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

Head : Médecin pharmacoépidémiologiste

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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	Patients recruited by cardiologists in interventional
sample selection.	cardiology centers
Database objective	cardiology centers
	Primary objective: evaluate the use of healthcare resources and cost in the 12 months following a
Database objective	Primary objective: evaluate the use of healthcare resources and cost in the 12 months following a percutaneous coronary intervention (PCI) combine with antiplatelet treatment in patients with acute coronary syndrome. Secondary objectives: quality of life and clinical
Database objective Main objective	<ul> <li>Primary objective: evaluate the use of healthcare resources and cost in the 12 months following a percutaneous coronary intervention (PCI) combine with antiplatelet treatment in patients with acute coronary syndrome.</li> <li>Secondary objectives: quality of life and clinical evolution.</li> <li>Be diagnosed with acute coronary syndrome, - treatment with percutaneous coronary intervention in routine care;</li> </ul>
Database objective Main objective Inclusion criteria	<ul> <li>Primary objective: evaluate the use of healthcare resources and cost in the 12 months following a percutaneous coronary intervention (PCI) combine with antiplatelet treatment in patients with acute coronary syndrome.</li> <li>Secondary objectives: quality of life and clinical evolution.</li> <li>Be diagnosed with acute coronary syndrome, - treatment with percutaneous coronary intervention in routine care;</li> </ul>
Database objective Main objective Inclusion criteria	<ul> <li>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.</li> <li>Secondary objectives: quality of life and clinical evolution.</li> <li>Be diagnosed with acute coronary syndrome, - treatment with percutaneous coronary intervention in routine care;</li> <li>start or continue antiplatelet treatment.</li> </ul> Adulthood (19 to 24 years) Adulthood (45 to 64 years) Elderly (65 to 79 years)

	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
Two of data collected	
Type of data collected	Clinical data Declarative data
Clinical data (detail)	
	Declarative data Direct physical measures
Clinical data (detail)	Declarative data Direct physical measures Medical registration
Clinical data (detail) Declarative data (detail)	Declarative data Direct physical measures Medical registration Paper self-questionnaire
Clinical data (detail) Declarative data (detail) Presence of a biobank	Declarative data Direct physical measures Medical registration Paper self-questionnaire No Health event/morbidity Health care consumption and services
Clinical data (detail) Declarative data (detail) Presence of a biobank Health parameters studied	Declarative data Direct physical measures Medical registration Paper self-questionnaire No Health event/morbidity Health care consumption and services Quality of life/health perception Hospitalization Medical/paramedical consultation
Clinical data (detail) Declarative data (detail) Presence of a biobank Health parameters studied Care consumption (detail)	Declarative data Direct physical measures Medical registration Paper self-questionnaire No Health event/morbidity Health care consumption and services Quality of life/health perception Hospitalization Medical/paramedical consultation

Details on monitoring of participants	1 year monitoring
Links to administrative sources	No
Promotion and access	
Promotion	
Link to the document	http://tinyurl.com/Pubmed-APTOR
Description	List of publications in Pubmed
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Report, poster and publication
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only