

# DéPasS - DEPRESSION ? ANXIETY AND DISABILITY IN A COHORT OF PATIENTS IN GENERAL MEDICINE PRACTICE

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## General

### Identification

Detailed name  
DEPRESSION ? ANXIETY AND DISABILITY IN A COHORT OF PATIENTS IN GENERAL MEDICINE PRACTICE

Sign or acronym  
DéPasS

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation  
CNIL n° 909006

### General Aspects

Medical area  
General practice  
Psychology and psychiatry

Keywords  
depression, anxiety, functional disability, Sheehan, general practice, antidepressant

### Scientific investigator(s) (Contact)

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Lundbeck

### Collaborations

### Funding

Funding status  
Private

Details	Lundbeck SAS Laboratory
<b>Governance of the database</b>	
Sponsor(s) or organisation(s) responsible	Lundbeck SAS
Organisation status	Private
<b>Additional contact</b>	
<b>Main features</b>	
<b>Type of database</b>	
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.	Selection of general practitioners: Study investigators were randomly selected from a list of 30 000 general practitioners, which in turn was derived from an exhaustive list of all general practitioners licensed by the national medical association in France and in active practice (CEGEDIM registry). All physicians were contacted by mail and invited to participate in the study. Selection of patients: each participating GP included around four eligible patients who spontaneously consulted for an anxiety or mood disorder during the three months following receipt of the study materials.
<b>Database objective</b>	
Main objective	To evaluate the change in self-reported personal, professional and social disability in patients with anxious or mood disorders three months after initiating antidepressant treatment.
Inclusion criteria	Patient of at least 18 years of age Patient having at least one of the 5 following

diagnoses:  
 major depressive episode (i.e. characterized)  
 Generalized anxiety disorder  
 Social anxiety disorder  
 Panic disorder with or without agoraphobia  
 Obsessive compulsive disorders  
 Patient initiating a new treatment with an  
 antidepressant  
 Patient being able to communicate and to evaluate  
 his quality of life  
 Patient having received written information on the  
 use of the medical data concerning himself within  
 the framework of the study.

## 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)	2009
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Date of last collection (YYYY or MM/YYYY)	2010
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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	8 029
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### Data

Database activity	Data collection completed
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Type of data collected	Clinical data Declarative data
Clinical data (detail)	Direct physical measures
Declarative data (detail)	Paper self-questionnaire
Presence of a biobank	No
Health parameters studied	Health event/morbidity Quality of life/health perception

## Procedures

**Data collection method**

Data collection was carried out at each visit on a voluntary and anonymous basis. Data were collected at the index consultation (inclusion visit) and at two follow-up consultations about 6 and 12 weeks after the index consultation. A window of  $\pm 2$  weeks was considered acceptable for the 6-week visit and a window of  $\pm 3$  weeks for the 12-week visit. These visits were programmed as part of the routine follow-up of the patients, and no additional protocol-specified study visit was imposed. Patients' disability was evaluated with the Sheehan Disability Scale (SDS) at baseline and after six and twelve weeks. The patient's symptomatology was also filled in by the patient himself using the HAD self-questionnaire at each of three collection times scheduled in the protocol. Other data were collected by the physician himself.

**Participant monitoring** Yes

**Details on monitoring of participants** 3 months

**Links to administrative sources** No

## Promotion and access

### Promotion

**Link to the document** <http://tinyurl.com/Hal-DEPASS>

**Description** List of publications in HAL

**Link to the document** <http://tinyurl.com/Pubmed-DEPASS>

**Description** List of publications in Pubmed

### Access

Terms of data access (charter for data provision, format of data, availability delay)

Publication in progress

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only