

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
Details	Institut National de la Santé et de la Recherche Médicale
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)	
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)	
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