☐ Protocol

Registration date
20/11/2015

Overall study status
Completed

☐ Statistical analysis plan

☒ Results

Last Edited
01/06/2018

Condition category
Mental and Behavioural Disorders

☐ Individual participant data

Plain English summary of protocol

Plain English summary under review.

Contact information

Type(s)

Public

Contact name

Mr Moise Roche

Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

20077

Study information

Scientific Title

Improving dementia care in African and Caribbean minority ethnic groups

Acronym

IDEMCARE

Study objectives

The aim of this study is to benefit patients and their family carers by increasing timely diagnosis of dementia in Black African and Caribbean groups and thus allowing choice, symptomatic treatment, reduce carers' depression, avoid crises, and delay care home admission.

Ethics approval required

Old ethics approval format

Ethics approval(s)

First Medical Research Ethics Committee, 31/03/2014, ref: 14/EE/0136

Study design

Randomised; Interventional; Design type: Process of Care

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Treatment

Participant information sheet

Not available in web format, please use the contact details below to request a patient information sheet

Health condition(s) or problem(s) studied

Topic: Primary Care; Subtopic: Dementias and neurodegeneration (DeNDRoN); Disease: All Diseases

Interventions

Leaflet, "Getting help for forgetfulness" leaflet has been developed through focus groups and interviews with members of the Black African and Caribbean ethnic minorities. It is a psychoeducation tool aimed changing attitudes to help-seeking for memory problems.

Intervention Type

Other

Primary outcome measure

APEND questionnaire; Timepoint(s): 2 weeks and 4 months

Secondary outcome measures

Not provided at time of registration

Overall study start date

16/11/2015

Completion date

30/04/2016

Eligibility**Key inclusion criteria**

Phase 1:

1. Aged 18 or over
2. Community group members who are from Black African or Caribbean Minority Ethnic groups who have given us permission to approach their members
3. Capacity to consent to participate

Phase 2:

1. Participants of phase 1 of the study who have agreed to be approached again
2. Family people with mild dementia
3. Capacity to consent to participate

Phase 3:

1. Black African or Caribbean people who are registered with a GP
2. Aged 50 years or over
3. Able to provide consent to take part in the study

Participant type(s)

Patient

Age group

Adult

Lower age limit

18 Years

Sex

Both

Target number of participants

Planned Sample Size: 94; UK Sample Size: 94

Key exclusion criteria

People who do not consent to participate or who lack capacity to consent to involvement in the study

Date of first enrolment

16/11/2015

Date of final enrolment

30/04/2016

Locations

Countries of recruitment

England

United Kingdom

Study participating centre

University College London

Division of Psychiatry

Maple House

149 Tottenham Court Road

London

United Kingdom

W1T 7NF

Sponsor information

Organisation

University College London

Sponsor details

Department of Molecular Pathology

Gower Street

London

England

United Kingdom

WC1E 6BT

Sponsor type

Hospital/treatment centre

ROR

<https://ror.org/02jx3x895>

Funder(s)

Funder type

Government

Funder Name

National Institute for Health Research

Alternative Name(s)

National Institute for Health Research, NIHR Research, NIHRresearch, NIHR - National Institute for Health Research, NIHR (The National Institute for Health and Care Research), NIHR

Funding Body Type

Government organisation

Funding Body Subtype

National government

Location

United Kingdom

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	01/08/2018		Yes	No
HRA research summary			28/06/2023	No	No