EUROCARE - European cross-sectionnal study on patients with cancer : survival, prevalence and care models

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General

Identification

Detailed name	European cross-sectionnal study on patients with cancer : survival, prevalence and care models
Sign or acronym	EUROCARE

General Aspects

Medical area	Cancer research
Health determinants	Iatrogenic
Keywords	care models, Europe, cancer, survival, prevalence

Scientific investigator(s) (Contact)

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Organization	Institut national de la Santé -

Collaborations

Funding

Funding status	Mixed
Details	Italian Ministry of HealthCompagnia di San Paolo di TorinoCARIPLO foundation

Governance of the database

Sponsor(s) or organisation(s) responsible	Institut national de la Santé - Italie

Organisation status	Public
Sponsor(s) or organisation(s) responsible	Centre national d'épidémiologie, de surveillance de la santé et de la promotion - Italie
Organisation status	Public

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Repeated cross-sectional studies (except case control studies)
Database recruitment is carried out by an intermediary	An administrative base or a register
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	From 117 cancer registries in 29 European countries.

Database objective

Main objective	To provide an updated description of cancer survival time trends and differences across European countries, to measure cancer prevalence, and to study patterns of care of cancer patients.
Inclusion criteria	Patients with cancer living in Europe.

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	International
Detail of the geography area	Europe :Denmark, Finland, Iceland, Norway, Sweden, England, Ireland, Northern Ireland, Scotland, Wales, Austria, Belgium, France, Germany, Netherlands, Switzerland, Croatia, Italy, Malta, Portugal, Slovenia, Spain, Bulgaria, Czech Republic, Estonia, Latvia, Lithuania, Poland, Slovakia.

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	1989

Size of the database

Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	22 000 000

Data

Database activity	Current data collection
Type of data collected	Clinical data Administrative data
Clinical data (detail)	Direct physical measures
Administrative data (detail)	Date of birth, country of residence
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption

Procedures

Participant monitoring	No

Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://www.eurocare.it/Publications/tabid/61/Default.aspx
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientific director.
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only