

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

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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)	
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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 - InVS Institut 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)
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Population covered	Sick population
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Gender	Woman
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Geography area	Departmental
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French regions covered by the database	Bourgogne Franche-Comté
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Detail of the geography area	Côte d'or
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Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1982
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Size of the database

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

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