

EOLE - Observational study of Long-term follow-up of post-myocardial infarction

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General	
Identification	
Detailed name	Observational study of Long-term follow-up of post-myocardial infarction
Sign or acronym	EOLE
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CCTIRS 05.484, CNIL 906042
General Aspects	
Medical area	Cardiology
Health determinants	Lifestyle and behavior Medicine Nutrition Occupation
Keywords	Acute Myocardial Infarction (AMI), lifestyle and dietary recommendations, cardiovascular treatments, secondary prevention, mortality, pharmacoepidemiology, cohort, Department of Pharmacology, Bordeaux
Scientific investigator(s) (Contact)	
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Organization	Université Bordeaux

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Unit	Service de Pharmacologie, CIC-P 0005-INSERM U657- Université Bordeaux Segalen
Organization	Université Bordeaux

Collaborations

Participation in projects, networks and consortia	No
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Funding

Funding status	Private
Details	Pierre Fabre Médicament (unconditional support)

Governance of the database

Sponsor(s) or organisation(s) responsible	Service de Pharmacologie, CIC-P 0005-INSERM U657- Université Bordeaux Segalen
Organisation status	Public
Presence of scientific or steering committees	Yes

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 care professionals
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Patients are included by cardiologists who are themselves selected from a random sample of hospital and non-hospital cardiologists in metropolitan France.
Database objective	
Main objective	The main objective is to assess in real situations of use the impact of cardiovascular medications and recommended lifestyle and dietary recommendations on all-cause mortality in secondary prevention of myocardial infarction.
Inclusion criteria	Patient with recent (?3 months) acute myocardial infarction (first or recurrent heart attack); Patient seen during the first or second consultation after acute myocardial infarction; Infarct defined by the presence at least of two among the following criteria: symptomatic (characteristic pain), electric (Q wave and / or ST elevation in at least two adjacent leads), enzymatic (elevated CPKMB and / or troponin greater than 2 twice the normal value); Patient who may be followed for 6 years; Patient without a non-cardiovascular disease that is life-threatening in the short term (?3 months); Patient not included in a clinical trial of a non marketed drug; Patient not affected by a language barrier (unable to read the information sheet or complete the self-administered questionnaires); Patient agreeing to participate.
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

Hospital and non-hospital cardiologists in metropolitan France

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)

2006

Date of last collection (YYYY or MM/YYYY)

2015

Size of the database

Size of the database (number of individuals)

[1000-10 000[individuals

Details of the number of individuals

5538

Data

Database activity

Data collection completed

Type of data collected

Clinical data
Declarative data
Administrative data

Clinical data (detail)

Direct physical measures

Declarative data (detail)

Paper self-questionnaire
Internet self-questionnaire
Phone interview

Details of collected declarative data

Questionnaire diet, physical activity before and after myocardial infarction, tobacco use, drugs taken, and hospitalisations after myocardial infarction of inclusion

Administrative data (detail)

Patient name , first name, date and place of birth, phone number and address, contact details of the general practitioner.

Presence of a biobank

No

Health parameters studied

Health event/morbidity
Health event/mortality
Health care consumption and services
Quality of life/health perception

Care consumption (detail)

Hospitalization
Medicines consumption

Procedures

Data collection method	Inclusion data are collected by cardiologists and patients via a paper questionnaire. Follow-up data are collected by cardiologists (only at 6 and 24 months of follow up in the first inclusion period) and by patients via an eCRF or a paper questionnaire.
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Participant monitoring	Yes
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Monitoring procedures	Monitoring by contact with the participant (mail, e-mail, telephone etc.)
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Details on monitoring of participants	The total follow-up is 6 years with an evaluation at 6 months, 24 months, 3 years 4 years 5 years 6 years via a self-administered questionnaire. Patients may fill-out these questionnaires via the internet (eCRF) or by postal mail using a paper questionnaire. The data collected in this questionnaire concern dietary habits since the AMI (only at 6 and 24 months of follow-up for patients included during the first inclusion period), treatments taken, physical activity, smoking, current health status, and hospitalizations. Follow-up by cardiologists concerns only patients included in the first inclusion period with an evaluation at 6 and 24 months (vital status, cardiovascular events, clinical and laboratory data, prescribed cardiovascular treatments). Vital status determination of included patients at 3.5 years and 6 years will be performed using the INSERM / INSEE procedure.
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Links to administrative sources	Yes
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Linked administrative sources (detail)	RNIPP, CéPIDC
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Promotion and access

Promotion

Link to the document	http://www.hal.inserm.fr/EOLE/
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Description	List of publications in HAL
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Confidential study reports are submitted to the Pharmaceutical company and health authorities. The study reports and scientific communications (posters, articles, ...) are validated by the study
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Scientific Committee. Ownership of study data is the subject of an agreement between the University Bordeaux Segalen and the pharmaceutical company. Terms for third-party access to the database are to be defined.

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only