- CHONSOL

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Last update : 08/19/2014 | Version : 1 | ID : 8130

General

Identification

Detailed name	CHONSOL
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL

General Aspects

Medical area	Cancer research Traumatology
Health determinants	Iatrogenic
Keywords	Solitary chondroma, chondrosarcoma, molecular markers, home, prognostic factors of onset., RESOS

Scientific investigator(s) (Contact)

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Unit	Service de chirurgie orthopédique et traumatologique du CHU Nantes
Organization	CHU

Collaborations

Funding

Funding status	Public

Details	Recherche de financements en cours
Governance of the database	
Sponsor(s) or organisation(s) responsible	CHU Nantes
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Longitudinal study (except cohorts)
Database recruitment is carried out by an intermediary	A selection of health institutions and services An administrative base or a register
Database recruitment is carried out as part of an interventional study	No
Database objective	
Main objective	The main objective of the study is to determine the incidence of chondrosarcoma at 5 years in a population presenting with solitary chondroma. The secondary objectives of the study are: - To determine the incidence of chondrosarcoma at 10 years a population presenting with solitary chondroma -To describe solitary chondroma diagnosis and follow-up modalities - To describe indications for biopsy, curettage or resection - To describe the level of pain and overall quality of life for patients with solitary chondroma as well as the impact of biopsy, curettage or resection -To describe the demographic, clinical, radiological, biological, molecular and genetic factors associated with, or predictive of, sarcomatous change. - To describe the demographic, clinical, radiological, biological, molecular and genetic factors predicting change (improvement or deterioration) in the overall quality of life and level of pain in patients with solitary chondroma - To evaluate the iatrogenic risks of biopsy, curettage or resection of solitary chondroma
Inclusion criteria	Inclusion criteria are: -Patients over 18 years of age

	-Patients with a single cartilaginous non-operative tumour -Patients fulfilling radiographic, scintigraphic, CT and/or MRI criteria for solitary chondroma -Patients who agreed to participate in the study and proposed follow-up and who have signed the informed consent for biocollection.
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	12 participating research centres (RESOS Network)
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	01/2015
Date of last collection (YYYY or MM/YYYY)	2030
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	200
Data	
Database activity	Current data collection
Type of data collected	Clinical data Declarative data Paraclinical data Biological data
Clinical data (detail)	Direct physical measures Medical registration

Declarative data (detail)	Paper self-questionnaire
Paraclinical data (detail)	Radiology
Biological data (detail)	Tissue, serum, DNA
Presence of a biobank	Yes
Contents of biobank	Serum Tissues DNA
Details of biobank content	Sample freezing will be carried out for storage at -80°C. Freezing multiple small quantities (a few milligrams) is preferred. Blood samples will be delivered and stored in biological resource centres and/or tumour banks and/or research units associated with the involved sites to undergo initial treatment for conservation.
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services Quality of life/health perception
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption

Procedures

Participant monitoring	Yes
Details on monitoring of participants	Follow-up over 10 years
Links to administrative sources	No

Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only