FIV-ABM - National Registry on In-Vitro Fertilization Attempts

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General	
Identification	
Detailed name	National Registry on In-Vitro Fertilization Attempts
Sign or acronym	FIV-ABM
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL approval.
General Aspects	
Medical area	Biology
Pathology (details)	medically-assisted procreation
Health determinants	latrogenic
Keywords	IVF, MAP, registry, sterility, in-vitro fertilisation, medically-assisted procreation, infertility
Scientific investigator(s) (Contact)	
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Organization	Agence de la
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Unit	Direction Procréation Embryologie Génétique humaine
Organization	Agence de la
Collaborations	
Funding	
Funding status	Public
Details	Ministry of Health and Social Affairs
Governance of the database	
Sponsor(s) or organisation(s) responsible	Agence de la Biomédecine
Organisation status	Public
Additional contact	
Additional contact Main features	
Main features	Morbidity registers
Main features Type of database	Morbidity registers Comprehensive
Main features Type of database Type of database Additional information regarding	

	attempt, throughout different centres if required, and to provide recommendations in order to improve clinical practice for the patient; To monitor pregnancy and to assess the child's health at birth through an after-birth study.
Inclusion criteria	 IVF attempts (with or without ICSI) and the transfer of frozen embryos; Intrauterine insemination or simple ovarian stimulation is not included.
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years)
Population covered	Sick population
Gender	Woman
Geography area	National
Detail of the geography area	France
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2008
Date of last collection (YYYY or MM/YYYY)	2012
Size of the database	
Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	50,000 per year.
Data	
Database activity	Current data collection
Type of data collected	Clinical data Declarative data Biological data
Clinical data (detail)	Direct physical measures Medical registration

Details of collected clinical data	patient health status, sterility assessment.
Declarative data (detail)	Paper self-questionnaire
Details of collected declarative data	Smoking.
Biological data (detail)	Sperm analysis.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health care consumption and services Others
Care consumption (detail)	Hospitalization Medical/paramedical consultation
Other (detail)	Labour/births.
Procedures	
Data collection method	Computer data is exchanged between MAP clinical biology centres.
Quality procedure(s) used	A data validation plan listing all quality checks for data from each progress report is established every year. This plan aims to define checks that may identify missing or inconsistent data. A data clarification request is systematically sent to centres so they may correct or clarify missing or inconsistent data.
Participant monitoring	Yes
Details on monitoring of participants	 Follow-up until the end of pregnancy. Child's health at birth - follow-up of MAP treatment plan for women: sequence of various attempts.
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Anonymous data, statistics available on website. Contact the scientist in charge for further information.
Access to aggregated data	Free access