METADAP - Do Antidepressants Induce Metabolic Syndromes?

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General

Identification

Detailed name	Do Antidepressants Induce Metabolic Syndromes?
Sign or acronym	METADAP
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL

General Aspects

Medical area	Endocrinology and metabolism
Health determinants	Intoxication
Others (details)	Metabolic syndromes
Keywords	criteria, definition, Health episodes, pharmacology, side effects

Scientific investigator(s) (Contact)

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Unit	INSERM U669 CHU DE BICETRE - SERVICE DE PSYCHIATRIE
Organization	INSERM - Institut National de la Santé et de la Recherche

Collaborations

Funding	
Funding status	Public
Details	Ministère de la Santé (PHRC NATIONAL)
Governance of the database	
Sponsor(s) or organisation(s) responsible	AP-HP
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 Inclusion cut-off date: 01/11/2011
Database objective	
Main objective	General objective: to assess the impact of antidepressants in terms metabolic syndromes. Secondary objective: to identify clinical, genetic and biological predictive factors concerning the onset of metabolic syndromes.
Inclusion criteria	Depressed patients requiring antidepressant treatment
Population type	
Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years)
Population covered	Sick population

Gender	Male Woman
Geography area	National
Detail of the geography area	Multicentric cohort throughout France (6 centres): 3 centres in the Paris region and 3 in the province.

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	11/2007

Size of the database

Size of the database (number of individuals)	< 500 individuals
Details of the number of individuals	248

Data

Database activity	Data collection completed
Type of data collected	Clinical data Declarative data Biological data
Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Paper self-questionnaire
Biological data (detail)	Type of samples taken: glucose level, cholesterol level, triglyceride level
Presence of a biobank	Yes
Contents of biobank	Serum Plasma DNA
Details of biobank content	Serum bank, plasma bank, DNA bank
Health parameters studied	Health event/morbidity Health event/mortality

Procedures

Data collection method	Self-administered questionnaire: direct input Interview: direct input Clinical examinations: direct input Biological analysis: direct input
Participant monitoring	Yes
Details on monitoring of participants	Follow-up duration: 6 months
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://www.ncbi.nlm.nih.gov/pubmed/18707937

Access

Terms of data access (charter for data provision, format of data, availability delay)	To be decided if data may be used by academic teams To be decided if 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