

# Coronary Artery Disease in Patients Older than 35 and Eligible for Cardiovascular Secondary Prevention: An Italian Retrospective Observational Analysis of Healthcare Administrative Databases

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Study

Finalised

## Administrative details

### EU PAS number

EUPAS1000000258

### Study ID

1000000258

### DARWIN EU® study

No

### Study countries

☐ Italy

## Study description

Background: This study describes patients with coronary artery disease (CAD) who are eligible for secondary prevention and assesses their healthcare consumption and costs from the perspective of the Italian National Health Service (SSN). Methods: From the Fondazione Ricerca e Salute's database, which collects Italian healthcare administrative data, all patients aged  $\geq 35$ , with  $\geq 1$  primary in-hospital CAD diagnosis and/or procedure on the coronary arteries, or with the specific disease exemption code, and who are suitable for long-term secondary prevention treatments, were identified in 2018 and analyzed. Demographics, comorbidities, one-year supplied drugs, hospitalizations, and costs were analyzed. Results: From  $>3$  million inhabitants aged  $\geq 35$ , 46,063 (1.3%) were identified (72.1% males, mean age 70 (12) years; approximately 50% with  $\geq 3$  comorbidities). During a one-year follow-up, 96.4% were treated with  $\geq 1$  drug for secondary prevention (mainly antiplatelets and lipid lowering agents), 69.4% with  $\geq 1$  concomitant cardiovascular drug, and 95.8% with  $\geq 1$  concomitant non-cardiovascular therapy. Within one year, 30.6% of patients were hospitalized at least once, mostly due to non-cardiovascular events. Calculated by mean, the INHS paid EUR 6078 per patient. Conclusions: This analysis confirms the relevant burden of CAD for patients with many comorbidities and who are frequently hospitalized, and the burden on the INHS. A multidisciplinary healthcare approach is encouraged to improve patients' outcomes and reduce costs for the INHS.

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## Study status

Finalised

## Research institutions and networks

### Institutions

## Fondazione ReS (Ricerca e Salute), CINECA partner

☐ Italy

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**Institution**

**Not-for-profit**

**ENCePP partner**

## Contact details

### Study institution contact

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**Study contact**

[calabria@fondazioneres.it](mailto:calabria@fondazioneres.it)

### Primary lead investigator

Letizia Dondi

**Primary lead investigator**

## Study timelines

### Date when funding contract was signed

Actual: 11/07/2019

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### Study start date

Actual: 12/09/2019

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### Date of final study report

Actual: 12/01/2020

## Sources of funding

- Pharmaceutical company and other private sector

## More details on funding

Acarpia Farmaceutici S.r.l, Milan, Italy

## Regulatory

**Was the study required by a regulatory body?**

No

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

Not applicable

## Methodological aspects

### Study type

### Study type list

**Study topic:**

Disease /health condition

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**Study type:**

Non-interventional study

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**Scope of the study:**

Drug utilisation

Healthcare resource utilisation

**Data collection methods:**

Secondary use of data

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**Study design:**

Retrospective longitudinal cohort study

**Main study objective:**

This study describes patients with coronary artery disease (CAD) who are eligible for secondary prevention and assesses their healthcare consumption and costs from the perspective of the Italian National Health Service (SSN)

## Study Design

**Non-interventional study design**

Cohort

## Study drug and medical condition

**Anatomical Therapeutic Chemical (ATC) code**

(B01AC) Platelet aggregation inhibitors excl. heparin

Platelet aggregation inhibitors excl. heparin

(C07A) BETA BLOCKING AGENTS

BETA BLOCKING AGENTS

(C09A) ACE INHIBITORS, PLAIN

ACE INHIBITORS, PLAIN

(C09B) ACE INHIBITORS, COMBINATIONS

ACE INHIBITORS, COMBINATIONS

(C09C) ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs), PLAIN

ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs), PLAIN

(C09D) ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs), COMBINATIONS

ANGIOTENSIN II RECEPTOR BLOCKERS (ARBs), COMBINATIONS

(C10) LIPID MODIFYING AGENTS

LIPID MODIFYING AGENTS

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### **Medical condition to be studied**

Coronary artery disease

## Population studied

### **Short description of the study population**

Among all patients older than 35 in the ReS database in 2018, and with at least a healthcare resource consumption in the charge of the SSN since 2015, subjects who were admitted to hospital at least once in 2018 (accrual) and whose hospital discharge contained a primary/secondary diagnosis of CAD and/or a procedure on coronary arteries, or patients with a CAD-specific cost sharing exemption code, were selected for the analysis.

## Study design details

### **Setting**

In-hospital and outpatient setting in public and affiliated with the SSN facilities

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## Summary results

From >3 million inhabitants aged  $\geq 35$ , 46,063 (1.3%) were identified (72.1% males, mean age 70 (12) years; approximately 50% with  $\geq 3$  comorbidities). During a one-year follow-up, 96.4% were treated with  $\geq 1$  drug for secondary prevention (mainly antiplatelets and lipid lowering agents), 69.4% with  $\geq 1$  concomitant cardiovascular drug, and 95.8% with  $\geq 1$  concomitant non-cardiovascular therapy. Within one year, 30.6% of patients were hospitalized at least once, mostly due to non-cardiovascular events. Calculated by mean, the SSN paid EUR 6078 per patient.

## Documents

### Study publications

[Coronary Artery Disease in Patients Older Than 35 and Eligible for Cardiovascu...](#)

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## 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)**

Database of Fondazione ReS

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**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

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**Check completeness**

Yes

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**Check stability**

Yes

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**Check logical consistency**

Yes

## Data characterisation

**Data characterisation conducted**

Yes