

# Cohort of Norway

<https://www.neurodegenerationresearch.eu/survey/cohort-of-norway/>

## Title of the cohort

Cohort of Norway

## Acronym for cohort

CONOR

## Name of Principal Investigator

Title Adviser and CONOR Coordinator

First name Kjersti

Last name Andersen Nerhus

## Address of institution where award is held

Institution The Norwegian Institute of Public Health

Street Address P.O.Box 4404 Nydalen

City Oslo

Postcode NO-0403

## Country

Norway

## Website

[http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

## Contact email

[email protected]

## Funding source

The Research Council of Norway and other national and international funding sources.

## 1. The cohort includes, or expects to include, incidence of the following conditions

- Neurodegenerative disease in general

## When studies on the above condition(s) are expected to become possible

Already possible

## 2a. Stated aim of the cohort

CONOR is a collection of health data and blood samples from several Norwegian health surveys. When the data collection is complete, CONOR will be a unique database with health data and

biological samples of about 200 000 individuals. The purpose of CONOR is investigating the causes of disease.

## **2b. Features distinguishing this cohort from other population cohorts**

### **3a. i) Number of publications that involve use of cohort to date**

0

### **3a. ii) Up to three examples of studies to date (PI, Institution, Title of Study)**

### **3b. Publication list/link to where data or publications are accessible (if available)**

### **3c. Information (i.e. research findings) expected to be gained from the population cohort**

### **4a. Study criteria: age range of participants at recruitment**

Age in years from:

To ('until death' if applicable): 100

### **4b. Study criteria: inclusion criteria**

[http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

### **4c. Study criteria: exclusion criteria**

[http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

## **5. Size of the cohort (i.e. number of participants enrolled)**

More than 15,000

## **6a. Measures used to characterise participants**

[http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

## **6b. Additional measures for participants with a clinical disorder**

## **6c. Are there defined primary and secondary endpoints (e.g. defined health parameters)**

[http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

## **7. Study design**

- Cross sectional survey
- Other (please specify)
- A collection of population cohorts

## **8. Cases matched by**

- Age
- Sex
- Co-morbidities
- Cognitive function
- Physical ability
- Other health assessment (specify) / N/A
- [http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4220:1](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4220:1)

## **9a. Does the study include a specialised subset of control participants**

Yes

**9b. If yes, description of specialised subset of control participants**

CONOR is a collection of population cohorts.

**10a. i) Data collection start date**

01-03-2011

**10a. ii) Data collection end date**

**10a. iii) Data collection for this study is**

- Data collection ongoing
- Data analysis ongoing

**10b. Plans to continue the cohort study beyond the current projected end date**

**11. Data collected**

- Through links to medical records
- Through links to other records or registers (such as dental records, police records etc). Please specify
- [http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4)

**12. System in place to enable re-contact with patients for future studies**

**13a. Format and availability of data stored in a database**

Language used:

**13b. Format and availability of data held as individual records**

Language used:

**14a. Are data available to other groups**

Yes

**14b. Access policy/mechanisms for access if data are available to other groups**

- Access independent of collaboration with PI
- Access Committee mechanism
- National access
- International access
- Applicant needs to provide separate external ethics approval

**15. Data sharing policy specified as a condition of use**

No requirement to make data publicly available

**16a. Are tissues/samples/DNA available to other groups**

Yes

**16b. i) Description of available tissues/samples/DNA**

- Living donors: DNA

**16b. ii) Form available tissues/samples/DNA are supplied in**

- Primary Samples: Stabilised samples (frozen or fixed)
- Secondary samples: DNA
- Other, please specify
- [http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea\\_5811&MainArea\\_5811=5903:0:15,4](http://www.fhi.no/eway/default.aspx?pid=238&trg=MainArea_5811&MainArea_5811=5903:0:15,4)

**16b. iii) Is the access policy/mechanism for obtaining samples the same as that for obtaining data**

Yes

**17. Is information on biological characteristics available to other groups**

Yes, for all the cohort