HOPE-EPI - HOPE-Epidemiology - Epidemiological Research in Pediatric Hemato-Oncology

Head :Clavel Jacqueline, UMRS-1018 Equipe 6

Last update : 08/07/2014 | Version : 2 | ID : 3121

General	
Identification	
Detailed name	HOPE-Epidemiology - Epidemiological Research in Pediatric Hemato-Oncology
Sign or acronym	HOPE-EPI
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL
General Aspects	
Medical area	Cancer research Hematology Pediatrics
Health determinants	Climate Genetic Geography Iatrogenic Medicine Pollution Social and psychosocial factors
Keywords	national tools, research infrastructure, health events, pediatric oncology, environnemental and genetic risk factors, social and territorial determinants, iatrogenic effects of treatments
Scientific investigator(s)	determinants, iatrogenic effects of treatments

Scientific	investigator(s)
(Contact)	

Name of the director	Clavel
Surname	Jacqueline
Address	UMRS-1018 Equipe 6, INSERM, Université Paris Sud, 16 avenue Paul Vaillant Couturier F-94807 Villejuif Cedex
Phone	0145595038

Fmail jacqueline.clavel@inserm.fr Unit UMRS-1018 Equipe 6 Organization **INSERM** Collaborations Participation in projects, Yes networks and consortia **Funding** Funding status Public **Details** ANR "Investissements d'avenir - Grand emprunt" Governance of the database Sponsor(s) or organisation(s) Inserm responsible Organisation status **Public** Additional contact Main features Type of database Type of database Study databases Study databases (details) Cohort study Database recruitment is carried A population file out by an intermediary Database recruitment is carried No out as part of an interventional study Database objective Main objective The objective of our project is to develop, from the existing national tools, a national research infrastructure pooling the work to collect data and validate information and health events in the service

of an epidemiological research program in the field

- the environnemental and genetic risk factors for

of pediatric oncology addressing:

childhood cancer.

	 the differences in management and their social and territorial determinants, the iatrogenic effects of treatments, and more generally, the state of health in the short-, intermediate- and long-term of people experiencing childhood cancer and the determinants of their quality of life. The project, closely connected with the pediatric oncology research of other disciplines, has, as a secondary objective, contributing to the sharing of data across disciplines.
Inclusion criteria	Children aged less than 18 years with cancer
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)
Population covered	Sick population
Gender	Male Woman
Geography area	National
Detail of the geography area	Metropolitan France
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2010
Date of last collection (YYYY or MM/YYYY)	2020
Size of the database	
Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	2010 : 170002020 : 35000
Data	

Type of data collected	Clinical data Paraclinical data Biological data Administrative data
Clinical data (detail)	Direct physical measures Medical registration
Details of collected clinical data	Diagnostic: Date of diagnosis, Histology, Cytology, Cytogenetics, Immunophenotype, Molecular biology, Topography, Staging (/classification) Coding (ICD-O and ICCC) Healthcare pathway. Traitment: Risk classification, Date treatment started, Protocol arm, Chemotherapy (date, drugs, cumulative doses), Surgery (date, type and site), Radiotherapy (date, type, site and dose), Stem cell transplantation (date, type and source)
Paraclinical data (detail)	Medical imaging
Biological data (detail)	
Administrative data (detail)	Location since birth, basic information,
Presence of a biobank	Yes
Contents of biobank	Whole blood
Details of biobank content	
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
Links to administrative sources	Yes
Linked administrative sources (detail)	SNIIR-AM
Promotion and access	
Promotion	

Link to the document	<u>Hope-Epi.pdf</u>
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientific board
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only