

# SurPass - Family, social and professional disability in major depressive disorder patients starting a treatment with antidepressant

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## General

### Identification

Detailed name	Family, social and professional disability in major depressive disorder patients starting a treatment with antidepressant
Sign or acronym	SurPass
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	Demande d'autorisation CNIL n°909245

### General Aspects

Medical area	Psychology and psychiatry
Health determinants	Occupation Social and psychosocial factors
Others (details)	Depression
Keywords	Antidepressants, functional effect, psychiatry

### Scientific investigator(s) (Contact)

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Organization	Lundbeck

## Collaborations

### Funding

Funding status Private

Details Lundbeck SAS

### Governance of the database

Sponsor(s) or organisation(s) responsible Lundbeck SAS

Organisation status Private

### Additional contact

### Main features

#### Type of database

Type of database Study databases

Study databases (details) Longitudinal study (except cohorts)

Database recruitment is carried out by an intermediary A selection of health care professionals

Database recruitment is made on the basis of: Medication(s) taken

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Psychiatrists selection: mailing of invitation letter sent to the whole liberal or mixed French psychiatrists. Patients selection: the first 5 patients seen by the psychiatrist and meeting inclusion criteria.

#### Database objective

Main objective Identify, in a cohort of major depressive disorder patients starting a treatment with any antidepressant to be followed for 6 months in an outpatient psychiatric setting, the predictive factors of an evolution of their family, social and professional disability and to establish a typology of patients according to their evolutive profile at 6 months after treatment initiation.

Inclusion criteria	18 years old or more patients who suffer from a major depressive episode according to DSM-IV, starting an antidepressants treatment in an outpatient psychiatric setting; patients which can be followed for 6 months; patients able to communicate and self-evaluate
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## 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)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	National
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Detail of the geography area	Metropolitan France
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## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY)	2010
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Date of last collection (YYYY or MM/YYYY)	2011
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### Size of the database

Size of the database (number of individuals)	[1000-10 000[ individuals
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Details of the number of individuals	4 300
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### Data

Database activity	Data collection completed
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Type of data collected	Clinical data Declarative data
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Clinical data (detail)	Direct physical measures
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Declarative data (detail)	Paper self-questionnaire
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Presence of a biobank	No
Health parameters studied	Health event/morbidity Quality of life/health perception
<b>Procedures</b>	
Data collection method	Data collected during spontaneous patients visits at inclusion and then at about 2 and 6 months. Data collected during each visit consists into a Sheehan disability scale at inclusion and at 6 months visits, filled by the patients, and a MADRS scale (depression severity) filled by the investigator. Patients meeting eligibility criteria and not included in the study are inscribed into a non-inclusion register which will allow to verify the representativeness of population included in the study. This register needs to be completed until the effective inclusion of 4 to 5 patients in the cohort.
Participant monitoring	Yes
Details on monitoring of participants	6 months
Links to administrative sources	No
<b>Promotion and access</b>	
<b>Promotion</b>	
<b>Access</b>	
Terms of data access (charter for data provision, format of data, availability delay)	Publication expected in a journal with reading committee
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only