# Antiplatelet Therapy during the First Year after Acute Coronary Syndrome in a Contemporary Italian Community of over 5 Million Subjects

First published: 08/05/2024 Last updated: 08/05/2024





# Administrative details

EU PAS number	
EUPAS1000000154	
Study ID	
100000154	
DARWIN EU® study	
No	
Study countries	
Italy	

**Study description** 

An observational retrospective non-interventional cohort study that has described patterns of antiplatelet therapy (APT) during the year following a hospital diagnosis of acute coronary syndrome (ACS) and possible implications in terms of revascularization rates, rehospitalizations, and costs for the Italian National Health Service (SSN).

#### **Study status**

**Finalised** 

# Research institutions and networks

## **Institutions**

Fondazione ReS (Ricerca e Salute), CINECA partner	
Italy	
First published: 05/07/2017	
<b>Last updated:</b> 01/10/2025	
Institution Not-for-profit ENCePP partner	

# Contact details

## **Study institution contact**

Silvia Calabria calabria@fondazioneres.it

Study contact

calabria@fondazioneres.it

**Primary lead investigator** 

# Letizia Dondi

**Primary lead investigator** 

# Study timelines

## Date when funding contract was signed

Actual: 08/01/2021

#### Study start date

Actual: 08/02/2021

#### Date of final study report

Actual: 08/06/2021

# Sources of funding

Pharmaceutical company and other private sector

# More details on funding

This research was partially funded by Sanofi Italy. No grant number has been generated. The funders had no role in the design of the study; in the collection, analyses, or interpretation of the data; in the writing of the manuscript, or in the decision to publish the results.

# Regulatory

Was the study required by a regulatory body?

## 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:

Drug utilisation

Healthcare resource utilisation

#### **Data collection methods:**

Secondary use of data

#### Study design:

Patients discharged (index date) with ACS diagnosis in 2017 were identified by an algorithm. Patients were characterized by revascularization rates at index

date, APT at one month and one year and rehospitalizations and healthcare costs during follow-up.

## Main study objective:

To investigate apparent discrepancies between guideline-recommended and real-world APT and the implications of such divergences for patients and healthcare systems.

# Study Design

## Non-interventional study design

Cohort

# Study drug and medical condition

## Medicinal product name, other

**Antiplatelets** 

## Study drug International non-proprietary name (INN) or common name

**ACETYLSALICYLIC ACID** 

**CLOPIDOGREL** 

**PRASUGREL** 

**TICAGRELOR** 

## **Anatomical Therapeutic Chemical (ATC) code**

(B01AC04) clopidogrel

clopidogrel

(B01AC05) ticlopidine

ticlopidine (B01AC06) acetylsalicylic acid acetylsalicylic acid (B01AC07) dipyridamole dipyridamole (B01AC22) prasugrel prasugrel (B01AC24) ticagrelor

ticagrelor

(B01AC30) combinations

combinations

#### Medical condition to be studied

Acute coronary syndrome

# Population studied

### **Age groups**

- Paediatric Population (< 18 years)
  - Preterm newborn infants (0 27 days)
  - Term newborn infants (0 27 days)
  - Infants and toddlers (28 days 23 months)
  - Children (2 to < 12 years)</li>
  - Adolescents (12 to < 18 years)</li>
- Adult and elderly population (≥18 years)
  - Adults (18 to < 65 years)</li>
    - Adults (18 to < 46 years)
    - Adults (46 to < 65 years)
  - Elderly (≥ 65 years)

- Adults (65 to < 75 years)</li>
- Adults (75 to < 85 years)
- Adults (85 years and over)

# Study design details

#### Setting

Inpatient and outpatient

#### **Summary results**

From the 2017 ReS database, 7966 (1.46x1000 inhabitants) were discharged alive with an ACS diagnosis. Most were >69 years and male. Of these, 83% (6640/7966) received at least 1 recommended antiplatelet agent within one month (treated group): 23% (1870/7966) as single and 60% (4770/7966) as dual APT. Among the 53% undergoing revascularization, 81% received dual APT at one month. Of the 78% with the same APT at one year, 66% showed appropriate coverage. For subjects treated and untreated with APT at one month, one-year rehospitalization rates were 54% and 66%, respectively, and mean per capita costs were EUR 14,316 and EUR 16,552, respectively (hospitalization driving >80% of costs).

# **Documents**

## **Study publications**

Antiplatelet Therapy during the First Year after Acute Coronary Syndrome in a C...

# 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 source(s), other

Fondazione ReS database

## **Data sources (types)**

Administrative healthcare records (e.g., claims)

# Use of a Common Data Model (CDM)

## **CDM** mapping

No

# Data quality specifications

#### **Check conformance**

Yes

## **Check completeness**

Yes

## **Check stability**

Yes

# **Check logical consistency**

Yes

# Data characterisation

## **Data characterisation conducted**

Yes