

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

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| 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 10/01/2020 | 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

Contact information

Type(s)

Scientific

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Additional identifiers

Protocol serial number

N/A

Study information

Scientific Title

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 the Institute of Psychiatry ethical committee, King's College London in April 2003 (ref: 076/03).

Study design

Randomised single-blind placebo-controlled cross-over study

Primary study design

Interventional

Study type(s)

Quality of life

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(s)

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

Key secondary outcome(s)

- 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.

Completion date

25/02/2006

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

Healthy volunteers allowed

No

Age group

Senior

Sex

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

04/03/2005

Date of final enrolment

25/02/2006

Locations

Countries of recruitment

United Kingdom

England

China

Study participating centre

Health Services and Population Research Department

London

United Kingdom

SE5 8AF

Sponsor information

Organisation

10/66 Dementia Research Group (UK)

ROR

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

Funder(s)

Funder type

Research organisation

Funder Name

World Health Organization (WHO) (Switzerland)

Alternative Name(s)

, , Всемирная организация здравоохранения, Organisation mondiale de la Santé, Organización Mundial de la Salud, WHO, , ВОЗ, OMS

Funding Body Type

Government organisation

Funding Body Subtype

International organizations

Location

Switzerland

Results and Publications

Individual participant data (IPD) sharing plan**IPD sharing plan summary****Study outputs**

| Output type | Details | Date created | Date added | Peer reviewed? | Patient-facing? |
|----------------------------------|-----------------|--------------|------------|----------------|-----------------|
| Protocol article | Study protocol: | 20/07/2007 | | Yes | No |
| Study website | Study website | 11/11/2025 | 11/11/2025 | No | Yes |