ECSA - Cohort of Alcohol-Dependent Patients Admitted for Withdrawal: Suicidal Behaviour Study

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General

Identification

Detailed name	Cohort of Alcohol-Dependent Patients Admitted for Withdrawal: Suicidal Behaviour Study
Sign or acronym	ECSA
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL: 25/10/2006

General Aspects

Medical area	Psychology and psychiatry
Pathology (details)	Alcoholism, suicidal risk, depression
Health determinants	Addictions Lifestyle and behavior Social and psychosocial factors
Keywords	suicide attempt, healthcare users, health events, dependency

Scientific investigator(s) (Contact)

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Unit	Unit 675 Psychiatry and Neuroscience Centre, Sainte Anne Hospital
Organization	Inserm - National Institute of Health and Medical

	Research
Collaborations	
Funding	
Funding status	Public
Details	PHRC (hospital clinical research programme); clinical research delegation (DRC)
Governance of the database	
Sponsor(s) or organisation(s) responsible	Inserm - National Institute of Health and Medical Research
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.	Inclusion method: Prospective Inclusion cut-off date: 01/08/2009 Number of required subjects: [500-1000] Number details: 800
Database objective	
Main objective	General aim: To measure the impact of suicidal thoughts, behaviour and death in alcohol dependency. Secondary aim: To assess clinical (psychiatric comorbidity), neuropsychological (computer IGT test) and genetic (blood sample) risk factors.
Inclusion criteria	Alcohol-dependent, admitted for withdrawal.
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	Regional
French regions covered by the database	Île-de-France
Detail of the geography area	Multicentric cohort throughout France (5 centres)

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	05/2007
Date of last collection (YYYY or MM/YYYY)	09/2011

Size of the database

Size of the database (number of individuals)	[500-1000[individuals
Details of the number of individuals	750

Data

Database activity	Current data collection
Type of data collected	Clinical data Declarative data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Clinical examination: At baseline and during follow-up. Examination frequency: 1 (year). Information collected during clinical examination: Depression, dependence, etc. (all psychiatric disorders).

Declarative data (detail)	Face to face interview
Details of collected declarative data	Interview questionnaire: At baseline and during follow-up. Interview frequency: 2 to 3 (years). Information collected during interview: Diagnostic interview for genetic studies (DIGS): semi-structured questionnaire. Other information sheet: During study. Frequency of other information sheet: ONCE AT ANALYSIS COMPLETION (Yearly). Information collected through other information sheet: Vital status. Who completes the other information sheet? Friends or relatives and/or general practitioner.
Biological data (detail)	Type of samples collected: Blood (for DNA)
Presence of a biobank	Yes
Contents of biobank	Plasma DNA
Details of biobank content	Biobank: Blood bank, DNA bank

Procedures

Quality procedure(s) used	Interviews: online entry; clinical examinations: online entry; biological examinations: online entry; consistency request: after electronic data is entered; management of missing data: refer back to patient; physician reminder for follow-up visits? yes; subject reminder for follow-up visits? yes; patients are informed about the use of their data (written).
Participant monitoring	Yes
Details on monitoring of participants	2 years
Links to administrative sources	No

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	Can data be used by academic teams? No. Can data be used by industrial teams? No.
Access to aggregated data	Access not yet planned

Access to individual data	No access