

COMAJ - Young Alzheimer's Disease Cohort

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General

Identification

Detailed name Young Alzheimer's Disease Cohort

Sign or acronym COMAJ

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation

CNIL

General Aspects

Medical area Geriatrics
Neurology

Health determinants Genetic
Social and psychosocial factors

Keywords neuropathological data, neurological diseases, cognitive impairment, young population

Scientific investigator(s) (Contact)

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Organization Assistance Publique - Hôpitaux de Paris

Collaborations

Funding

Funding status Mixed

Details Alzheimer's Foundation, ANR

Governance of the database

Sponsor(s) or organisation(s) responsible Assistance Publique - Hôpitaux de Paris AP-HP

Organisation status Public

Additional contact

Main features

Type of database

Type of database Study databases

Study databases (details) Cohort study

Database recruitment is carried out by an intermediary A selection of health institutions and services

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Patients are recruited from 9 research centres (Lille, Paris, Rouen, etc.)

Database objective

Main objective To study neuropathological data in neurological disorders with cognitive impairment.

Inclusion criteria

- male or female
- adult
- having a neurological condition resulting in cognitive and/or behavioural impairment (Alzheimer's disease, Lewy body dementia, vascular dementia, etc.)
- disorder began before age 60
- has identified caregiver

Population type

Age Adulthood (45 to 64 years)

Population covered Sick population

Gender	Male Woman
Geography area	National
Detail of the geography area	Memory evaluation - CHU Rouen and 8 peripheral centres
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	10/2009
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	245
Data	
Database activity	Current data collection
Type of data collected	Clinical data Paraclinical data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Medical-social record Lumbar puncture
Paraclinical data (detail)	MRI, Single photon emission computed tomography
Presence of a biobank	Yes
Contents of biobank	Fluids (saliva, urine, amniotic fluid, ?)
Details of biobank content	Cerebrospinal fluid
Health parameters studied	Health event/morbidity Health event/mortality
Procedures	
Participant monitoring	Yes
Details on monitoring of participants	Neurologist consultation every 6 months with neuropsychological tests every 12 months.

Links to administrative sources No

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay) Contact the scientist in charge.

Access to aggregated data Access on specific project only

Access to individual data Access on specific project only