

# Cohort – Suleyman Demirel University

<https://neurodegenerationresearch.eu/survey/cohort-suleyman-demirel-university/>

## Title of the cohort

Cohort – Suleyman Demirel University

## Acronym for cohort

## Name of Principal Investigator

Title Prof

First name Serpil

Last name Demirci

## Address of institution where award is held

Institution Suleyman Demirel University

Street Address

City

Postcode

## Country

Turkey

## Website

N/A

## Contact email

srpildemirci@yahoo.com

## Funding source

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

- Alzheimer's disease and other dementias

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

#### 2a. Stated aim of the cohort

N/A

#### 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: 00

To ('until death' if applicable):

**Age in years from: 40 To ('until death' if applicable): until death**

N/A

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

N/A

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

1,000 – 5,000 participants

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

N/A

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

N/A

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

No

**7. Study design**

- Other (please specify)
- N/A

**8. Cases matched by**

- Other health assessment (specify) / N/A
- N/A

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

No

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

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

14-05-2011

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

14-05-2011

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

- At the planning stage

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

- No

**11. Data collected**

**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**

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

No policy exists

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

No

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

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

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

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

- No