

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
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Promotion and access

Promotion

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