- New Caledonia Cancer Registry (Certified Registry 2013-2015)

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General

Identification

Detailed name	New Caledonia Cancer Registry (Certified Registry 2013-2015)
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL 998250 (08/06/1999)

General Aspects

Medical area	Cancer research
Health determinants	Geography
Keywords	standardised incidence rate, epidemiology

Scientific investigator(s) (Contact)

Name of the director	Laumond
Surname	Sylvie
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Organization	Direction des affaires sanitaires et sociales de la Nouvelle-Calédonie -

Collaborations

Funding

Funding status	Public
Details	New Caledonia

Governance of the database

Sponsor(s) or organisation(s) responsible	Direction des affaires sanitaires et sociales de la Nouvelle-Calédonie - DASS
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Morbidity registers
Additional information regarding sample selection.	All primary invasive tumours, haematological malignancies, benign tumours or unpredictable developments in the central nervous system. Superficial non-invasive tumours of the bladder and in situ tumours of the colon, breast, cervix and melanoma.

Database objective

Main objective	To identify all new cancer cases in individuals living in New Caledonia.
Inclusion criteria	- Male and female; - Living in New Caledonia; - 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	General population
Gender	Male Woman
Geography area	Local
Detail of the geography area	New Caledonia.

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1977

Size of the database

Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	252,000

Data

Database activity	Current data collection
Type of data collected	Clinical data Administrative data
Clinical data (detail)	Direct physical measures
Details of collected clinical data	A standardised sheet is completed for each new cancer case that is reported and verified.
Administrative data (detail)	Date of birth, sex, ethnicity, place of residence.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Data collected by public and private anatomical pathology laboratories, as well as public and private physicians.
Quality procedure(s) used	Ongoing quality procedure.
Participant monitoring	No
Links to administrative sources	Yes
Linked administrative sources (detail)	- Network of registries in the Pacific region - South Pacific Commission - International Agency for Research on Cancer (IARC).

Promotion and access

Promotion

Link to the document	http://www.dass.gouv.nc/portal/page/portal/dass/observatoire_sante/Le%20registre%20du%20cancer
Link to the document	http://tinyurl.com/PUBMED-RCNC
Description	Liste des publications dans Pubmed
Link to the document	http://tinyurl.com/HAL-RCNC
Description	Liste des publications dans HAL
Access	
Terms of data access (charter for data provision, format of data, availability delay)	The data is only avaiable by request to the health medical inspector, president of the Cancer Registry of the committee. The Pasteur Institute has no role in the management of the registry.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only