

# The Consortium for the Early Identification of Alzheimer's Disease

<https://neurodegenerationresearch.eu/survey/the-consortium-for-the-early-identification-of-alzheimers-disease/>

## Title of study

The Consortium for the Early Identification of Alzheimer's Disease

## Acronym for cohort

CIMA-Q

## Name of Principal Investigator - Title

Prof

## Name of Principal Investigator - First name

Jean-Paul

## Name of Principal Investigator - Last name

Soucy

## Address of institution -Institution

McGill University

## Address of institution - Street address

## Address of institution - City

## Address of institution - Postcode

## Country

Canada

## Website

## Contact email

[jean-paul.soucy@mcgill.ca](mailto:jean-paul.soucy@mcgill.ca)

## Funding source

CQDM and Brain Canada (with financial support of Health Canada)

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

Alzheimer's disease and other dementias

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

To advance knowledge about Alzheimer's disease with the mission to develop tools permitting its earliest detection.

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

Direct comparison of brain and retinal amyloid content.

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

0

**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**

**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**

We hope to validate an easy-to-implement retinal scanning technique as an early test for AD detection.

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

55

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

85

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

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

**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?**

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

Clinical, cognitive, neuroimaging measures, retinal fluorescence imaging

**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)?**

**If YES please specify**

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

Cross sectional survey

**Q8. Are your cases matched by**

Cognitive function

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

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

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

Data collection ongoing

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

No

**Q11. Are data collected**

Only through the study

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

**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)**

No

**% Available**

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

No

**% Available**

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

No

**% 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**

No

**% Available**

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

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

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

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

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

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

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

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

**Types:**

Case Control Studies

**Member States:**

Canada

**Diseases:**

Alzheimer's disease & other dementias

**Years:**

2016

**Database Categories:**

N/A

**Database Tags:**

N/A