- General Cancer Registry of French Polynesia

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General

Identification

Detailed name	General Cancer Registry of French Polynesia
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL

General Aspects

Medical area	Cancer research
Health determinants	Geography
Keywords	Polynesia, incidence, diagnosis, screening

Scientific investigator(s) (Contact)

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Organization	Institut National de la Santé et de la Recherche

Collaborations

Funding

Funding status	Public
Details	Health Directorate of French Polynesia

Governance of the database

Sponsor(s) or organisation(s) responsible	Direction de la Santé de la Polynésie française
Organisation status	Public
Sponsor(s) or organisation(s) responsible	Office de Protection contre les Rayonnements Ionisants
Organisation status	Public
Sponsor(s) or organisation(s) responsible	Institut National de la Santé et de la Recherche Médicale
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Morbidity registers
Additional information regarding sample selection.	Cancer cases are identified based on reports from private and public physicians, examination reports from anatomopathology laboratories throughout the territory, PMSI data from the Mammao Regional Hospital Centre (Centre Hospitalier Territorial de Mammao), medical evacuations and death certificates.

Database objective

Main objective	To identify all new cancer cases among individuals living in French Polynesia. To conduct descriptive, analytic and evaluative epidemiological work with this data. Assessment of healthcare and diagnostic practices. Assessment of screening practices. Participation in etiological and prognostic research.
Inclusion criteria	- male and female - living in French Polynesia at time of diagnosis - subject with cancer

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) Great age (80 years and more)
Population covered	Sick population
Gender	Male Woman
Geography area	Departmental
Detail of the geography area	French Polynesia

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1985

Size of the database

Size of the database (number of individuals)	[10 000-20 000[individuals
Details of the number of individuals	475/an/year

Data

Database activity	Current data collection
Type of data collected	Clinical data
Clinical data (detail)	Direct physical measures
Details of collected clinical data	Patient sociodemographic characteristics (surname, maiden name, first name, age, sex, place of residence, date and place of birth, vital status) Cancer morphology and topography. Patient's vital status.
Presence of a biobank	No
Health parameters studied	Health event/morbidity

	Health event/mortality

Procedures

Data collection method	Data are gathered through a report form, including patient identification, topographical characteristics and tumour histology, treatment, monitoring and report source.
Quality procedure(s) used	- Internal quality control for data validation by establishing quality indicators once a year. - Using quality criteria from the FRANCIM network and French National Committee of Registries. - Data validation protocol: comprehensive consultation of data sources from laboratories and DIMs (Medical Information Departments); missing and duplicated data control, inconsistency control performed using IARC Tools and corrections carried out.
Participant monitoring	Yes
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://onlinelibrary.wiley.com/doi/10.1046/j.1365-3156.2000.00624.x/full
Link to the document	http://tinyurl.com/PUBMED-CRFP
Description	Liste des publications dans Pubmed

Access

Terms of data access (charter for data provision, format of data, availability delay)	Data can only be disseminated through direct involvement with those in charge of the registries.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only