

# A multi-database cohort study to assess the incidence rates of colorectal hyperplasia among hypertensive patients

**First published:** 27/12/2014

**Last updated:** 23/04/2024

Study

Finalised

## Administrative details

### EU PAS number

EUPAS8295

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### Study ID

17564

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### DARWIN EU® study

No

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### Study countries

- ☐ Italy
  - ☐ Netherlands
  - ☐ Spain
  - ☐ United Kingdom
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## Study description

This study will estimate the frequency of colorectal polyps, cysts and neoplasms among hypertensive patients. Age- and sex-specific incidence rates will be estimated as well as the time from hypertension diagnosis to colorectal polyps, colorectal cysts and colorectal neoplasms. This multi-database dynamic cohort study, in four European primary care databases (CPRD, IPCI, CPRD, SIDIAP), includes all adult patients (aged 18-79 years) with an incident diagnosis of arterial hypertension. Incidence rates will be calculated by dividing the total number of incident cases over the total number of person-years at risk within the study population for each database separately. Standardization of incidence rates will be performed using either a European reference population or by internal validation using one of the four databases as reference population to which the other three databases will be compared to. Kaplan Meier analyses will be performed to estimate the survival time from arterial hypertension diagnosis date until one of the primary safety endpoints (colorectal polyps, cysts or neoplasms) but also for each of the endpoints separately.

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## Study status

Finalised

# Research institutions and networks

## Institutions

**Novartis Pharmaceuticals**

**First published:** 01/02/2024

**Last updated:** 01/02/2024

## Contact details

### Study institution contact

Clinical Disclosure Officer Novartis  
trialandresults.registries@novartis.com

Study contact

[trialandresults.registries@novartis.com](mailto:trialandresults.registries@novartis.com)

### Primary lead investigator

Clinical Disclosure Officer Novartis

Primary lead investigator

## Study timelines

### Date when funding contract was signed

Planned: 01/01/2015

Actual: 09/01/2015

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### Study start date

Planned: 01/01/2015

Actual: 15/05/2015

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### Date of final study report

Planned: 01/01/2016

Actual: 14/12/2015

## Sources of funding

- Pharmaceutical company and other private sector

## More details on funding

Novartis Pharma AG

## Study protocol

[CSPP100A2417-Redacted-Protocol.pdf](#) (637.02 KB)

## Regulatory

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

No

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

EU RMP category 3 (required)

## Other study registration identification numbers and links

CSPP100A2417

## Methodological aspects

### Study type

### Study type list

**Study topic:**

Disease /health condition  
Human medicinal product

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**Study type:**

Non-interventional study

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**Scope of the study:**

Disease epidemiology

**Data collection methods:**

Secondary use of data

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**Main study objective:**

This study will estimate the frequency of colorectal polyps, cysts and neoplasms among hypertensive patients. Age- and sex-specific incidence rates will be estimated as well as the time from hypertension diagnosis to colorectal polyps, colorectal cysts and colorectal neoplasms.

## Study Design

**Non-interventional study design**

Cohort

## Study drug and medical condition

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

ALISKIREN HEMIFUMARATE

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

(C09XA) Renin-inhibitors

Renin-inhibitors

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## **Medical condition to be studied**

Essential hypertension

Accelerated hypertension

Hypertension

Blood pressure increased

Blood pressure orthostatic increased

Blood pressure measurement

HELLP syndrome

Blood pressure ambulatory increased

Blood pressure diastolic increased

Blood pressure systolic increased

## **Population studied**

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

Adult patients (aged 18-80 years) in the individual databases with an incident diagnosis of arterial hypertension between January 01, 2000 and December 31, 2012.

Patients with continuous enrollment in the database for at least 1 year prior to the start of follow-up, aged 18 years or older at the start of follow-up and diagnosis of arterial hypertension were included.

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### **Age groups**

- Adults (18 to < 46 years)
- Adults (46 to < 65 years)

- Adults (65 to < 75 years)
- Adults (75 to < 85 years)

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### **Special population of interest**

Other

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### **Special population of interest, other**

Arterial hypertension patients

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### **Estimated number of subjects**

10000

## Study design details

### **Outcomes**

Incidence of colorectal polyps, cysts and neoplasms among hypertensive patients.

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### **Data analysis plan**

Incidence rates will be calculated by dividing the total number of incident cases over the total number of person-years at risk within the study population for each database separately.. Standardization of incidence rates will be performed using either a European reference population or by internal validation using one of the four databases as reference population to which the other three databases will be compared to. Kaplan Meier analyses will be performed to estimate the survival time from arterial hypertension diagnosis date until one of the primary safety endpoints (colorectal polyps, cysts or neoplasms) but also for each of the endpoints separately.

## Documents

## Study results

[SPP100A2417-Redacted-Final-Study-Report.pdf](#) (1.1 MB)

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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)

Clinical Practice Research Datalink

Health Search/IQVIA Health Longitudinal Patient Database

The Information System for Research in Primary Care (SIDIAP)

German Pharmacoepidemiological Research Database

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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**

Unknown

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**Check completeness**

Unknown

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**Check stability**

Unknown

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**Check logical consistency**

Unknown

Data characterisation

**Data characterisation conducted**

Unknown