

# PROGENE - National Study on Prostate Cancer

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## General

### Identification

Detailed name National Study on Prostate Cancer

Sign or acronym PROGENE

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL: 27/06/2001

### General Aspects

Medical area Cancer research

Health determinants Genetic  
Social and psychosocial factors

Others (details) Prostate cancer

Keywords anatomopathological, data, weight, height, Health episodes, clinical, family history

### Scientific investigator(s) (Contact)

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Unit CEREPP

Organization APHP

### Collaborations

### Funding

Funding status	Mixed
Details	AP-HP CEREPP
Governance of the database	
Sponsor(s) or organisation(s) responsible	CEREPP
Organisation status	Private
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Cohort study
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.	Inclusion method: Prospective Other bodies active in creating this cohort: CHU, CHG, INDEPENDENT PHYSICIANS, CNAM
Database objective	
Main objective	General objective: To initiate a collection of informative families with at least one member with prostate cancer and apparently health subjects. Secondary objectives: - To identify genes for predisposition and susceptibility to prostate cancer - To identify genetic markers associated with recurrence after treatment - To develop a genetic screening test for these pathologies
Inclusion criteria	Men with histologically verified prostate cancer Healthy subjects with PSA levels below 4 ng/ml
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years)

Population covered	Sick population
Gender	Male
Geography area	National
Detail of the geography area	France
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	06/2001
Size of the database	
Size of the database (number of individuals)	[1000-10 000[ individuals
Details of the number of individuals	5965
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Declarative data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Paper self-questionnaire
Biological data (detail)	Type of samples taken: Blood and/or saliva and/or urine
Presence of a biobank	Yes
Contents of biobank	Serum DNA
Details of biobank content	Serum bank, DNA bank
Health parameters studied	Health event/morbidity Health event/mortality
Procedures	
Data collection method	Self-administered questionnaire: Entry from a paper

questionnaire (Manual input) Clinical examination:  
Handwritten (Manual input) Biological Analysis:  
Handwritten (Manual input)

Participant monitoring Yes

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/?term>

### Access

Terms of data access (charter for data provision, format of data, availability delay) Data may be used by academic teams. Access based on contract collaboration. Data may not be used by industrial teams

Access to aggregated data Access on specific project only

Access to individual data Access on specific project only