

APTOR - Prospective observational study of patients with acute coronary syndrome managed by PCI and antiplatelet treatment: Antiplatelet Treatment Observational Registry

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General	
Identification	
Detailed name	Prospective observational study of patients with acute coronary syndrome managed by PCI and antiplatelet treatment: Antiplatelet Treatment Observational Registry
Sign or acronym	APTOR
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL N°906309
General Aspects	
Medical area	Cardiology
Keywords	Resource use, acute coronary syndrome, percutaneous angioplasty, antiplatelet treatment
Scientific investigator(s) (Contact)	
Name of the director	Médecin pharmacoépidémiologiste
Email	PHARMACOEPI_FRMAIL@LILLY.COM
Organization	Eli Lilly
Collaborations	
Funding	
Funding status	Private
Details	Eli Lilly and Company
Governance of the database	
Sponsor(s) or organisation(s)	Eli Lilly

responsible	
Organisation status	Private
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 is made on the basis of:	Another treatment or procedure
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Patients recruited by cardiologists in interventional cardiology centers
Database objective	
Main objective	Primary objective: evaluate the use of healthcare resources and cost in the 12 months following a percutaneous coronary intervention (PCI) combined with antiplatelet treatment in patients with acute coronary syndrome. Secondary objectives: quality of life and clinical evolution.
Inclusion criteria	Be diagnosed with acute coronary syndrome, - treatment with percutaneous coronary intervention in routine care; - start or continue antiplatelet treatment.
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 International

Detail of the geography area France, Spain and Great Britain

Data collection

Dates

Date of first collection (YYYY or MM/YYYY) 2007

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 1525

Data

Database activity Data collection completed

Type of data collected Clinical data
Declarative data

Clinical data (detail) Direct physical measures
Medical registration

Declarative data (detail) Paper self-questionnaire

Presence of a biobank No

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

Care consumption (detail) Hospitalization
Medical/paramedical consultation
Medicines consumption

Procedures

Data collection method Study data collection form

Participant monitoring Yes

Details on monitoring of participants	1 year monitoring
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Links to administrative sources	No
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Promotion and access

Promotion

Link to the document	http://tinyurl.com/Pubmed-APTOR
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Description	List of publications in Pubmed
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Report, poster and publication
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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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