- PGRx: Rheumatoid polyarthritis suspicion

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General

Identification

Detailed name	PGRx: Rheumatoid polyarthritis suspicion
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL : 907.309

General Aspects

Medical area	Internal medicine Rare diseases Rheumatology
Health determinants	Genetic
Others (details)	Rheumatoid polyarthritis (PAR) suspicion
Keywords	pharmaco-epidemiology

Scientific investigator(s) (Contact)

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Collaborations

Funding

Funding status	Mixed
Details	LA-SER

Governance of the database

Sponsor(s) or organisation(s) responsible	LA-SER
Organisation status	Private

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Case control study
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Cases of central demyelination are recruited by a network of internal medicine, rheumatology and pediatric centers, and the referral agents by a network of general practitioners, distributed all across the territory.

Database objective

Main objective	Surveillance and evaluation of the risk of an initial rheumatoid polyarthritis, associated with drug or vaccine exposure under real treatment situations.
Inclusion criteria	For cases: patient, men or women, aged from 18 to 79 included, having a clinico-biological presentation evocative of rheumatoid polyarthritis declared by the specialist, the early symptoms of the episode of which date back less than 12 months. For controls: patient, men or women aged from 18 to 79 included, having consulted a general practitioner.

Population type

Age	Adolescence (13 to 18 years) Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years)
Population covered	General population

Gender	Male Woman
Geography area	International
Detail of the geography area	France, Italy, Spain, United Kingdom, Canada

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	04/2008

Size of the database

Size of the database (number of individuals)	[10 000-20 000[individuals
Details of the number of individuals	- 151 cas/cases - 11125 témoins/referents

Data

Database activity	Current data collection
Type of data collected	Clinical data Declarative data
Clinical data (detail)	Direct physical measures
Declarative data (detail)	Phone interview
Presence of a biobank	No
Health parameters studied	Health event/morbidity

Procedures

Participant monitoring	No
Links to administrative sources	No

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	Terms of data access to be defined.

Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only