

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

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

**Last updated:** 22/04/2024

Study

Ongoing

## 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**

Ongoing

## Research institutions and networks

### Institutions

## University of Oslo

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

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

**Institution**

## 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**

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

☐ Norway

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

**Last updated:** 08/11/2016

**Institution**

**Educational Institution**

**ENCePP partner**

## Leibniz Institute for Prevention Research and Epidemiology - BIPS

☐ Germany

**First published:** 29/03/2010

**Last updated:** 26/02/2024

**Institution**

Not-for-profit

ENCEPP partner

## Fundació Institut Català de Farmacologia (FICF)

☐ Spain

**First published:** 29/03/2010

**Last updated:** 17/09/2019

**Institution**

Educational Institution

Hospital/Clinic/Other health care facility

Not-for-profit

ENCEPP partner

## 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:** 08/02/2023

**Institution**

**Educational Institution**

**Hospital/Clinic/Other health care facility**

**Not-for-profit**

**ENCePP partner**

## Instituto Aragonés de Ciencias de la Salud (IACS)

☐ Spain

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

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

**Institution**

**Educational Institution**

## Aarhus University Hospital

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

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

**Institution**

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

☐ Spain

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

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

**Institution**

University of Oslo

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

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

**Institution**

University of Oslo Norway, Aarhus University  
Hospital Denmark, University of Copenhagen  
Denmark, FISABIO Spain, IACS Spain, Swansea  
University UK

## Networks

EU Pharmacoepidemiology and Pharmacovigilance  
(PE&PV) Research Network

☐ Netherlands

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

**Last updated:** 26/11/2024

**Network**

## Contact details

### Study institution contact

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Study contact

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### 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

## 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

## 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

Drug utilisation

**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

## 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**

Hepatic impaired

Immunocompromised

Pregnant women

Renal impaired

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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 publications

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

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

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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)**

Danish registries (access/analysis)

German Pharmacoepidemiological Research Database

ARS Toscana

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**Data source(s), other**

Danish Registries (access/analysis), NorPD, GePaRD, ARS

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**Data sources (types)**

Administrative healthcare records (e.g., claims)

Disease registry

Drug dispensing/prescription data

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