OVR - Familial Retinal Vein Occlusion

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General

Identification

Detailed name	Familial Retinal Vein Occlusion
Sign or acronym	OVR
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	banque d'ADN de l'institut de la vision

General Aspects

Medical area	Cardiology Ophthalmology
Health determinants	Genetic Occupation
Others (details)	Familial retinal vein occlusion
Keywords	urban population, Health episodes, rural population, hospitalisation, health system

Scientific investigator(s) (Contact)

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Unit	CIC 503 INSERM
Organization	INSERM - Institut National de la Santé et de la Recherche

Collaborations

Funding	
Funding status	Public
Details	PHRC
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)	Not-repeated cross-sectional studies (except case control studies)
Database recruitment is carried out by an intermediary	An administrative base or a register
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Retrospective Number of required subjects: [100-500]
Database objective	
Main objective	General objective: to phenotype forms of familial retinal vein occlusion Secondary objective: to collect DNA
Inclusion criteria	At least 3 cases of retinal vein occlusion (RVO) in the immediate family (two cases if one occurs before the age of 30).
Population type	
Age	Great age (80 years and more)
Population covered	General population
Gender	Male Woman

Geography area	Local
Detail of the geography area	Multicentric cohort throughout France (2 centres)

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	01/2000

Size of the database

Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	40 (03/05/2013)

Data

Database activity	Data collection completed
Type of data collected	Clinical data Declarative data Paraclinical data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Face to face interview
Paraclinical data (detail)	Imaging
Biological data (detail)	Type of samples taken: DNA
Presence of a biobank	Yes
Contents of biobank	DNA
Details of biobank content	DNA bank
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Self-administered questionnaire: input from a paper questionnaire; Interviews: direct input; Clinical examinations: handwritten Biological analysis: handwritten

Participant monitoring	No
Details on monitoring of participants	(Indefinite duration)
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://www.ncbi.nlm.nih.gov/pubmed/17173011
Link to the document	http://www.ncbi.nlm.nih.gov/pubmed/20703046
Link to the document	http://www.ncbi.nlm.nih.gov/pubmed/19701812

Access

Terms of data access (charter for data provision, format of data, availability delay)	To be decided if data may be used by academic teams To be decided if data may be used by industrial teams
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only