MEMENTO: Determining Factors and The Progression of The Onset of Alzheimer's Disease and Cognitive Impairment - Cohort of Patients with A Cognitive Complaint

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General

Identification

Detailed name	Cohort of Patients with A Cognitive Complaint
Sign or acronym	MEMENTO: Determining Factors and The Progression of The Onset of Alzheimer's Disease and Cognitive Impairment
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CPP: 2010-A01394-35 (15/12 2010); AFSSAPS: B101404-30 (6/12/2010).

General Aspects

Medical area	Geriatrics Neurology
Health determinants	Social and psychosocial factors
Keywords	loss of independence, cognitive decline, quality of life

Scientific investigator(s) (Contact)

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Name of the director	Dufouil

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Organization	Institut de la santé et de la recherche médicale

Collaborations

Funding

Funding status	Mixed
Details	Alzheimer's Foundation Plan (Fondation Plan Alzheimer), PHRC

Governance of the database

Sponsor(s) or organisation(s) responsible	CHU de Bordeaux
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 recruited from memory resource and research centres (CM2R) in France; regional expertise and clinical research centres for cognitive impairment.

Database objective

Main objective	To study the onset of early signs (cognitive complaints, psychometric test performances lower than normal) that may indicate Alzheimer's disease or a related illness.
Inclusion criteria	- Male or female;

	- Adult; - With a cognitive complaint.

Population type

Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years) Great age (80 years and more)
Population covered	Sick population
Gender	Male Woman
Geography area	National
Detail of the geography area	France

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	2011

Size of the database

Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	1,928 (enrolment on 21/02/14).

Data

Database activity	Current data collection
Type of data collected	Clinical data Paraclinical data Biological data
Clinical data (detail)	Direct physical measures
Details of collected clinical data	----
Paraclinical data (detail)	Neuropsychological tests, positron emission tomoscintigraphy and MRI.
Biological data (detail)	Lumbar puncture.

Presence of a biobank	Yes
Contents of biobank	Fluids (saliva, urine, amniotic fluid, ?)
Details of biobank content	Cerebrospinal fluid.
Health parameters studied	Health event/mortality Quality of life/health perception

Procedures

Data collection method	Clinical, biological, psychological, sociological and brain imaging data will be collected throughout the study.
Classifications used	----
Quality procedure(s) used	----
Participant monitoring	Yes
Monitoring procedures	Monitoring by convocation of the participant
Details on monitoring of participants	----
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	[Memento.pdf]()

Access

Terms of data access (charter for data provision, format of data, availability delay)	The cohort is a translational research platform open to sub-studies and ancillary studies submitted by any researcher in good faith. One main data set will be available 6 months after each wave of study monitoring and data cleaning. Please contact the scientist in charge of the study for access to more specific data.
Access to aggregated data	Free access
Access to individual data	Access on specific project only