

# - Doubs and Territoire de Belfort primitive malignant tumors register

Responsable(s) :Woronoff Anne-Sophie

Date de modification : 23/01/2015 | Version : 1 | ID : 210

## Général

### Identification

Nom détaillé	Doubs and Territoire de Belfort primitive malignant tumors register
Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.)	903417

### Thématiques générales

Domaine médical	Cancer research
Autres, précisions	cancer
Mots-clés	tumor, collection, cancer diagnoses, cancer risk, surveys

### Responsable(s) scientifique(s)

Nom du responsable	Woronoff
Prénom	Anne-Sophie
Adresse	CHU Saint-Jacques 2 place Saint-Jacques - 25030 Besançon cedex
Téléphone	03 81 21 83 12
Email	asworonoff@chu-besancon.fr
Organisme	Registre des tumeurs du Doubs et du Territoire de

### Collaborations

### Financements

Financements	Public
--------------	--------

Précisions	Le registre est financé par l'InVS, l'INCa, le CHU de Besançon et l'ARS de Franche-Comté. D'autres financements ponctuels interviennent sur projets (Ligue contre le cancer ...).
------------	-----------------------------------------------------------------------------------------------------------------------------------------------------------------------------------

## Gouvernance de la base de données

Organisation(s) responsable(s) ou promoteur

CHU de Besançon

Statut de l'organisation

Secteur Public

## Contact(s) supplémentaire(s)

## Caractéristiques

## Type de base de données

Type de base de données

Morbidity registers

Origine du recrutement des participants

A selection of health institutions and services

Le recrutement dans la base de données s'effectue dans le cadre d'une étude interventionnelle

No

Informations complémentaires concernant la constitution de l'échantillon

Selection of subjects having the required inclusion criteria.

Several sources are used:

- Anatomopathology laboratories
- Hematological cytology laboratories
- Genetic departments of training hospitals
- Medical information departments
- Health Insurance Funds
- Private-practicing and hospital clinicians (+ GPs when necessary)
- Summaries of Multidisciplinary Consultation Meetings (RCPs)
- Organized screening structure
- Haut-Rhin and Bas-Rhin Registry
- National Registry of Hematological Diseases in children
- Registry of Solid Tumors in children

## Objectif de la base de données

Objectif principal

The registry carries out a continuous and exhaustive collection of all new cancer diagnoses in the general population living in a given département. It has the twofold objective of describing and monitoring the cancer risk and conducting research from the analysis of data gathered or specific surveys.

Objectives of the Registry in terms of public health:

- Participate in the epidemiological surveillance of

cancers at local and national level under the Francim network by publishing incidence indicators by cancer location, gender, age and year of diagnosis. - Help to evaluate primary and secondary prevention initiatives (organized cancer screening), patient treatment and healthcare needs in the general population.

#### Critères d'inclusion

All primitive malignant tumors diagnosed in patients living, at the time of diagnosis, in the Doubs or Territoire-de-Belfort départements, irrespective of where they are treated, are recorded - including stratum basale skin carcinomas (specific to the Doubs Registry). In situ primitive tumors of the breast, colon-rectum, cervix, bladder and excretory ducts, as well as in situ melanomas. Benign tumors and tumors of intermediate malignancy of the central nervous system and bladder are also recorded, as are borderline tumors of the ovary.

#### Type de population

##### 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 concernée

Sick population

##### Sexe

Male  
Woman

##### Champ géographique

Departmental

##### Régions concernées par la base de données

Bourgogne Franche-Comté

##### Détail du champ géographique

Doubs (25) and Territoire de Belfort (90)

#### Collecte

##### Dates

##### Année du premier recueil

1978

##### Année du dernier recueil

2008

## Taille de la base de données

Taille de la base de données (en nombre d'individus) Greater than 20 000 individuals

Détail du nombre d'individus 1978-2008: 79100 / Doubs 2007-2008: 2312 / Territoire de Belfort

## Données

Activité de la base Data collection completed

Type de données recueillies Clinical data  
Paraclinical data  
Biological data  
Administrative data

Données cliniques, précisions Direct physical measures

Données paracliniques, précisions Any additional examination providing information for tumor coding

Données biologiques, précisions Any additional examination providing information for tumor coding

Données administratives, précisions Identification data (Last name, first name, date of birth, town of birth, address)

Existence d'une bibliothèque No

Paramètres de santé étudiés Health event/morbidity  
Health event/mortality

## Modalités

Mode de recueil des données Active or passive collection (CNIL authorization): - Receipt of anatomopathological and cytological reports, lists of hospital stays (PMSI) and chronic disease diagnoses with regard to health insurance reimbursement. Lists by encrypted electronic dispatch. - Active search for regular complementary information in medical records

Suivi des participants Yes

Détail du suivi Vital status, death, date of last news - Sample-based follow-up is carried out (High-Resolution Francim surveys). - Passive follow-up is carried out using information from the Registry (anatomocytology reports).

Appariement avec des sources administratives Yes

Sources administratives  
appariées, précisions

PMSI, Chronic disease lists, Organized screening  
structure (CNIL authorization)

## Valorisation et accès

### Valorisation et accès

#### Accès

Charte d'accès aux données  
(convention de mise à  
disposition, format de données  
et délais de mise à disposition)

Incidence data is available in aggregate form, by  
cancer location, year, gender and age. This data is  
published at regular intervals at local and national  
level (see ""Find out more"").

The data recorded in the Registry may be used for  
specific research. Project requests are analyzed by  
the Registry's Strategic and Scientific Advisory  
Board. The arrangements for cooperation must be  
defined.

Accès aux données agrégées

Free access

Accès aux données individuelles

Access on specific project only