

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)

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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)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	National
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Detail of the geography area	Multicentric cohort throughout France (97 centres)
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Data collection

Dates

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

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

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