

10/66 Dementia Research Group randomised controlled trial: helping carers to care - Argentina

Submission date 20/04/2007	Recruitment status No longer recruiting	<input type="checkbox"/> Prospectively registered <input checked="" type="checkbox"/> Protocol
Registration date 11/06/2007	Overall study status Completed	<input type="checkbox"/> Statistical analysis plan <input type="checkbox"/> Results
Last Edited 06/11/2019	Condition category Mental and Behavioural Disorders	<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

Study website

<http://www.alz.co.uk/1066>

Contact information

Type(s)

Scientific

Contact name

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Contact details

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

N/A

Study information

Scientific Title

10/66 Dementia Research Group randomised controlled trial: helping carers to care - Argentina

Acronym

10/66 Dementia Caregiver Intervention

Study objectives

The primary hypothesis is that an intervention focusing upon education and training of caregivers (10/66 intervention) will be associated with a reduction in caregiver psychological strain (the 20-item Self-Reporting Questionnaire [SRQ-20] score). We further hypothesise that the intervention will be associated with an improvement in the quality of life of both caregivers and people with dementia, and in the distress experienced by caregivers arising from behavioural and psychological symptoms in the person with dementia.

Ethics approval required

Old ethics approval format

Ethics approval(s)

The study has been approved by:

1. The Institute of Psychiatry ethical committee, King's College London in April 2003 (ref: 076/03)
2. The Ethical committee of the Mental Health Research Centre of the Russian Academy of Medical Sciences in October 2003

Study design

Randomised single-blind placebo-controlled cross-over study

Primary study design

Interventional

Secondary study design

Randomised controlled trial

Study setting(s)

Other

Study type(s)

Quality of life

Participant information sheet

Health condition(s) or problem(s) studied

Dementia syndrome

Interventions

60 caregivers of people with dementia in each centre were randomised to receive the intervention immediately (n = 30) or six months later (n = 30). The 10/66 intervention targets the main carer, but includes members of the immediate and extended family. The aim is to provide basic education about dementia and specific training on managing problem behaviours. The three simple, manualised modules are delivered over five, weekly, half hour sessions.

1. Module one: assessment (one session):

- 1.1. Cognitive/functional impairment
- 1.2. Carer's knowledge and understanding of dementia
- 1.3. Care arrangements:
 - 1.3.1. Who are the family members?
 - 1.3.2. Who lives with the person with dementia?
 - 1.3.3. How do they assist the main carer?
 - 1.3.4. Which behavioural problems present most difficulties?
 - 1.3.5. How burdened do they feel?

2. Module two: basic education (two sessions):

- 2.1. General introduction to the illness
- 2.2. What to expect in the future
- 2.3. What causes/does not cause dementia?
- 2.4. Locally available care and treatment

3. Module three: training on problem behaviours (two sessions): up to eight problem behaviours identified in the assessment are addressed:

- 3.1. Personal hygiene
- 3.2. Dressing incontinence
- 3.3. Repeated questioning
- 3.4. Clinging
- 3.5. Aggression
- 3.6. Wandering
- 3.7. Apathy

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

Caregiver psychological distress (SRQ-20), which is assessed at baseline and after six months.

Secondary outcome measures

- 1. Caregiver:
 - 1.1. Zarit Burden Interview
 - 1.2. Quality of Life, measured with the World Health Organisation Quality of Life Assessment (WHO-QoL BREF)

2. Person with dementia:

2.1. Behavioural and Psychological symptoms (Neuropsychiatric Inventory Questionnaire [NPI-Q])

2.2. Quality of life, measured using the Dementia Quality Of Life instrument (DEMQOL)

All secondary outcomes will be measured at baseline and six months.

Overall study start date

01/06/2007

Completion date

01/06/2008

Eligibility

Key inclusion criteria

1. Aged 65 years and over

2. Meet Diagnostic and Statistical Manual of mental disorders fourth edition (DSM-IV) criteria for Dementia syndrome

Participant type(s)

Patient

Age group

Senior

Sex

Not Specified

Target number of participants

60 participants (30 in each arm)

Key exclusion criteria

1. Serious intercurrent illness (e.g. terminal illness) in the person with dementia, where the intervention might seem pointless to the family

2. A complete absence of family caregivers

Date of first enrolment

01/06/2007

Date of final enrolment

01/06/2008

Locations

Countries of recruitment

Argentina

England

United Kingdom

Study participating centre
Health Services and Population Research Department
London
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Sponsor information

Organisation
10/66 Dementia Research Group (UK)

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Sponsor type
Research organisation

Website
<http://www.iop.kcl.ac.uk/iopweb/departments/home/default.aspx?locator=403>

ROR
<https://ror.org/04jmzkq74>

Funder(s)

Funder type
Research organisation

Funder Name
Alzheimers Association (USA)

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?
Protocol article	Study protocol:	01/12/2007		Yes	No