

WP6 negative control study: The risk of myocardial infarction not associated with the use of antibiotics: A study using a US database

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Study

Finalised

Administrative details

EU PAS number

EUPAS3025

Study ID

28276

DARWIN EU® study

No

Study countries

 United States

Study status

Finalised

Research institutions and networks

Institutions

Sanofi


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Institution

Networks

PROTECT


 Belgium


 Denmark

 France


 Germany

 Italy


 Netherlands

 Poland

 Spain

 Sweden

 Switzerland

 United Kingdom

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Network

Contact details

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Primary lead investigator

Stéphanie Tcherny-Lessenot

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 02/02/2012

Actual: 02/02/2012

Study start date

Planned: 01/11/2012

Actual: 15/11/2012

Data analysis start date

Planned: 01/11/2012

Actual: 15/11/2012

Date of final study report

Planned: 31/12/2013

Actual: 21/07/2014

Sources of funding

- Pharmaceutical company and other private sector

More details on funding

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Study protocol

[PROTECT_WP6_protocol_Antibiotics_MI_negative control_invision datamart_final_02022012.pdf](#) (137.95 KB)

Regulatory

Was the study required by a regulatory body?

No

Is the study required by a Risk Management Plan (RMP)?

Not applicable

Methodological aspects

Study type

Study type list

Study topic:

Disease /health condition

Human medicinal product

Study type:

Non-interventional study

Scope of the study:

Assessment of risk minimisation measure implementation or effectiveness

Other

If 'other', further details on the scope of the study

Negative control

Data collection methods:

Secondary use of data

Main study objective:

to assess the absence of association between antibiotics use and myocardial infarction by replicating a nested case-control design in a US claims database (LabRx)

Study Design

Non-interventional study design

Case-control

Study drug and medical condition

Anatomical Therapeutic Chemical (ATC) code

(J01) ANTIBACTERIALS FOR SYSTEMIC USE

Medical condition to be studied

Myocardial infarction

Population studied

Short description of the study population

Patients of all ages with an active or died registration status during the study period of January 1st, 2004 to December 31st, 2009 in the Invision Data Mart. Patients must have attained one year of enrolment in the database at the beginning of the study period.

Age groups

- Adults (18 to < 46 years)
 - Adults (46 to < 65 years)
 - Adults (65 to < 75 years)
 - Adults (75 to < 85 years)
 - Adults (85 years and over)
-

Estimated number of subjects

80000

Study design details

Outcomes

To estimate the risk of myocardial infarction associated with antibiotics exposure (users and non-users) To estimate the risk of myocardial infarction

associated with various antibiotics classes To estimate the risk of myocardial infarction associated with specific individual antibiotics To assess the effect of dose and duration of use for specific individual antibiotics, To replicate the analysis using a population-based case-control design

Data analysis plan

We will compute odds ratios (OR) and 95% confidence intervals of first occurrence of acute myocardial infarction associated with current use of antibiotics (as a group and different classes and individual drugs when possible) as compared to non-use with conditional logistic regression. Age, sex, calendar year, and other variables will be introduced in the model to control for potential confounding. Also, dose and duration-relationships will be examined.

Documents

Study results

[PROTECT WP6 ATB AMI REPORT 21072014_final.pdf](#) (217.03 KB)

Data management

ENCePP Seal

The use of the ENCePP Seal has been discontinued since February 2025. The ENCePP Seal fields are retained in the display mode for transparency but are no longer maintained.

Data sources

Data sources (types)

Administrative healthcare records (e.g., claims)

Use of a Common Data Model (CDM)

CDM mapping

No

Data quality specifications

Check conformance

Unknown

Check completeness

Unknown

Check stability

Unknown

Check logical consistency

Unknown

Data characterisation

Data characterisation conducted

Unknown