

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
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Health parameters studied	Health event/morbidity Quality of life/health perception
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Procedures

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.
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Participant monitoring	Yes
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Details on monitoring of participants	6 months
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Links to administrative sources	No
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Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	Publication expected in a journal with reading committee
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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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