

# CONSIGN study: COVID-19 infection and medicines in pregnancy - a multinational registry based study

**First published:** 10/02/2021

**Last updated:** 04/12/2025

Study

Finalised

## Administrative details

### EU PAS number

EUPAS39438

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### Study ID

48598


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### DARWIN EU® study

No

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### Study countries

 Denmark

 France


 Germany

 Italy

 Norway

 Spain

 Sweden

 United Kingdom

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## **Study description**

A retrospective multi-database dynamic cohort study, conducted during the years 2018 to 2020, including a period of SARS-CoV-2 circulation in Europe. The study population includes women of reproductive age (12-55 years), pregnant women and their children. The study will include data from 9 electronic health care registries in 8 European countries. Descriptive analysis will focus on 3-monthly prevalence rates of medication use, incidence rates of COVID-19 outcomes and prevalence of pregnancy outcomes. The primary objectives are: 1) To estimate the prevalence of medicines used, by trimester of pregnancy, and compare this among pregnant women with COVID-19, pregnant women without COVID-19, and non-pregnant women with COVID-19. 2) To describe severity and clinical outcomes of COVID-19 disease in pregnant women with COVID-19, according to treatments received during pregnancy, and compare these data with those of nonpregnant women of reproductive age with COVID-19. 3) To assess and compare the rates of adverse maternal and neonatal outcomes in pregnant women with and without COVID-19, using different medicines.

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## **Study status**

Finalised

## **Research institutions and networks**

### **Institutions**

## Pharmacoepidemiology and Drug Safety Research Group (PharmaSafe), University of Oslo

 Norway

**First published:** 19/10/2016


**Last updated:** 06/11/2025

**Institution**

**Educational Institution**

**ENCePP partner**

## Division of Pharmacoepidemiology & Clinical Pharmacology (PECP), Utrecht Institute for Pharmaceutical Sciences (UIPS), Utrecht University

 Netherlands

**First published:** 01/03/2010

**Last updated:** 23/05/2024

**Institution**

**Educational Institution**

**ENCePP partner**

## Health Services Research and Pharmacoepidemiology Unit (HSRP Unit) FISABIO

 Spain

**First published:** 30/11/2023

**Last updated:** 30/11/2023

Institution

Other

ENCePP partner

## EpiChron Research Group on Chronic Diseases, Aragon Health Sciences Institute (IACS)

 Spain

**First published:** 17/02/2017


**Last updated:** 02/04/2024

Institution

Educational Institution

ENCePP partner

## Aarhus University & Aarhus University Hospital DEPARTMENT OF CLINICAL EPIDEMIOLOGY

 Denmark


**First published:** 20/07/2021

**Last updated:** 08/05/2026

Institution

Educational Institution

## University Medical Center Utrecht (UMCU)

 Netherlands

**First published:** 24/11/2021

**Last updated:** 22/02/2024


Institution

Educational Institution

Hospital/Clinic/Other health care facility

ENCePP partner

## Centre for Pharmacoepidemiology, Karolinska Institutet (CPE-KI)

 Sweden

**First published:** 24/03/2010

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

Institution


Educational Institution

Laboratory/Research/Testing facility


Not-for-profit


ENCePP partner


## RTI Health Solutions (RTI-HS)


 France

 Spain

 Sweden

 United Kingdom

 United Kingdom (Northern Ireland)

 United States

**First published:** 21/04/2010


**Last updated:** 13/03/2025

Institution

Not-for-profit

ENCePP partner

## Bordeaux PharmacoEpi, University of Bordeaux

 France

**First published:** 07/02/2023

**Last updated:** 08/12/2025

**Institution**


Educational Institution

Hospital/Clinic/Other health care facility

Not-for-profit

ENCePP partner

## Swansea University Medical School

 United Kingdom

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

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

**Institution**

Educational Institution

Hospital/Clinic/Other health care facility

## Agenzia regionale di sanità della Toscana (ARS Toscana)

 Italy

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

**Last updated:** 23/03/2026

**Institution**

EU Institution/Body/Agency

ENCePP partner

## Vall d'Hebron University Hospital - Vall d'Hebron Research Institute (HUVH/VHIR)

 Spain

**First published:** 01/07/2025

**Last updated:** 24/07/2025


**Institution**

Hospital/Clinic/Other health care facility

Laboratory/Research/Testing facility

## Networks

### EU Pharmacoepidemiology and Pharmacovigilance (PE&PV) Research Network

 Netherlands

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

**Last updated:** 24/09/2025

**Network**

## Contact details

### Study institution contact

Hedvig Nordeng h.m.e.nordeng@farmasi.uio.no

## Study contact

[h.m.e.nordeng@farmasi.uio.no](mailto:h.m.e.nordeng@farmasi.uio.no)

### Primary lead investigator

Hedvig Nordeng

## Primary lead investigator

## Study timelines

### Date when funding contract was signed

Planned: 17/07/2020

Actual: 17/07/2020

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### Study start date

Planned: 01/01/2021

Actual: 01/01/2021

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### Date of interim report, if expected

Planned: 14/10/2021

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### Date of final study report

Planned: 28/07/2023

Actual: 07/11/2023

## Sources of funding

- Other

## More details on funding

EMA

## Study protocol

[ProtocolCONSIGNWP1v1.0\\_EUPASS.pdf](#) (2.11 MB)

[EUPAS39438 CONSIGN WP1 protocol including amendment 1.pdf](#) (2.31 MB)

## Regulatory

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

Yes

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

Not applicable

## Other study registration identification numbers and links

[Link to EU PE&PV website](#)

## 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:**

Disease epidemiology

Drug utilisation

**Data collection methods:**

Secondary use of data

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**Main study objective:**

To estimate prevalence of medicines use in pregnant COVID-19 patients and to describe severity of clinical outcomes as well as pregnancy and neonatal outcomes

## Study Design

**Non-interventional study design**

Case-control

Cohort

## Study drug and medical condition

**Medical condition to be studied**

COVID-19

## Population studied

## Age groups

- Adolescents (12 to < 18 years)
  - Adults (18 to < 46 years)
  - Adults (46 to < 65 years)
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## Special population of interest

Pregnant women

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## Estimated number of subjects

1000000

# Study design details

## Outcomes

Use of medication Severity of COVID-19 Pregnancy and neonatal outcomes

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## Data analysis plan

Descriptive analysis, matched analysis of cohorts and sensitivity analysis will be conducted.

# Documents

[Study report on Zenodo](#)

## Study publications

[Hurley, Eimir, Sturkenboom, Miriam, Poblador-Plou, Beatriz, Sanfelix-Gimeno, Ga...](#)

[CONSIGN community on Zenodo \(includes all publications currently available with...](#)

## 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)**

Danish registries (access/analysis)

German Pharmacoepidemiological Research Database

ARS Toscana

Norwegian Linked Health registry at University of Oslo

Système National des Données de Santé (French national health system main database)

EpiChron Cohort

The Valencia Health System Integrated Database

SAIL Databank

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### **Data source(s), other**

Swedish registries

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## **Data sources (types)**

Administrative healthcare records (e.g., claims)

Disease registry

Electronic healthcare records (EHR)

Other

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## **Data sources (types), other**

Prospective patient-based data collection, Exposure registry

# 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