

# - PGRx : Central demyelination

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## General

### Identification

Detailed name PGRx : Central demyelination

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL : 907.069

### General Aspects

Medical area Neurology

Others (details) First central demyelinating event, Demyelination, Multiple Sclerosis (MS)

Keywords pharmaco-epidemiology

### Scientific investigator(s) (Contact)

Name of the director Grimaldi - Bensouda

Surname Lamiae

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### Collaborations

#### Funding

Funding status Private

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 as part of an interventional study	No
Additional information regarding sample selection.	Cases of central demyelination are recruited by a network of neurology and pediatric centers and the referents by a network of general practitioners, distributed all across the territory.

### Database objective

Main objective	Surveillance and assessment of the risk of central demyelination associated with drug or vaccine use in real life situations.
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Inclusion criteria	For the cases: patients, males and females aged from 0 to 79 years, having a first central demyelinating event reported by the neurologist, for which the first symptoms of the episode occurred less than 12 months ago. For the controls: patients, males and females aged from 0 to 79 years, having consulted a general practitioner.
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### Population type

Age	Newborns (birth to 28 days) Infant (28 days to 2 years) Early childhood (2 to 5 years) Childhood (6 to 13 years) 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)
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Population covered	General population
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Gender	Male Woman
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Geography area	International
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Detail of the geography area	France, Italy, Spain, United Kingdom, Canada
<b>Data collection</b>	
<b>Dates</b>	
Date of first collection (YYYY or MM/YYYY)	2007
<b>Size of the database</b>	
Size of the database (number of individuals)	[10 000-20 000[ individuals
Details of the number of individuals	517 cas et 11125 référents
<b>Data</b>	
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
<b>Procedures</b>	
Participant monitoring	No
Links to administrative sources	No
<b>Promotion and access</b>	
<b>Promotion</b>	
<b>Access</b>	
Terms of data access (charter for data provision, format of data, availability delay)	Methods for accessing the database are currently being defined
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only

