

# EPIMART - Cross-sectional Study on Epileptic Patients Living in Martinique

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## General

### Identification

Detailed name Cross-sectional Study on Epileptic Patients Living in Martinique

Sign or acronym EPIMART

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

### General Aspects

Medical area Neurology

Health determinants Geography  
Lifestyle and behavior

Keywords French West Indies, Martinique, incidence

### Scientific investigator(s) (Contact)

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Organization Hôpital Cantonal de

### Collaborations

### Funding

Funding status Public

Details	Cantonal Hospital of Geneva
Governance of the database	
Sponsor(s) or organisation(s) responsible	CEA
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Not-repeated cross-sectional studies (except case control studies)
Database recruitment is carried out by an intermediary	A selection of health care professionals
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Individuals attending a hospital or neurologist on the island after their first spontaneous epileptic seizure were recruited for the study.
Database objective	
Main objective	To determine the overall incidence of epileptic seizures within the island population and to identify clinical seizure type and factors.
Inclusion criteria	<ul style="list-style-type: none"> <li>- Male and female;</li> <li>- Living in Martinique;</li> <li>- Experienced an epileptic seizure between 01/05/1994 and 31/04/1995.</li> </ul>
Population type	
Age	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)
Population covered	Sick population

Gender	Male Woman
Geography area	Departmental
French regions covered by the database	Martinique
Detail of the geography area	Island of Martinique.
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	05/1994
Date of last collection (YYYY or MM/YYYY)	04/1995
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	309
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Declarative data Paraclinical data
Clinical data (detail)	Direct physical measures
Declarative data (detail)	Paper self-questionnaire
Paraclinical data (detail)	- Electroencephalography (EEG) - computed tomography.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality
Procedures	
Participant monitoring	No

Links to administrative sources	No
Promotion and access	
Promotion	
Link to the document	<a href="http://onlinelibrary.wiley.com/doi/10.1111/j.1528-1157.1999.tb00826.x/pdf">http://onlinelibrary.wiley.com/doi/10.1111/j.1528-1157.1999.tb00826.x/pdf</a>
Link to the document	<a href="http://www.ncbi.nlm.nih.gov/pubmed/9773072">http://www.ncbi.nlm.nih.gov/pubmed/9773072</a>
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only