

# MEMO study: Mental health in Elderly Maintained with Omega-3

<b>Submission date</b> 12/09/2005	<b>Recruitment status</b> No longer recruiting	<input checked="" type="checkbox"/> Prospectively registered
<b>Registration date</b> 12/09/2005	<b>Overall study status</b> Completed	<input type="checkbox"/> Protocol
<b>Last Edited</b> 14/02/2008	<b>Condition category</b> Mental and Behavioural Disorders	<input type="checkbox"/> Statistical analysis plan
		<input type="checkbox"/> Results
		<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

**EudraCT/CTIS number**

**IRAS number**

**ClinicalTrials.gov number**

**Secondary identifying numbers**  
2005\_05/08

# Study information

## Scientific Title

## Acronym

MEMO

## Study objectives

Counteract the process of mental deterioration in elderly people through enhancement of their EPA-DHA status

## 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)

Treatment

## Participant information sheet

## Health condition(s) or problem(s) studied

Depressive disorders, Depression, Cognitive decline

## Interventions

1. 400 mg EPA-DHA in capsules
2. 1.8 g EPA-DHA in capsules
3. Placebo oil capsules

## Intervention Type

Drug

## Phase

Not Specified

## Drug/device/biological/vaccine name(s)

EPA-DHA

**Primary outcome measure**

Cognitive function and mental well-being

**Secondary outcome measures**

Quality of life

**Overall study start date**

01/11/2005

**Completion date**

01/07/2007

## **Eligibility**

**Key inclusion criteria**

1. Men and women
2. Aged 65 years and over
3. Informed consent signed

**Participant type(s)**

Patient

**Age group**

Senior

**Sex**

Both

**Target number of participants**

300

**Key exclusion criteria**

1. A score of > 16 on the CES-D (Centre for Epidemiological Studies-Depression Scale)
2. A score of < 21 points on MMSE (Mini-Mental State Examination)
3. Current or recent (<4 weeks) use of fish oil supplements or intake of more than 4 times fish/week; 24.35 g of EPA-DHA from fish per month (800 mg/day) as judged by a fish consumption questionnaire
4. Current use of pharmacological antidepressants
5. Current use of dementia (Alzheimer) medication
6. Serious liver disease
7. Use of more than 4 glasses of alcohol per day
8. Unable to participate as judged by the responsible medical physician
9. Allergy to fish(oil)
10. Swallowing problems
11. Participation in another clinical trial less than 2 months before the start of the trial or at the same time

**Date of first enrolment**

01/11/2005

**Date of final enrolment**

01/07/2007

## Locations

**Countries of recruitment**

Netherlands

**Study participating centre**

**Division of Human Nutrition**

Wageningen

Netherlands

6700 EV

## Sponsor information

**Organisation**

Wageningen University (Netherlands)

**Sponsor details**

Division of Human Nutrition

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**Sponsor type**

University/education

**Website**

<http://www.wur.nl/UK/>

**ROR**

<https://ror.org/04qw24q55>

## Funder(s)

**Funder type**

Research council

**Funder Name**

ZON-MW, The Netherlands Organization for Health Research and Development

**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