COREVE - Contraception and recurrent venous event

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General

Identification

Detailed name	Contraception and recurrent venous event
Sign or acronym	COREVE
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL

General Aspects

Health determinants	Addictions Occupation
Others (details)	venous thrombosis
Keywords	Evaluation, types of hormonal contraception, recurrence, impact, risk

Scientific investigator(s) (Contact)

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Unit	INSERM U1018 CENTRE DE RECHERCHE EN ÉPIDÉMIOLOGIE ET SANTÉ DES POPULATIONS
Organization	INSERM

Collaborations

Funding

Funding status	Mixed

Details	INSERM, PIERRE FABRE MEDICAMENT
Governance of the database	
Sponsor(s) or organisation(s) responsible	INSERM - Institut National de Santé et Recherche Médicale
Organisation status	Public
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.	Prospective Other bodies active in creating this cohort: PIERRE-YVES SCARABIN; GENEVIEVE PLU BUREAU; JACQUELINE CONARD; MARIE HELENE HORELLOU.
Database objective	
Main objective	General objective: to study whether or not different forms of hormonal contraception contribute to the risk of venous thrombosis recurrence for women at high risk of venous thromboembolism.
Inclusion criteria	Women aged between 18 and 45 years attending Hôtel-Dieu de Paris Hospital for the first documented instance of a venous thrombotic event (Doppler ultrasound, computed tomography angiography scan, angiogram, etc.) and/or thrombophilia (factor V Leiden, prothrombin 20210a mutation, ATIII deficiency, PC, PS).
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years)
Population covered	Sick population

Gender	Woman
Geography area	Local
French regions covered by the database	Île-de-France
Detail of the geography area	Paris and surrounding region

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	2010

Size of the database

Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	2500

Data

Database activity	Current data collection
Type of data collected	Clinical data Declarative data Paraclinical data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Paper self-questionnaire Face to face interview
Paraclinical data (detail)	Medical Imaging
Biological data (detail)	Type of samples collected: Systematic testing for thrombophilia as part of a specialist consultation.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Self-administered questionnaire: double data entry

	from paper questionnaire Interview: double data entry from paper questionnaire Clinical Examinations: handwritten with double data entry
Participant monitoring	Yes
Links to administrative sources	No
Promotion and access	
Promotion **Access**	
Terms of data access (charter for data provision, format of data, availability delay)	Data may be used by academic teams 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