

3GENSSC - GFRS- 3 GENERATIONS-SCLERODERMA

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General

Identification

Detailed name GFRS- 3 GENERATIONS-SCLERODERMA

Sign or acronym 3GENSSC

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation N°de protocole RBM-04-60 (20-09-2004) et collection N°DC-2008-327 (2008)

General Aspects

Health determinants Genetic

Keywords HLA genotyping, family members, compatibility, microchimerism, autoimmunity markers

Scientific investigator(s) (Contact)

Name of the director Lambert

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UMRs 1097

Organization INSERM - Institut National de la Santé et de la
Recherche

Name of the director Farge

Surname Dominique

Collaborations

Funding

Funding status Mixed

Details PRO-A INSERM

Governance of the database

Sponsor(s) or organisation(s) responsible INSERM - Institut National de la Santé et de la Recherche Médicale

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

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. New subjects are enrolled from active patient files monitored by recruitment centres

Database objective

Main objective To form the first French cohort of 3 generations of subjects who have scleroderma (SSc). To carry out HLA (human leukocyte antigen) genotyping for all subjects and family members so as to understand the relationship between HLA compatibility between SSc patients with and members of their family and to further study the role of microchimerism and maternal-foetal compatibility. To continue studies previously undertaken on the preponderance of autoimmune disease in women and to research new markers of SSc on a larger number of subjects.

Inclusion criteria Women whose parents are still alive and whose children wish to participate in the study.

Population type	
Age	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	France
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	10/2004
Size of the database	
Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	400
Data	
Database activity	Current data collection
Type of data collected	Clinical data
Clinical data (detail)	Direct physical measures
Presence of a biobank	Yes
Contents of biobank	Whole blood
Details of biobank content	Whole blood from subjects will be collected in EDTA vacutainer tubes. An aliquot of 350 ul will be retained and DNA will be extracted using a kit (EZ1 DNA Blood Kit, Qiagen, Hilden, Germany) on a BIOROBOT EZ1 according to instructions.

Health parameters studied	Health event/morbidity Health event/mortality
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Procedures

Data collection method	Family members will be contacted by post by a Clinical Research representative (research engineer). To facilitate geographic difficulties for dispersed families, mouth wash will be sent to collect buccal cells for DNA extraction, or an appointment will be made in accordance with the subject in an analysis laboratory close to the subject's home.
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Participant monitoring	Yes
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Links to administrative sources	No
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Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	To be defined
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Access to aggregated data	Access on specific project only
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Access to individual data	Access on specific project only
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