NETSARC - NATIONAL PROSPECTIVE COHORT ON SARCOMAS/GIST/DESMOID AND CONNECTIVE TISSUE TUMOURS

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General	
Identification	
Detailed name	NATIONAL PROSPECTIVE COHORT ON SARCOMAS/GIST/DESMOID AND CONNECTIVE TISSUE TUMOURS
Sign or acronym	NETSARC
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	Accord CNIL DR-2013-383, CCTIRS 16/09/2010
General Aspects	
Medical area	Cancer research
Health determinants	Genetic Geography Iatrogenic
Keywords	Sarcoma, rare cancers, network, clinical, biological samples, ISKS project, genetic, treatment
Scientific investigator(s) (Contact)	
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Unit	INSERM U570
Organization	Centre Léon Bérard

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Unit	EAM Santé Individu Société 4129 Universités Lyon 1/ Lyon 2 / Lyon 3
Organization	Centre Léon Bérard
Collaborations	
Participation in projects, networks and consortia	Yes
Funding	
Funding status	Mixed
Details	Pour le projet ISKS spécifique : - INFO SARCOMES (association de patients)- LIDDY SHRIVER (association de patients)Recherche de financements additionnels
Governance of the database	
Sponsor(s) or organisation(s) responsible	Centre Léon Bérard
Organisation status	Private
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
Database recruitment is carried out as part of an interventional study	No

Additional information regarding sample selection.	 patients with sarcoma or connective tissue tumours (all ages and histologies) discussed in multidisciplinary meetings (RCP) in regional centres of expertise specialising in the treatment of sarcomas
Database objective	
Main objective	The objectives of this cohort are: real-time integration of patients into a database- to generate real-time national and regional descriptive data for all patients with sarcoma - to provide national indicators of patient care and equal access to care and innovative treatments - to improve knowledge of rare tumours - to investigate the link between sarcomas and genetics.
Inclusion criteria	 patients with sarcoma or connective tissue tumours (all ages and histologies) discussed in multidisciplinary meetings (RCP) in regional centres of expertise specialising in the treatment of sarcomas
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	National
Detail of the geography area	The national NETSARC clinical network is composed of 28 regional centres of expertise specialising in sarcoma treatment and research.
Data collection	
Dates	
Date of first collection (YYYY or	01/2010

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Presence of a biobank Yes Contents of biobank DNA	Biological data (detail)	DNA
Contents of biobank DNA	Administrative data (detail)	
	Presence of a biobank	Yes
Details of biobank content DNA	Contents of biobank	DNA
	Details of biobank content	DNA

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
Procedures	
Data collection method	Standardised reports from regional centres specialising in the treatment of sarcomas included in the network.
Participant monitoring	Yes
Details on monitoring of participants	Frequent return to medical record while patient is undergoing treatment and follow-up in the centre. Return every 2 years during follow-up outside centres via mail addressed to correspondents.
Links to administrative sources	No
Promotion and access	
Promotion	
Promotion Link to the document	<u>NetSarc.pdf</u>
	<u>NetSarc.pdf</u>
Link to the document	NetSarc.pdf Shared database is accessible online and developed by two network I.T. professionals. Data is hosted by a professional hosting provider. Users can connect with a unique password. Different rights are allocated per profile (clinician, clinical research associate, quality manager, administrator). Website terms of use must be previously signed. Non-identifiable patient data (HASH system is being put in place). A scientific committee consisting of representatives clinicians and pathologists centres to large volume of patients in the network is responsible for controlling, analysing and authorising requests for studies from the shared data base.
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