SAH - Treatment for Subarachnoid Haemorrhage from Ruptured Aneurysm: National Cohort Study

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Last update : 07/30/2014 | Version : 1 | ID : 60057

General

Identification

Detailed name	Treatment for Subarachnoid Haemorrhage from Ruptured Aneurysm: National Cohort Study
Sign or acronym	SAH
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL : 10/01/2008

General Aspects

Medical area	Neurology
Others (details)	Subarachnoid haemorrhage
Keywords	Neurological moribidity, death, hospitalisation

Scientific investigator(s) (Contact)

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Organization	CHU de
Name of the director	AUDIBERT
Unit	CENTRE D'EPIDEMIOLOGIE CLINIQUE CIC-EC INSERM CIE 6, NANCY

Name of the director	PUYBASSET
Collaborations	
Funding	
Funding status	Mixed
Details	CHU DE NANCY, ANARFL
Governance of the database	
Sponsor(s) or organisation(s) responsible	CHU de Nancy
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: 01/10/2009
Database objective	
Main objective	Objective general: to identify prognostic factors of mortality and neurological morbidity in patients suffering a subarachnoid haemorrhage (SAH) from an aneurysmal rupture who arrive alive at hospital. - to describe characteristics, medical treatment and neurologic outcome for these subjects; - to define poor prognostic factors.
Inclusion criteria	Adult patients hospitalised for a subarachnoid haemorrhage from an aneurysmal rupture.
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	Multicentric cohort (30 centres): France (including overseas territories)

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	10/2007
Date of last collection (YYYY or MM/YYYY)	10/2010

Size of the database

Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	2100

Data

Database activity	Data collection completed
Type of data collected	Clinical data
Clinical data (detail)	Direct physical measures Medical registration
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Cinical examinations: manual input

Participant monitoring	Yes
Details on monitoring of participants	Follow-up duration: 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)	Data may not be used by academic teams Data may not be used by industrial teams
Access to aggregated data	Access on specific project only
Access to individual data	No access