

- Bas-Rhin Cancer Registry (Certified Registry)

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Général

Identification

Nom détaillé	Bas-Rhin Cancer Registry (Certified Registry)
Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.)	CNIL 998044.

Thématiques générales

Domaine médical	Cancer research
Mots-clés	tool, epidemiological research, assessment, morbidity, mortality, incidence, prevalence, survival

Responsable(s) scientifique(s)

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Organisme	Faculté de Médecine

Collaborations

Participation à des projets, des réseaux, des consortiums	Yes
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Précisions	European Network of Cancer Registries (ENCR), International Association of Cancer Registries (IACR) and compliance with operating rules established by the International Agency for Research on Cancer (IARC). French Network of Cancer Registries (FRANCIM Network) and collaboration with the French Institute for Public
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Health Surveillance (InVS) and National Cancer Institute (INCA), as part of the national cancer surveillance programme.

Financements

Financements Public

Précisions InVS

Gouvernance de la base de données

Organisation(s) responsable(s) ou promoteur Faculté de Médecine Starsbourg

Statut de l'organisation Secteur Public

Contact(s) supplémentaire(s)

Caractéristiques

Type de base de données

Type de base de données Morbidity registers

Informations complémentaires concernant la constitution de l'échantillon Selection of subjects meeting the required inclusion criteria. Several sources were used: - Anatomical pathology laboratories - university hospital centre - CLCC - European cancer treatment centres - hospital - private clinics - health insurance fund.

Objectif de la base de données

Objectif principal

- Registry objectives regarding public health: The registry was created in 1974 with two main aims:
 1. To create a useful tool for epidemiological research and evaluation, modelled on tools established in various other countries;
 2. To verify morbidity data of the unfavourable situation in Alsace with regards to high cancer mortality.

The Bas-Rhin cancer registry is responsible for estimating cancer incidence, prevalence and survival.

The registry also aims to participate in national and international studies in these areas: comparing incidence rates to locate different geographical regions and compare them in a database; to develop existing work to identify factors that explain variations in cancer incidence, prevalence and survival. The cancer registry is also a tool for

assessing the effectiveness of preventive measures, especially screening campaigns throughout the département for breast cancer since 1989, cervical and colon cancer since 1994 and colon and rectal cancer since 2008. As such, the trend in incidence rate can be monitored and the proportion of false negatives can be documented.

- Registry research objectives:

The registry is developing research in the field of estimating cancer incidence in geographical areas with no registry, as well as estimating survival through relative survival and healing models in order to obtain the best "current" survival estimates.

Critères d'inclusion

All diagnosed malignant or primitive in situ tumours are registered for subjects residing in the Bas-Rhin département at the time of diagnosis, regardless of their treatment location. Benign and intermediate malignant tumours in the bladder and central nervous system are also registered. Basocellular skin tumours are excluded. Multiple tumours in the same subject are registered, regardless if they occur simultaneously in paired organs or sub-locations of the same organ.

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

Alsace Champagne-Ardenne Lorraine

Détail du champ géographique

Bas-Rhin département (67).

Collecte

Dates

Année du premier recueil 1975

Taille de la base de données

Taille de la base de données (en nombre d'individus) [1000-10 000[individuals

Détail du nombre d'individus 1975-2004: 122,617 reported cases.

Données

Activité de la base Current data collection

Type de données recueillies Clinical data
Administrative data

Données cliniques, précisions Direct physical measures

Données administratives, précisions Identification data.

Existence d'une bibliothèque No

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

Modalités

Mode de recueil des données Consultation of files or clinical records on-site or electronically.

Suivi des participants Yes

Détail du suivi - Vital status - a sample follow-up is carried out. Organised as part of the "high resolution" studies conducted by the FRANCIM network, where the objective is to study the development of cases (recurrence, metastases, complications) and treatment based on one or two large anatomical localisations per year with an average rotation period of five years.

Appariement avec des sources administratives Yes

Sources administratives appariées, précisions PMSI

Valorisation et accès

Valorisation et accès

Lien vers le document <http://www.ncbi.nlm.nih.gov/pubmed?term>

Accès

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

Contact the scientist in charge.

Accès aux données agrégées

Access on specific project only

Accès aux données individuelles

Access on specific project only