- Registre des cancers du sein et autres cancers gynécologiques de Côte d'Or (registre qualifié)

Head :Arveux Patrick, Centre d'Epidémiologie des Populations - Equipe d'accueil EA 4184

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General

Identification

Detailed name	Registre des cancers du sein et autres cancers gynécologiques de Côte d'Or (registre qualifié)

General Aspects

Medical area	Cancer research Gynecology/ obstetrics
Health determinants	Geography
Keywords	epidemiology

Scientific investigator(s) (Contact)

Name of the director	Arveux
Surname	Patrick
Address	1 rue du Professeur Marion ? B.P. 77980 ? 21000 DIJON
Phone	+ 33 (0)3 80 73 75 11
Email	parveux@cgfl.fr
Unit	Centre d'Epidémiologie des Populations - Equipe d'accueil EA 4184
Organization	Centre Georges-François

Collaborations

Funding

Funding status	Public
Details	Institut de veille sanitaire - InVSInstitut national du cancer - INCa

Governance of the database

Sponsor(s) or organisation(s) responsible	Centre Georges-François LECLERC (Centre Régional de lutte contre le cancer)
Organisation status	Both

Additional contact

Main features

Type of database

Type of database	Morbidity registers
Additional information regarding sample selection.	Selection of subjects having the required inclusion criteria (date of diagnosis, département, cancerous tumor, etc.) Several sources are used to identify cases: - anatomopathology laboratories (cancer prevention centers, teaching hospitals, private) - medical information departments of cancer prevention centers - Radiotherapy center (private) - Teaching hospital gynecology department - Gynecology department of hospitals based on the outskirts of the département - Gynecology department of public and private institutions in the neighboring département - Multidisciplinary Consultation Meetings of the cancer prevention center

Database objective

Main objective	The Côte d'Or registry for breast cancer and gynecological cancers, set up in 1982 and relaunched by a new team in 2005 is the only registry to specialize in these diseases in France. It now has a precise and reliable database on patients, the diseases and all treatments. In addition to the conventional work of descriptive epidemiology, useful to the health authorities for public health purposes, the registry's team develops research projects on the treatment of elderly patients, social inequalities concerning access to cancer screening on the one hand and patient treatment on the other. The setup and management of the national cohort CANTO by the registry team is a key asset for developing research projects involving population data and cohort findings.
Inclusion criteria	All cases of breast cancer - in situ and invasive - diagnosed in women living in the département at the time of diagnosis. Cases of lobular neoplasia are also listed.

	Recording of all gynecological cancers (of the uterus, ovaries, etc.) diagnosed in women living in the département at the time of diagnosis.

Population type

Age	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	Woman
Geography area	Departmental
French regions covered by the database	Bourgogne Franche-Comté
Detail of the geography area	Côte d'or

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1982

Size of the database

Size of the database (number of individuals)	[10 000-20 000[individuals
Details of the number of individuals	1982-2008: 9482 tumeurs du sein (8540 cancers infiltrants, 873 cancers in situ, 69 néoplasies lobulaires)

Data

Database activity	Current data collection
Type of data collected	Clinical data Paraclinical data Biological data Administrative data
Clinical data (detail)	Direct physical measures
Paraclinical data (detail)	diagnostic circumstances (screening, etc.)

Biological data (detail)	hormone receptors, HER2 assay
Administrative data (detail)	Identification data (last name, first name, date of birth, exact address, etc.)
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services
Care consumption (detail)	Medicines consumption

Procedures

Data collection method	Active collection - Collection of anatomocytopathological report listings then visit to each of the laboratories - Consultation of paper and/or electronic medical records
Classifications used	ICD-O-3
Participant monitoring	Yes
Details on monitoring of participants	- consultation of the medical record for the vital status or date of last visit (administrative data)- RNIPP query- letter from the physician
Links to administrative sources	Yes
Linked administrative sources (detail)	RNIPP

Promotion and access

Promotion

Link to the document	Resume des 5 publi significatives du registre.doc
Link to the document	http://tinyurl.com/PUBMED-BGCR
Description	Liste des publications dans Pubmed
Link to the document	http://tinyurl.com/HAL-BGCR
Description	Liste des publications dans HAL

Access

Terms of data access (charter for data provision, format of data, availability delay)	data collected in the Francim database (network of registries) data made available as part of collaborative

	research (by asking the manager)
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only