

# Early biomarker changes in frontotemporal dementia

<https://www.neurodegenerationresearch.eu/survey/early-biomarker-changes-in-frontotemporal-dementia/>

## **Title of study**

Early biomarker changes in frontotemporal dementia

## **Acronym for cohort**

FTD-RisC

## **Name of Principal Investigator - Title**

Prof

## **Name of Principal Investigator - First name**

John

## **Name of Principal Investigator - Last name**

van Swieten

## **Address of institution -Institution**

Erasmus MC

## **Address of institution - Street address**

s Gravendijkwal 230

## **Address of institution - City**

Rotterdam

## **Address of institution - Postcode**

3015 CE

## **Country**

Netherlands

## **Website**

[www.alzheimercentrumzwn.nl](http://www.alzheimercentrumzwn.nl)

## **Contact email**

[email protected]

## Funding source

Memorabel

**Q1a. Please indicate below if your cohort includes or expects to include, incidence of the following conditions?**

Motor neurone diseases| Alzheimer's disease and other dementias

**Q2a. In a single sentence what is the stated aim of the study? (Maximum 30 words)**

Aim to find early disease-related differences and biomarkers in MRI, blood, cerebrospinal fluid and neuropsychological assessment in presymptomatic familial frontotemporal dementia

**Q2b. What distinguishes this case-control study from other studies?**

This is the first and the largest single-center longitudinal cohort study of presymptomatic familial frontotemporal dementia

**Q3a. i) Number of publications that involve use of your cohort to date**

6

**Q3a. ii) Please give up to three examples of studies to date (PI, Institution, Title of Study)**

**Q3b. If data on research outputs are already available please paste the publication list/other data or provide a link to where these data are publicly available**

Dopper et al., 2014 – Structural and functional brain connectivity in presymptomatic familial frontotemporal dementia - Neurology Rohrer et al., 2015 – Presymptomatic cognitive and neuroanatomical changes in genetic frontotemporal dementia in the Genetic Frontotemporal dementia Initiative (GENFI) study: a cross-sectional analysis – Lancet Neurology Dopper et al., 2016 – Cerebral blood flow in presymptomatic MAPT and GRN mutation carriers: A longitudinal arterial spin labeling study – NeuroImage Clinical Jiskoot et al., 2016 – Presymptomatic cognitive decline in familial frontotemporal dementia: A longitudinal study – Neurology Meeter et al., 2016 – Neurofilament light chain: a biomarker for genetic frontotemporal dementia – Annals of Clinical and Translational Neurology Meeter et al., 2016 – Progranulin levels in plasma and cerebrospinal fluid in granulin mutation carriers – Dementia and Geriatric Cognitive disorders Extra

**Q3c. If no research has been done as yet, please explain in a few sentences what information (i.e. research findings) you expect will be gained from the case-control study**

**Q4a. Study criteria: what is the age range of participants at recruitment? Age in years From:**

18

**Q4a. Study criteria: what is the age range of participants at recruitment? To:**

until death

**Q4b. Study criteria: what are the inclusion criteria?**

Subjects are 50% at risk for familial frontotemporal dementia by a first-degree relative with a known pathogenic familial FTD mutation

**Q4c. Study criteria: what are the exclusion criteria?**

Previous stroke or other neurological conditions that may affect cognitive functions

**Q5a. What is the size of the cohort (i.e. how many participants have enrolled)?**

1-1,000

**Q5b. What is the expected number of control participants?**

200-500

**Q6a. Please describe what measures are used to characterise participants**

MRI, DNA, RNA, plasma, serum, CSF, NPA, skin biopsy

**Q6b. Are there additional measures for participants with the clinical disorder?**

No

**Q6c. Are there defined primary and secondary endpoints (e.g. defined health parameters)?**

No

**If YES please specify**

**Q7. What is the study design?**

Prospective cohort

**Q8. Are your cases matched by**

Age| Sex

**Q9a. Does your study includes a specialised subset of control participants?**

Yes

**Q9b. If your study includes a specialised subset of control participants please describe**

Control participants are 50% at-risk subjects, without carrying the gene mutation

**Q10a. Is data collection for this study**

Data collection ongoing| Data analysis ongoing

**Q10b. If data collection is ongoing, are there plans to continue the cohort study beyond the current projected end date?**

Yes - intend to apply for funding

**Q11. Are data collected**

Only through study

**Q12. Is there a system in place to enable re-contact with patients for future studies?**

No

**Q13a. Please give information on data stored in a database (1)**

Data summarised in database

**% Available**

100

**Q13a. Please give information on data stored in a database (2)**

Database is web-based

**% Available**

90

**Q13a. Please give information on data stored in a database (3)**

Database on spreadsheet (e.g. excel)

**% Available**

100

**Q13a. Please give information on data stored in a database (4)**

Database on paper

**% Available**

**Q13a. Please give information on data stored in a database (5)**

No

**% Available**

**Please specify language used**

**% Available**

100

**Q13b. Please give information on how data is held as individual records**

Data is web-based

**% Available**

90

**Q14a. Are data available to other groups?**

Yes

**Q14b. If data is available to other groups what is the access policy/mechanisms for access?**

Apply to PI or co-ordinator at resource| Access through collaboration with PI only| Local/ regional access| National access| International access| Access for pilot studies permitted| Resource has own ethics approval so usually no need for separate external ethics approval

**Q15. What data sharing policy is specified as a condition of use?**

No policy exists

**Q16a. Are tissues/samples/DNA available to other groups?**

Yes

**Q16b i) If yes, please describe below**

Living donors: blood| Living donors: blood derivatives| Living donors: DNA| Living donors: cerebro-spinal fluid| Living donors: other (Skin biopsy)

**Q16b. ii) In what form are tissues/samples/DNA supplied?**

Primary Samples: Stabilised samples (frozen or fixed)| Secondary samples:(derivatives of primary samples)| Secondary samples: plasma| Secondary samples: DNA| Secondary samples: RNA

**Q16b iii) Is the access policy/mechanism for obtaining samples the same as that for obtaining data (Q14 above)?**

Yes

**Q17. Is information on biological characteristics available to other groups?**

Yes, for all the cohort

**Types:**

## Case Control Studies

### **Member States:**

Netherlands

### **Diseases:**

Alzheimer's disease & other dementias, Motor neurone diseases

### **Years:**

2016

### **Database Categories:**

N/A

### **Database Tags:**

N/A