

- 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)

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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	<ul style="list-style-type: none"> - Male and female; - Living in New Caledonia; - With cancer.
Population type	
Age	<p>Newborns (birth to 28 days)</p> <p>Infant (28 days to 2 years)</p> <p>Early childhood (2 to 5 years)</p> <p>Childhood (6 to 13 years)</p> <p>Adolescence (13 to 18 years)</p> <p>Adulthood (19 to 24 years)</p> <p>Adulthood (25 to 44 years)</p> <p>Adulthood (45 to 64 years)</p> <p>Elderly (65 to 79 years)</p> <p>Great age (80 years and more)</p>
Population covered	General population
Gender	<p>Male</p> <p>Woman</p>
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 available 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