MFV - Cross-sectional study on Migrations-Family-Ageing in French overseas departments

Head :Claude-Valentin Marie

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General

Identification

Detailed name	Cross-sectional study on Migrations-Family-Ageing in French overseas departments
Sign or acronym	MFV
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL

General Aspects

Medical area	General practice
Health determinants	Geography Healthcare system and access to health care services Lifestyle and behavior Social and psychosocial factors
Keywords	French overseas department, Demography, Intergenerational, Language practices, Religious practices, Migrations, Discrimination, Family, Ageing, Health

Scientific investigator(s) (Contact)

Name of the director	Claude-Valentin
Surname	Marie
Phone	+33 (0)1 56 06 21 83
Email	claude-valentin.marie@ined.fr
Organization	INED

Collaborations

Funding

Funding status	Public
Details	INED

Governance of the database

Sponsor(s) or organisation(s) responsible	INED
Organisation status	Public
Sponsor(s) or organisation(s) responsible	Insee
Organisation status	Public
Presence of scientific or steering committees	Yes

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 population file
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	The survey was conducted among men and women between the ages of 18 and 79, in a representative sample of 4,000 households in each department, i.e. a total of 16,000 for the four French overseas departments.

Database objective

Main objective	The MFV survey was carried out in a coordinated way in all the French overseas departments and focuses on the changes in family life, the sources and obstacles of social itineraries, the impact of migration and the evolution of intergenerational

	solidarity
Inclusion criteria	Three main categories of key player were asked: * Native people of the department * Returning migrants: native people of the department who have had an emigration experience of more than six consecutive months and who were resettled there at the time of the survey * Immigrants: the non-native people of the department who were settled there at the time of the survey, whether French or foreign
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years)
Population covered	General population
Pathology	
Gender	Male Woman
Geography area	Local
French regions covered by the database	Guadeloupe Guyane Martinique La Réunion
Detail of the geography area	Guadeloupe (GP) French Guiana (GY) Martinique (MQ) Réunion (RE)
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2009
Date of last collection (YYYY or MM/YYYY)	2010
Size of the database	
Size of the database (number of individuals)	[10 000-20 000[individuals
Details of the number of individuals	15 770

Data	
Database activity	Data collection completed
Type of data collected	Declarative data
Declarative data (detail)	Face to face interview
Details of collected declarative data	THE EXTENDED FAMILY MODULE AND FAMILY SOLIDARITES (Members of the Ego family currently migrating, Solidarity and caregivers, State of health and access to health care) ; THE FAMILY MODULE (The children of Ego and his spouse, The present union and past unions of Ego, Mastery and the future of fertility); THE MODULE ON THE PROFESSIONAL AND MIGRATORY TRAJECTORIES OF NATIVE PEOPLE AND IMMIGRANTS (Social and professional trajectories, The professional and migratory trajectories of the native people of the department, The migratory and professional trajectories of people born outside the department, Experiences of rejection or discrimination in the department); THE CULTURAL RESOURCES MODULE (Citizen life, Languages, Religion)
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health care consumption and services Quality of life/health perception
Care consumption (detail)	Medicines consumption
Procedures	
Data collection method	CAPI questionnaire
Participant monitoring	No
Followed pathology	
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of	Contact the scientist in charge

data, availability delay)	
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only