

SNAC - Study of NorpregnAnes Coagulation

Head :

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General

Identification

Detailed name	Study of NorpregnAnes Coagulation
Sign or acronym	SNAC
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CCPPRB Bicêtre (21/12/2005), CNIL (19/12/2005)

General Aspects

Medical area	Biology Endocrinology and metabolism Internal medicine
Pathology (details)	Hormone therapy for menopause
Health determinants	Medicine
Keywords	Oestrogen, progestin, menopause, haemostasis, thrombosis

Scientific investigator(s)
(Contact)

Collaborations

Funding

Funding status	Public
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Details	Institut National de la Santé et de la Recherche Médicale
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Governance of the database

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	A population file
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Samples selected on a voluntary basis
Database objective	
Main objective	Impact of different types of HRT on activated protein C resistance and haemostatic parameters.
Inclusion criteria	- Women - between 45 and 70 years of age - menopausal
Population type	
Age	Adulthood (45 to 64 years)
Population covered	General population
Gender	Woman
Geography area	Regional
French regions covered by the database	Île-de-France
Detail of the geography area	Centre d'Investigations Préventives et Cliniques de Paris: SS medical examination centre
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2006
Date of last collection (YYYY or MM/YYYY)	2007
Size of the database	
Size of the database (number of	< 500 individuals

individuals)

Details of the number of individuals

300- 100 femmes non traitées/women no treated-
100 femmes estrogènes transdermiques +
progestérone- 100 femmes estrogènes
transdermiques + NOMAC

Data

Database activity

Data collection completed

Type of data collected

Clinical data
Declarative data
Paraclinical data
Biological data

Clinical data (detail)

Direct physical measures

Declarative data (detail)

Face to face interview

Paraclinical data (detail)

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Biological data (detail)

- blood - DNA

Presence of a biobank

Yes

Contents of biobank

Serum
Plasma
DNA

Details of biobank content

500 microL plasma aliquots collected with citrate,
heparin or EDTA

Health parameters studied

Others

Other (detail)

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Procedures

Data collection method

Standardised questionnaire

Participant monitoring

No

Links to administrative sources

No

Promotion and access

Promotion

Access

Terms of data access (charter

Contact the scientist in charge

for data provision, format of data, availability delay)

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only