

ARCHIMED HOPITAL - Prevention of venous thromboembolic events with Arixtra 2.5mg for medical ill patients in hospital

Head :Leclerc-Zwirn Christel, Laboratoire GSK

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General

Identification

Detailed name Prevention of venous thromboembolic events with Arixtra 2.5mg for medical ill patients in hospital

Sign or acronym ARCHIMED HOPITAL

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL : 906317

General Aspects

Medical area Hematology

Health determinants Medicine

Keywords pharmaco-epidemiology, thromboprophylaxis, fondaparinux, arixtra

Scientific investigator(s) (Contact)

Name of the director Leclerc-Zwirn

Surname Christel

Phone +33 (0)1 39 17 86 96

Email christel.c.leclerc-zwirn@gsk.com

Unit Laboratoire GSK

Collaborations

Funding

Funding status Private

Details GSK laboratory

Governance of the database	
Sponsor(s) or organisation(s) responsible	LABORATOIRE GSK
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 made on the basis of:	Medication(s) taken
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	<p>All public and private hospitals that purchase ARIXTRA® 2.5 mg in metropolitan France will be solicited to participate in this study (source: database from the GlaxoSmithKline laboratory). Investigators will be the pharmacists of the hospitals that have accepted to participate in the study. Using the registry kept, the pharmacist will include the first 10 patients responding to the following criteria:</p> <ul style="list-style-type: none"> ? Patient of at least 18 years of age, ? hospitalized in one of the establishment's care departments (excluding surgery departments), for whom a treatment via ARIXTRA® 2.5 mg is initiated with thromboprophylaxis
Database objective	
Main objective	The main objective of this study is to evaluate the average duration of treatment via ARIXTRA® 2.5 mg at the hospital within the framework of the thromboprophylaxis indication
Inclusion criteria	<p>Using the registry kept, the pharmacist will include the first 10 patients responding to the following criteria:</p> <ul style="list-style-type: none"> ? Patient of at least 18 years of age,

? hospitalized in one of the establishment's care departments (excluding surgery departments),
 ? for whom a treatment via ARIXTRA® 2.5 mg is initiated with thromboprophylaxis

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	France
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Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	2009
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Date of last collection (YYYY or MM/YYYY)	2011
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Size of the database

Size of the database (number of individuals)	[500-1000[individuals
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Details of the number of individuals	680
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Data

Database activity	Data collection completed
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Type of data collected	Clinical data Declarative data
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Clinical data (detail)	Medical registration
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Declarative data (detail)	Face to face interview
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Presence of a biobank	No
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Health parameters studied	Health event/morbidity Health care consumption and services
Care consumption (detail)	Hospitalization Medicines consumption
Procedures	
Data collection method	During the first month following the set up of the study in the center, the pharmacist will log in a registry all of the prescription requests for ARIXTRA® 2.5 mg in thromboprophylaxis carried out by his establishment's care departments (excluding surgery departments). The pharmacist will identify the patients that should be included in the study according to the defined eligibility criteria. He will complete a specific inclusion questionnaire for the first 10 patients that meet the eligibility criteria. The pharmacist will complete for each patient included a follow-up questionnaire when released from the hospital.
Participant monitoring	Yes
Details on monitoring of participants	The data that will be collected in the follow-up questionnaire are: ? Premature interruption of the treatment, ? total duration of the administration of the treatment, ? prescriptions for prophylactic purposes after the treatment
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Publications in progress
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only