

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
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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