Efficacy of a multicomponent support programme for caregivers of disabled persons: a randomised controlled study

Submission date Recruitment status Prospectively registered 04/05/2007 No longer recruiting [] Protocol [] Statistical analysis plan Registration date Overall study status 19/09/2007 Completed [X] Results [] Individual participant data **Last Edited** Condition category Other 11/06/2019

Plain English summary of protocol

Not provided at time of registration

Contact information

Type(s)

Scientific

Contact name

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Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers

Study information

Scientific Title

Efficacy of a multicomponent support programme for caregivers of disabled persons: a randomised controlled study

Acronym

CAREGIVER

Study objectives

A tailored support programme for caregivers of the disabled persons is effective in helping them to continue their caregiving.

Ethics approval required

Old ethics approval format

Ethics approval(s)

Ethics approval received from the Ethical Committee of Kymenlaakso Central Hospital, Kotka on the 17th December 2001 (addition 17th February 2003).

Study design

A randomised controlled trial

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

Caregiving of a disabled person

Interventions

Tailored two weeks training period in a rehabilitation centre, with tailored physical exercise, lessons about caregivers role to give psychological support and increase caregivers' knowledge and checking for medical treatments.

The control group received usual care.

The follow-up time was two years for mortality and continuation of caregiving, one year for quality of life and Zung's depression scale.

Intervention Type

Other

Phase

Not Specified

Primary outcome measure

- 1. Mortality of the care receiver
- 2. Continuity of caregiving
- 3. Placement of the care receiver into an institutional care

Primary outcomes were measured at two years.

Secondary outcome measures

- 1. Health related quality of life of the caregiver (15D Measure of Health-Related Quality of Life)
- 2. Depressive symptoms of the caregiver (Zung's depression scale)

Secondary outcomes were measured at one year.

Overall study start date

26/02/2003

Completion date

31/12/2006

Eligibility

Key inclusion criteria

- 1. 135 caregiver care receiver dyads randomly divided into an intervention group (n = 66) and a control group (n = 69)
- 2. Caregiver of a disabled adult person living in Kuusankoski and Kouvola region (south-east Finland)
- 3. Caregiving continued at least six months
- 4. Ability to leave the care receiver to temporary institutional care
- 5. Ability to live in spa-hotel conditions during the intervention

Participant type(s)

Patient

Age group

Adult

Sex

Both

Target number of participants

Total final enrolment

135

Key exclusion criteria

- 1. Serious mental or physical disease of the caregiver
- 2. Caregiving only occasional
- 3. Not able to leave the care receiver
- 4. Not able to live in a spa-hotel during the intervention

Date of first enrolment

26/02/2003

Date of final enrolment

31/12/2006

Locations

Countries of recruitment

Finland

Study participating centre University of Kuopio

Kuopio Finland 70100

Sponsor information

Organisation

The Social Insurance Institution of Finland (KELA) (Finland)

Sponsor details

c/o Dr Paavo Rissanen PO Box 450 Helsinki Finland 00101

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paavo.rissanen@kela.fi

Sponsor type

Government

Website

http://www.kela.fi/in/internet/english.nsf

ROR

https://ror.org/057yw0190

Funder(s)

Funder type

Government

Funder Name

The Social Insurance Institution of Finland (KELA) (Finland)

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	01/07/2013	11/06/2019	Yes	No