ENDOCARDITIS - Repeated cross-sectional study for patients with infective endocarditis: surgical impact

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General

Identification

Detailed name	Repeated cross-sectional study for patients with infective endocarditis: surgical impact
Sign or acronym	ENDOCARDITIS
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL: 01/01/2009

General Aspects

Medical area	Cardiology Infectious diseases
Keywords	Recurrence, valve surgery, hospitalisation, death

Scientific investigator(s) (Contact)

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Collaborations

Participation in projects, networks and consortia	Yes
Details	Involvement in a cohort: ICE

Funding

Funding status	Public
Details	PHRC

Governance of the database

Sponsor(s) or organisation(s) responsible	CHU Nancy
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Repeated cross-sectional studies (except case control studies)
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.	Prospective
Database objective	
Main objective	To assess the impact of surgical strategy on medium to long-term morbidity and mortality. Secondary objective: descriptive and analytic.
Inclusion criteria	Patients over 18 years of age treated in hospital for definite (Duke) infective endocarditis (IE) in 2008.
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years)
Population covered	Sick population
Gender	Male Woman
Geography area	National
Detail of the geography area	100 centres: Rhône-Alpes, Paris and immediate suburbs, Lorraine, Franche-Comté, Marne, Languedoc-Roussillon, Ille-et-Vilaine.
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	1991
Date of last collection (YYYY or MM/YYYY)	2008
Size of the database	
Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	2008: 497 patients
Data	
Database activity	Data collection completed
Type of data collected	Declarative data

	Paraclinical data Biological data
Declarative data (detail)	Face to face interview
Details of collected declarative data	Information sheet at baseline and during follow-up supplied by the physicians.
Paraclinical data (detail)	Imaging
Biological data (detail)	Micro-organisms
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption
Procedures	
Data collection method	Clinical exams: manual input. Biological analysis: manual input.
Quality procedure(s) used	Request for consistency when data is processed electronically and after data is processed electronically. Missing data is managed by returning to source file or third party. Physicians contacted again for follow-up visits. Internal quality audit performed annually. SOP (Standard Operating Procedure) quality procedure. Patients are informed of the use of their data in writing.
Participant monitoring	No
Links to administrative sources	Yes
Linked administrative sources (detail)	CépiDc
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Data may be used by academic teams. To be decided if data may be used by industrial teams.

Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only