

PREG-Med - Case-Control Study on the Impact of Pregnancy on Prescribing Psychotropic Medication

Head :Apter Gisèle, Research Unit for Psychiatry and Psychopathology
Becquemont Laurent
Devouche Emmanuel

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General	
Identification	
Detailed name	Case-Control Study on the Impact of Pregnancy on Prescribing Psychotropic Medication
Sign or acronym	PREG-Med
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL
General Aspects	
Medical area	Geriatrics Psychology and psychiatry
Health determinants	Intoxication Lifestyle and behavior Medicine
Keywords	psychotropic medication prescription; post-partum; antenatal; pregnancy
Scientific investigator(s) (Contact)	
Name of the director	Apter
Surname	Gisèle
Phone	+33 (0)1 46 74 33 99
Email	gisele.apter@gmail.com
Unit	Research Unit for Psychiatry and Psychopathology
Organization	Erasme
Name of the director	Becquemont

Surname	Laurent
Phone	+33 (0)1 45 21 27 22
Organization	Bicêtre Hospital
Name of the director	Devouche
Surname	Emmanuel
Phone	+33 (0)1 55 20 59 33
Email	emmanuel.devouche@parisdescartes.fr
Organization	Institute of Psychology
Collaborations	
Funding	
Funding status	Public
Details	Erasme, Institute of Psychology
Governance of the database	
Sponsor(s) or organisation(s) responsible	Erasme
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Case control study
Database recruitment is carried out by an intermediary	An administrative base or a register
Base or register (detail)	Member of a social security scheme
Database recruitment is carried out as part of an interventional study	No
Database objective	

Main objective	To determine if and how the prescription of psychotropic medication changes during pregnancy.
Inclusion criteria	Women covered by a social security scheme.
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years) Elderly (65 to 79 years) Great age (80 years and more)
Population covered	General population
Gender	Woman
Geography area	National
Detail of the geography area	France
Data collection	
Dates	
Size of the database	
Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	87,213 (pregnant women); 87,213 controls
Data	
Database activity	Data collection completed
Type of data collected	Administrative data
Administrative data (detail)	Prescription of psychotropic medication
Presence of a biobank	No
Health parameters studied	Health event/morbidity
Procedures	
Participant monitoring	Yes
Details on monitoring of participants	4 months before pregnancy; 9 months during pregnancy and 4 months after pregnancy.

Links to administrative sources	Yes
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Linked administrative sources (detail)	CRAMIF database
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Promotion and access

Promotion

Link to the document	http://www.journal-therapie.org/articles/therapie/abs/first/therapie140012/therapie140012.html
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Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge (risk that data may be removed after two years due to lack of archiving space in a rolling system).
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Access to aggregated data	Access on specific project only
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Access to individual data	Access on specific project only
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