- Finistère Registry for Digestive Tumours (Certified Registry 2013-2016)

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General	
Identification	
Detailed name	Finistère Registry for Digestive Tumours (Certified Registry 2013-2016)
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	908270
General Aspects	
Medical area	Cancer research Gastroenterology et hepatology
Scientific investigator(s) (Contact)	
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Unit	Coordinatrice
Organization	CHUR
Collaborations	
Funding	
Funding status	Public
Details	French Institute for Public Health Surveillance - INVS National Cancer Institute - InCA League
Governance of the database	
Sponsor(s) or organisation(s) responsible	CHU Brest
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Morbidity registers
Additional information regarding sample selection.	Selection of subjects meeting the inclusion criteria. Several sources are used to identify cases:- Clinical departments - University Hospital Centres (UHC)- Other public clinical departments- Private clinical departments- Private gastroenterologist practices ?

General practitioner surgeries - Medical Information Department (DIM) of the UHC - DIM of other institutions - Anatomopathology laboratory -Regional oncology networks ? Civil registry -National PMSI database - Local medical departments of various health insurance funds (CPAM in Finistère, MSA, RSI, etc.).

Database objective

Main objective	Objectives of the registry in terms of public health (surveillance, evaluation): Thanks to the multi-source collection of digestive cancer cases, the registry's aim is to produce quality data for calculating health indicators: it allows for the exact measurement of incidence trends for digestive cancers depending on the tumour site or anatomopathological type as well as patient survival. By recording incident cancer cases, in the years to come, we will be able to evaluate the efficacy of Hémoccult screening for colorectal cancer, conducted in the département since June 2004 (Cancer Screening Association ADEC 29).
	- Objectives of the registry in terms of research: The digestive tumour registry has already been used for medical theses and publications. Locally, incidence and survival studies are undertaken on a regular basis by the registry's team. We also have projects with local partners: the Brest University Hospital Centre team, University of West Brittany (prognostic factor of statin prescription on the occurrence of cancers).
	The registry's data covers the entire validated period (1984-2006) and is sent annually to the national cancer incidence base managed by the FRANCIM network, the French Institute for Public Health Surveillance and Lyon Civil Hospices. As such, we are participating in the national cancer surveillance programme and contributing to national research.
Inclusion criteria	The registry identifies all patients affected with a primitive malignant tumour, including in situ cancer of the digestive tract, liver, pancreas, biliary tracts, peritoneum and retroperitoneal tissue. The date of incidence must fall after 01 January 1984. Patients must live in Finistère and consent to the

registration of their cases.

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	Bretagne
Detail of the geography area	Finistère and bordering départements.
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	1984
Size of the database	
Size of the database (number of	
individuals)	Greater than 20 000 individuals
	Greater than 20 000 individuals 1984-2010: 27371 reported cases.
individuals) Details of the number of	
individuals) Details of the number of individuals	
individuals) Details of the number of individuals Data	1984-2010: 27371 reported cases.
individuals) Details of the number of individuals Data Database activity	1984-2010: 27371 reported cases. Current data collection Clinical data Paraclinical data Biological data

Biological data (detail)	Tumour markers.
Administrative data (detail)	Identification data.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality
Procedures	
Data collection method	Active, the registry shall research information from different sources.
Classifications used	ICD-O'3
Participant monitoring	Yes
Details on monitoring of participants	- Systematic for invasive malignant tumour and in situ carcinoma cases - vital status.
Links to administrative sources	Yes
Links to administrative sources Linked administrative sources (detail)	Yes RNIPP.
Linked administrative sources	
Linked administrative sources (detail)	
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