

COPART - Longitudinal Study on Patients Hospitalised for Artery Disease

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General

Identification

Detailed name Longitudinal Study on Patients Hospitalised for Artery Disease

Sign or acronym COPART

General Aspects

Medical area Cardiology

Health determinants Genetic

Keywords systolic pressure index, coronary and cerebrovascular events, hospitalisation

Scientific investigator(s) (Contact)

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Unit U1048 Institut des maladies métaboliques et cardiovasculaires (I2MC)Équipe/activité : Production et fonction plaquettaire, signalisation et phosphoinositides

Organization CHU Toulouse ;

Collaborations

Funding

Funding status Public

Details	CHU
Governance of the database	
Sponsor(s) or organisation(s) responsible	CHU Toulouse
Organisation status	Public
Sponsor(s) or organisation(s) responsible	Institut national de la santé et de la recherche médicale - Inserm
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Longitudinal study (except cohorts)
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.	All patients with peripheral artery disease that were hospitalised in the vascular medicine department of the Rangueil university hospital between 01/06/2004 and 31/07/2006 were consecutively included.
Database objective	
Main objective	To form a database on peripheral artery disease of the lower limbs in hospitalised patients that is atheromatous in origin, which shall allow the treatment, monitoring and prognosis of affected patients to be assessed. This study aims to compare prescribed treatment upon discharge from hospital and French National Health Authority recommendations and to evaluate the trends in prescribed treatment over time.
Inclusion criteria	<ul style="list-style-type: none"> - male and female - adult - patients with peripheral artery disease hospitalised

in the vascular medicine department
Patients that died during hospitalisation were excluded from the study.

Population type

Age
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
Local

French regions covered by the database
Aquitaine Limousin Poitou-Charentes
Languedoc-Roussillon Midi-Pyrénées

Detail of the geography area
Toulouse

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)
06/2004

Date of last collection (YYYY or MM/YYYY)
06/2005

Size of the database

Size of the database (number of individuals)
< 500 individuals

Details of the number of individuals
400

Data

Database activity
Current data collection

Type of data collected
Declarative data
Paraclinical data
Administrative data

Declarative data (detail)
Paper self-questionnaire

Paraclinical data (detail)
Restenosis or bypass obstruction diagnosis will be by ultrasound or imaging.

Administrative data (detail)	Information on the patient's vital status will be requested from the town/city hall where the patient was born or residing. This is applicable for all patients.
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Presence of a biobank	No
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Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services
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Care consumption (detail)	Hospitalization Medical/paramedical consultation
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Procedures

Participant monitoring	Yes
Details on monitoring of participants	Follow-up is annual. Each patient should have information regarding their outcome at one year. Events that occurred during follow-up will be listed

Links to administrative sources	No
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Promotion and access

Promotion

Link to the document	http://www.portailvasculaire.fr/espace-sfmv/etude-copart
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Link to the document	http://tinyurl.com/HaI-COPART
Description	List of publications in HAL

Link to the document	http://www.ncbi.nlm.nih.gov/pubmed/?term=copart
Description	List of publications in Pubmed

Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge.
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Access to aggregated data	Access on specific project only
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Access to individual data	Access on specific project only
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