

CARDIORENAL - Longitudinal observational study on cardiovascular and renal risk in patients with hypertension and at high CV risk

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General

Identification

Detailed name Longitudinal observational study on cardiovascular and renal risk in patients with hypertension and at high CV risk

Sign or acronym CARDIORENAL

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL N°903103

General Aspects

Medical area Cardiology
Neurology
Urology, andrology and nephrology

Health determinants Genetic
Social and psychosocial factors

Others (details) Myocardial infarction, stroke, cardiovascular death

Keywords High-risk patients, cardiovascular event, monitoring

Scientific investigator(s) (Contact)

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Collaborations

Funding

Funding status Private

Details Bristol-Myers Squibb France / Sanofi-Aventis France

Governance of the database

Sponsor(s) or organisation(s) responsible Bristol-Myers Squibb France (BMS)

Organisation status Private

Sponsor(s) or organisation(s) responsible Sanofi Aventis

Organisation status Private

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. Random sampling in clusters

Database objective

Main objective	Evaluate, over a cohort of patients with a high cardiovascular risk according to the ANAES definition and monitored by a general practitioner, the incidence of the major cardiovascular event (MI, CVA and cardiovascular death) according to the various sub-classes of risks defined by ANAES in the "high risk" level
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Inclusion criteria	M/F ≥ 18 years, examined in ambulatory consultation by general practitioners and cardiologists, having a treated essential HBP, or such that SBP/DBP $> 140/90$ mmHg in the absence of treatment, monitored in a normal manner by the investigating doctor
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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)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	National
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Detail of the geography area	Metropolitan France
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Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	2003
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Date of last collection (YYYY or MM/YYYY)	2007
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Size of the database

Size of the database (number of individuals)	[10 000-20 000[individuals
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Details of the number of individuals	16600
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Data

Database activity	Data collection completed
Type of data collected	Clinical data
Clinical data (detail)	Direct physical measures
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	Paper observation notebook
Participant monitoring	Yes
Details on monitoring of participants	Y1, Y2, Y3
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Publications
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only