Development, execution, analysis and utilisation of a survey on long-term formal professional home care and institutional long term nursing care faculties in Europe.

No registrations found.

Ethical review	Positive opinion
Status	Recruiting
Health condition type	-
Study type	Observational non invasive

Summary

ID

NL-OMON22946

Source
NTR

Brief title
RightTimePlaceCare

Health condition

Dementia

Sponsors and support

Primary sponsor: Fundació Privada Clinic per la Recerca Biomedica, Hospital Clinic of Barcelona;
Gerontopole, University of Toulouse;
Lund University;
Maastricht University;
University of Manchester;
University of Turku;
University of Tartu;
Witten/Herdecke University;

Source(s) of monetary or material Support: European Commision, Seventh Framework Programme, (project 242153).

Intervention

Outcome measures

Primary outcome

1. People with dementia: Quality of life and quality of care;

2. Informal caregivers: Quality of life and caregiver burden.

Secondary outcome

1. People with dementia: Functional status, cognition, medication use, neuropsychiatric symptoms, comorbidity, use of services/resources;

2. Informal caregivers: Psychological wellbeing, positive and negative aspects of caregiving, experiences on quality of care.

Study description

Background summary

N/A

Study objective

N/A

Study design

Baseline assessment and follow-up after three months.

Intervention

Group 1: People with dementia who are recently admitted to an institutional nursing care facility (max. 3 months) and their informa caregivers.

Group 2: People with dementia who live at home, receive professional home care and who

are at risk for institutionalisation and their informal caregivers.

Contacts

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Eligibility criteria

Inclusion criteria

1. A formal diagnosis of dementia as diagnosed by an expert assessment (i.e. physician, psychiatrist, neurologist, geriatrician or general practitioner depending on countries' specific diagnostic procedures) and recorded in the medical record;

2. An MMSE score of 24 or below;

3. The presence of an informal caregiver who visits at least twice a month.

Exclusion criteria

1. Patients < 65 years;

2. Patients wth a primary diagnosis of Korsakoff's disease or primary psychiatric diagnosis;

3. Patients whose informal caregiver cannot be identified.

Study design

Design

Study type:	Observational non invasive
Intervention model:	Parallel
Allocation:	Non controlled trial
Masking:	Open (masking not used)
Control:	N/A , unknown

Recruitment

NL

Recruitment status:	Recruiting
Start date (anticipated):	01-11-2010
Enrollment:	2200
Type:	Anticipated

Ethics review

Positive opinion

Date:	23-12-2011
Application type:	First submission

Study registrations

Followed up by the following (possibly more current) registration

No registrations found.

Other (possibly less up-to-date) registrations in this register

No registrations found.

In other registers

Register	ID
NTR-new	NL3065
NTR-old	NTR3213
Other	European Commision, Seventh Framework Programme : 242153
ISRCTN	ISRCTN wordt niet meer aangevraagd.

Study results

Summary results
N/A