FDRK - Roadmap for Colon Cancer

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General

Identification

Detailed name	Roadmap for Colon Cancer
Sign or acronym	FDRK
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL: 16/03/2006

General Aspects

Medical area	Cardiology
Health determinants	Nutrition
Others (details)	Colon cancer
Keywords	tumour progression, metastasis, primary metastatic disease, local tumour recurrence, Health episodes, death

Scientific investigator(s) (Contact)

Name of the director	Abenhaim
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Unit	INSERM U657
Organization	LASER

Collaborations

Funding

Funding status	Mixed

Details	APPEL D'OFFRE ROCHE

Governance of the database

Sponsor(s) or organisation(s) responsible	Laser
Organisation status	Private

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Cohort study
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Inclusion method: Prospective Other bodies active in creating this cohort: CHU, CHG, CANCER TREATMENT CENTRES, PRIVATE NON-PROFIT HOSPITALS PARTICIPATING IN PUBLIC HOSPITAL SERVICE (PSPH) AND PRIVATE CLINICS.

Database objective

Main objective	General Objective 1. To describe treatment methods in France for 1,360 patients with colorectal cancer diagnosed between 2006 and 2008 2. To describe overall survival at 5 years according to initial treatment methods Secondary objectives: - To describe target populations and chemotherapy usage patterns; - To describe usage patterns for different centres; - To describe the "outcomes" after 12, 36 and 60 months: progression-free survival and quality of life.
Inclusion criteria	- Primary adenocarcinoma of the colon or upper rectum with histological confirmation (> 10 cm from the anal margin, subperitoneal upper rectum included); - Incident cases (histological diagnosis of colon cancer)

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	Male Woman
Geography area	National
Detail of the geography area	Multicentric cohort throughout France (97 centres)

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	06/2006
Date of last collection (YYYY or MM/YYYY)	05/2013

Size of the database

Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	1360

Data

Database activity	Data collection completed
Type of data collected	Clinical data Declarative data
Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Paper self-questionnaire
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Self-administered questionnaire: Entry from a paper questionnaire (Manual input) Clinical examination: Handwritten (Manual input)
Participant monitoring	Yes
Details on monitoring of participants	Follow-up duration: 5 years
Links to administrative sources	Yes
Linked administrative sources (detail)	Database(s) used: CépiDc

Promotion and access

Promotion

Link to the document	http://www.nature.com/bjc/journal/v81/n1/abs/6690651a.html
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Data may be used by academic teams: access for all doctoral students accepted into a high-level team (INSERM). Data may be used by industrial teams: Access to aggregate and anonymised data at ROCHE
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only