

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

Head :Laumond Sylvie

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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

Address BP N4 98851 Nouméa cedex

Phone 00 687 24 37 15 / 00 687 27 20 64

Email sylvie.laumond@gouv.nc

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
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Organisation status	Public
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## Additional contact

## Main features

## Type of database

Type of database	Morbidity registers
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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.
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## Database objective

Main objective	To identify all new cancer cases in individuals living in New Caledonia.
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Inclusion criteria	<ul style="list-style-type: none"><li>- Male and female;</li><li>- Living in New Caledonia;</li><li>- With cancer.</li></ul>
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## 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)
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Population covered	General population
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Gender	Male Woman
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Geography area	Local
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Detail of the geography area	New Caledonia.
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## 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	<a href="http://www.dass.gouv.nc/portal/page/portal/dass/observatoire_sante/Le%20registre%20du%20cancer">http://www.dass.gouv.nc/portal/page/portal/dass/observatoire_sante/Le%20registre%20du%20cancer</a>
Link to the document	<a href="http://tinyurl.com/PUBMED-RCNC">http://tinyurl.com/PUBMED-RCNC</a>
Description	Liste des publications dans Pubmed
Link to the document	<a href="http://tinyurl.com/HAL-RCNC">http://tinyurl.com/HAL-RCNC</a>
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