

SAFETY-VAC: Phenotype proposal and rates of immunocompromised populations in real-world data sources.

First published: 24/02/2025

Last updated: 23/10/2025

Study

Planned

Administrative details

EU PAS number

EUPAS1000000288

Study ID

1000000288

DARWIN EU® study

No

Study countries

☐ Denmark

☐ Finland

☐ France

☐ Italy

- ☐ Norway
 - ☐ Spain
 - ☐ United Kingdom
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Study description

This SAFETY-VAC project aims to provide a consolidated approach to identify and characterise immunocompromised populations in real-world data sources and to produce incidence and prevalence rates for these populations using electronic healthcare records databases.

Study status

Planned

Research institutions and networks

Institutions

Aarhus University & Aarhus University Hospital
DEPARTMENT OF CLINICAL EPIDEMIOLOGY

☐ Denmark

First published: 20/07/2021

Last updated: 02/04/2024

Institution

Educational Institution

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

Fundació Institut Universitari per a la Recerca a l'Atenció Primària de Salut Jordi Gol i Gurina, IDIAPJGol

☐ Spain

First published: 05/10/2012

Last updated: 23/05/2025

Institution

Educational Institution

Laboratory/Research/Testing facility

Not-for-profit

ENCEPP partner

Bordeaux PharmacoS, 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

Instituto Aragonés de Ciencias de la Salud (IACS)

☐ Spain

First published: 01/02/2024

Last updated: 02/04/2024

Institution

Educational Institution

Agenzia regionale di sanità della Toscana (ARS)

☐ Italy

First published: 01/02/2024

Last updated: 12/03/2024

Institution

EU Institution/Body/Agency

ENCePP partner

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

☐ Spain

First published: 01/02/2024

Last updated: 31/10/2025

Institution

Agencia Española de Medicamentos y Productos Sanitarios (Spanish Agency for Medicines and Medical Devices, AEMPS)

☐ Spain

First published: 01/02/2024

Last updated: 04/09/2024

Institution

EU Institution/Body/Agency

Not-for-profit

Regulatory Authority

ENCePP partner

University of Oslo

First published: 01/02/2024

Last updated: 01/02/2024

Institution

Teamit Institute

☐ Spain

First published: 12/03/2024

Last updated: 12/03/2024

Institution

Other

ENCePP partner

Utrecht University (UU), University of Eastern
Finland (UEF)

Networks

Vaccine monitoring Collaboration for Europe (VAC4EU)

- ☐ Belgium
- ☐ Denmark
- ☐ Finland
- ☐ France
- ☐ Germany
- ☐ Italy
- ☐ Netherlands
- ☐ Norway
- ☐ Spain
- ☐ United Kingdom

First published: 22/09/2020

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Network

Outdated

ENCePP partner

EU Pharmacoepidemiology and Pharmacovigilance
(PE&PV) Research Network

☐ Netherlands

First published: 01/02/2024

Last updated: 24/09/2025

Network

Pedianet network (So.Se.Te)

☐ Italy

First published: 23/10/2025

Last updated: 23/10/2025

Network

Contact details

Study institution contact

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Study contact

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Primary lead investigator

Carlos Durán Salinas

Primary lead investigator

Study timelines

Date when funding contract was signed

Planned: 15/01/2024

Actual: 06/02/2024

Study start date

Planned: 15/05/2024

Date of final study report

Planned: 16/09/2024

Sources of funding

- EMA

Regulatory

Was the study required by a regulatory body?

Yes

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

Not applicable

Methodological aspects

Study type

Study type list

Study topic:

Disease /health condition

Study type:

Non-interventional study

Scope of the study:

Disease epidemiology

Method development or testing

Data collection methods:

Secondary use of data

Study design:

- Scoping review aimed at describing and identifying immunocompromised populations in epidemiological studies using EHRs.
- Retrospective cohort study from January 1, 2017, to the last available data aimed at identifying and characterizing these populations using RWD.

Main study objective:

- To describe what operational definitions are used to identify immunocompromised populations when conducting epidemiological research in EHR databases and propose a phenotype to identify immunocompromised individuals in database studies correctly.
- To estimate the incidence and one-year prevalence rates of immunocompromised status based on the agreed phenotype and different durations of episodes.
- To assess the contribution of different provenances of data and durations to the prevalence of the immunocompromised status.

Study Design

Non-interventional study design

Cohort

Population studied

Short description of the study population

Electronic health records of people from 10 data sources in 7 countries in Europe from January 1st, 2017, until the last data availability were selected.

Persons were included in the study population when they had:

- Information on age and gender available,
- At least one day of follow in the study period (1/1/2017- latest availability).

Follow-up started at the latest date of any of the following dates: day that one year of lookback is available during the study period, or at 1/1/2017 when the person is born in the data base.

Follow-up finished at the earliest of the following dates: death, disenrollment, end of study period, or recommended end date.

Age groups

- **Paediatric Population (< 18 years)**
- **Adult and elderly population (≥ 18 years)**

Study design details

Setting

The study will use data from 10 secondary electronic health record databases that are population-based in 7 countries in Europe (UK, Spain, Denmark, Finland, Norway, Italy and France).

The study will be conducted during the period from January 1st, 2017, to the last data availability, specific for each data source.

The source population comprises all persons in the data sources who can be potentially included in the study.

Documents

Study report

[SAFETY-VAC_Immunocompromised_population_D4_Report.pdf](#) (1009.31 KB)

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)

Pedianet

Norwegian Health Registers

The Valencia Health System Integrated Database

EpiChron Cohort

Danish Health Data Registries

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

BIFAP - Base de Datos para la Investigación Farmacoepidemiológica en el
Ámbito Público (Pharmacoepidemiological Research Database for Public Health
Systems)

Clinical Practice Research Datalink (CPRD) GOLD

The Information System for Research in Primary Care (SIDIAP)

Data source(s), other

Finnish national data registers

Use of a Common Data Model (CDM)

CDM mapping

Yes

Data quality specifications

Check conformance

Yes

Check completeness

Yes

Check stability

Yes

Check logical consistency

Yes

Data characterisation

Data characterisation conducted

Yes