- Cancer Registry of the Manche (qualified registry)

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General

Identification

Detailed name	Cancer Registry of the Manche (qualified registry)
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	332500

General Aspects

Medical area	Cancer research
Others (details)	Cancer

Scientific investigator(s) (Contact)

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Collaborations

Funding

Funding status	Public
Details	InVS, l'INCa et l'ARKM

Governance of the database

Sponsor(s) or organisation(s) responsible	Centre Hospitalier Public du Cotentin
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Morbidity registers
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Selection of subjects having the required inclusion criteria. Several sources are used to identify cases: - Anatomopathological laboratories - Information Departments (DIM: PMSI data) - Radiotherapy centers - Permanent cancer survey (EPC) of the CLCC (cancer research centers) - Healthcare institutions (ES) - Specialists (private-practicing and hospital-practicing) and general practitioners - Specialized registries - Summaries of the Pluridisciplinary Concertation Meetings (RCP)

Database objective

Main objective	The Registry of Cancers for la Manche was created in 1994. It stems from a local medical need, in order to supplement the incidence rates measured in the neighboring département of Calvados and to answer the public health questions raised by the industrial activity in the département (the nuclear industry predominantly and the use of asbestos in civil and military ship building). The registry performs a permanent and complete collection of all of the new diagnoses of cancer in the general population residing in the département of la Manche. It has a dual objective of describing and monitoring the risk of cancer and of conducting research using the analysis of the data collected or one-off surveys. Objectives of the Registry in terms of public health: - Participate in the epidemiological monitoring of cancers on a local and national level within the framework of the Francim network, through

	publishing incidence indicators by cancer location, sex, age and year of diagnosis. - Contribute to assessing primary and secondary prevention initiatives (organized cancer screenings), caring for patients and the care needs for the general population.
Inclusion criteria	All malignant tumors diagnosed in patients residing in the département of la Manche at the time of the diagnosis. These are invasive malignant tumors, in-situ, benign tumors and intermediate malignancy of the bladder and of the central nervous system. Skin tumors of the basal cell type are excluded.

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
French regions covered by the database	Normandie
Detail of the geography area	Department of La Manche

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1994
Date of last collection (YYYY or MM/YYYY)	2008

Size of the database

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

Details of the number of individuals	1994-2008 : 45 078

Data

Database activity	Data collection completed
Type of data collected	Clinical data Administrative data
Clinical data (detail)	Direct physical measures
Administrative data (detail)	Identification data
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Active or passive collection (CNIL Authorization): - Reception of the anatomocytopathological reports, lists of stays (PMSI). Lists via encrypted computer transmission. - Active search for additional regular information in the medical records.
Participant monitoring	Yes
Details on monitoring of participants	- Vital status, date of latest news - An active follow-up is performed on a sample (High Resolution Francim surveys). - A passive follow-up is conducted using information received at the Registry.
Links to administrative sources	Yes

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	The incidence data is available in an aggregate form, by cancer location, year, sex and age. These are published on a regular basis on a local and national level (cf. Find out more). Regional documents can be downloaded from the registry website. The data recorded in the Registry database can be used for specific studies. The methods for collaboration have to be defined.

Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only