

# Risk of Herpes Zoster in individuals diagnosed with COVID19 infection in the Valencia region of Spain: a retrospective cohort population-based study (AIV\_HZ\_2021\_05\_COVIDHZ\_AOS)

**First published:** 26/10/2021

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Study

Planned

## Administrative details

### EU PAS number

EUPAS43810

### Study ID

43811

### DARWIN EU® study

No

### Study countries

☐ Spain

## Study description

Based on the potential association between SARS-CoV-2 and HZ, we propose here a study to estimate the risk of HZ in subjects with SARS-CoV-2 infection using Real World Data (RWD). This population-based retrospective dynamic cohort study (meaning that members can leave or be added over time) will be based on eHR databases/registries from the Valencia Region (Valencia 7 Health System Integrated Database (VID) (27). VID allows linking socio-demographic, inpatients, outpatients, specialists, medication, and microbiology databases (among others) at individual level. The study population will consist of all population covered by the Public Health System (over 98%), representing about 5 million persons. As of March 8th, 2021, more than 381,919 confirmed cases of SARS-CoV-2 have been registered in the Valencian Community (4). Together with the VID makes the region of Valencia the ideal candidate to test the proposed hypothesis.

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## Study status

Planned

## Research institutions and networks

### Institutions

The Foundation for the Promotion of Health and Biomedical Research of Valencia Region (FISABIO)

☐ Spain

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**Last updated:** 05/11/2024

Institution

## Contact details

### Study institution contact

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Study contact

[alejandro.orrico@fisabio.es](mailto:alejandro.orrico@fisabio.es)

### Primary lead investigator

Alejandro Orrico-Sánchez

Primary lead investigator

## Study timelines

### Date when funding contract was signed

Planned: 24/09/2021

Actual: 24/09/2021

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### Study start date

Planned: 31/03/2022

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### Date of final study report

Planned: 01/10/2022

## Sources of funding

- Pharmaceutical company and other private sector

## More details on funding

GSK

## Study protocol

[PROTOCOL\\_AIV\\_V5.0.pdf](#)(1.06 MB)

## 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 type:**

Non-interventional study

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**Scope of the study:**

Assessment of risk minimisation measure implementation or effectiveness

Disease epidemiology

**Main study objective:**

Compare the risk of HZ among individuals 50 years and older with and without laboratory confirmed SARS-CoV2 infection. Compare the risk of HZ among individuals 18 years and older with and without laboratory confirmed SARS-CoV2 infection. Compare the risk of HZ in overall population older than 18 years old in the pandemic period against pre-pandemic period.

## Study Design

### **Non-interventional study design**

Cohort

## Study drug and medical condition

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

Herpes zoster

SARS-CoV-2 test positive

## Population studied

### **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)

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## **Special population of interest**

Renal impaired

Hepatic impaired

Immunocompromised

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## **Estimated number of subjects**

4000000

# **Study design details**

## **Outcomes**

Compare the risk of HZ among individuals 50 years and older with and without laboratory confirmed SARS-CoV2 infection. Compare the risk of HZ among individuals 18 years and older with and without laboratory confirmed SARS-CoV2 infection. Compare the risk of HZ in overall population older than 18 years old in the pandemic period against pre-pandemic period. Describe the likelihood of developing HZ after laboratory-confirmation of SARS-CoV2 according to severity of disease.

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## **Data analysis plan**

To develop the primary objective, we will compare the incidence of HZ among SARS-Cov-2 free and Sars-Cov-2 laboratory-confirmed subjects 50 years and older during the pandemic period. For this purpose, models using individual patient data or grouped will be implemented. The risk of HZ in subjects with Sars-cov-2 respect to subjects without Sars-Cov-2 will be estimated by a multivariate Poisson or negative binomial model according to applicability assumptions. Individual data will be grouped by SARS-Cov-2 free and Sars-Cov-2 laboratory-confirmed exposure, age, sex and comorbidities. Individual and grouped data can both be analyzed with the Poisson (or Negative binomial) distribution. Given the usual large size of the cohort, estimation when using

grouped data is considerably quicker than when using individual level data and the results for both approaches are remarkably similar. The number of HZ cases by aggregation unit will be compared among SARS-Cov-2 free and lab-confirmed subjects.

## 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 \(e.g., claims\)](#)

[Disease registry](#)

[Drug dispensing/prescription data](#)

[Electronic healthcare records \(EHR\)](#)

## 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**

No