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(Cost)effectiveness of timely reactivation of psychogeriatric patients suffering from psychiatric function disorders compared to usual care towards a patient-specific intervention

Submission date	Recruitment status
01/11/2004	No longer recruiting
Registration date	**Overall study status**
09/12/2004	Completed
Last Edited	**Condition category**
08/11/2019	Mental and Behavioural Disorders

[ ] Prospectively registered
[ ] Protocol
[ ] Statistical analysis plan
[X] Results
[ ] Individual participant data

Plain English summary of protocol
Not provided at time of registration

Contact information

Type(s)
Scientific

Contact name
Mr Ton J.E.M. Bakker

Contact details
P.O. Box 4023
Schiedam
Netherlands
3102 GA
+31 (0)10 4278163
tbakker@zorggroepwaterwegnoord.nl

Additional identifiers

EudraCT/CTIS number

IRAS number

ClinicalTrials.gov number

Secondary identifying numbers
1365.0004(98-10-004)

Study information

Scientific Title
(Cost)effectiveness of timely reactivation of psychogeriatric patients suffering from psychiatric function disorders compared to usual care towards a patient-specific intervention

Study objectives
Not provided at time of registration

Ethics approval required
Old ethics approval format

Ethics approval(s)
Not provided at time of registration

Study design
Randomised controlled trial

Primary study design
Interventional

Secondary study design
Randomised controlled trial

Study setting(s)
Not specified

Study type(s)
Not Specified

Participant information sheet

Health condition(s) or problem(s) studied
Psychogeriatric patients with psychiatric function disorders.

Interventions
1. Psychogeriatric Reactivation, inpatient interdisciplinary programme, duration three months
2. Usual care as the control condition

Intervention Type
Other

Phase
Not Specified

Primary outcome measure

Not provided at time of registration

Secondary outcome measures
Not provided at time of registration

Overall study start date
01/06/2001

Completion date
01/11/2005

Eligibility

Key inclusion criteria
Psychogeriatric patients (65+) with mild to moderate cognitive impairment and functional-psychiatric symptoms.

Eligibility criteria:
1. Mini Mental State Examination: scores >18 - <27
2. Neuropsychiatric Inventory: >3 positive items
3. Barthel-index (Activities of Daily Living): scores >5 - <19
4. Diagnostic and Statistical Manual of Mental Disorders, Fourth Edition (DSM-IV) classification of dementia, delirium excluded

Participant type(s)
Patient

Age group
Not Specified

Sex
Not Specified

Target number of participants
Not provided at time of registration

Key exclusion criteria
Not provided at time of registration

Date of first enrolment
01/06/2001

Date of final enrolment
01/11/2005

Locations

Countries of recruitment
Netherlands

Study participating centre
P.O. Box 4023
Schiedam
Netherlands
3102 GA

Sponsor information

Organisation
Netherlands Organisation for Health Research and Development (ZonMw)

Sponsor details
Laan van Nieuw Oost Indië 334
P.O. Box 93245
The Hague
Netherlands
2509 AE
+31 (0)70 3495111
info@zonmw.nl

Sponsor type
Not defined

ROR
https://ror.org/01yaj9a77

Funder(s)

Funder type
Not defined

Funder Name
ZonMW, The Hague

Funder Name
Foundation Scientific Fund, Schiedam

Funder Name

De Groot Fund

Funder Name
SWBV (Foundation Support Scientific Research in Nursing Homes)

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	30/09/2013		Yes	No
Other publications	secondary analysis	01/03/2019	08/11/2019	Yes	No