

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

Responsable(s) : Crochard Anne

Date de modification : 01/01/2020 | Version : 2 | ID : 178

Général

Identification

Nom détaillé DEPRESSION ? ANXIETY AND DISABILITY IN A COHORT OF PATIENTS IN GENERAL MEDICINE PRACTICE

Sigle ou acronyme DéPasS

Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.) CNIL n° 909006

Thématisques générales

Domaine médical General practice
Psychology and psychiatry

Mots-clés depression, anxiety, functional disability, Sheehan, general practice, antidepressant

Responsable(s) scientifique(s)

Nom du responsable Crochard

Prénom Anne

Adresse 37-45 Quai du Président Roosevelt 92445 Issy-Les-Moulineaux

Téléphone +33 (0)1 79 41 28 51

Email acro@lundbeck.com

Organisme Lundbeck

Collaborations

Financements

Financements Private

Précisions Lundbeck SAS Laboratory

Gouvernance de la base de données

Organisation(s) responsable(s) ou promoteur Lundbeck SAS

Statut de l'organisation Secteur Privé

Contact(s) supplémentaire(s)

Caractéristiques

Type de base de données

Type de base de données Study databases

Base de données issues d'enquêtes, précisions Longitudinal study (except cohorts)

Origine du recrutement des participants A selection of health care professionals

Critère de sélection des participants Medication(s) taken

Le recrutement dans la base de données s'effectue dans le cadre d'une étude interventionnelle No

Informations complémentaires concernant la constitution de l'échantillon 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.

Objectif de la base de données

Objectif principal 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.

Critères d'inclusion 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.

Type de population

| | |
|-----|-----------------------------------------------------------------------------------------------------------------------------------------------------|
| 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 concernée | Sick population |
|----------------------|-----------------|

| | |
|------|---------------|
| Sexe | Male Woman |
|------|---------------|

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| Champ géographique | National |
|--------------------|----------|

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|------------------------------|---------------------|
| Détail du champ géographique | Metropolitan France |
|------------------------------|---------------------|

Collecte

Dates

| | |
|--------------------------|------|
| Année du premier recueil | 2009 |
|--------------------------|------|

| | |
|--------------------------|------|
| Année du dernier recueil | 2010 |
|--------------------------|------|

Taille de la base de données

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|------------------------------------------------------|---------------------------|
| Taille de la base de données (en nombre d'individus) | [1000-10 000] individuals |
|------------------------------------------------------|---------------------------|

| | |
|------------------------------|-------|
| Détail du nombre d'individus | 8 029 |
|------------------------------|-------|

Données

| | |
|---------------------|---------------------------|
| Activité de la base | Data collection completed |
|---------------------|---------------------------|

| | |
|-----------------------------|-----------------------------------|
| Type de données recueillies | Clinical data Declarative data |
|-----------------------------|-----------------------------------|

| | |
|----------------------------------------------|--------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------------|
| Données cliniques, précisions | Direct physical measures |
| Données déclaratives, précisions | Paper self-questionnaire |
| Existence d'une biothèque | No |
| Paramètres de santé étudiés | Health event/morbidity Quality of life/health perception |
| Modalités | |
| Mode de recueil des données | 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 ± 2 weeks was considered acceptable for the 6-week visit and a window of ± 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. |
| Suivi des participants | Yes |
| Détail du suivi | 3 months |
| Appariement avec des sources administratives | No |
| Valorisation et accès | |
| Valorisation et accès | |
| Lien vers le document | http://tinyurl.com/Hal-DEPASS |
| Description | List of publications in HAL |
| Lien vers le document | http://tinyurl.com/Pubmed-DEPASS |
| Description | List of publications in Pubmed |
| Accès | |
| Charte d'accès aux données | Publication in progress |

(convention de mise à disposition, format de données et délais de mise à disposition)

Accès aux données agrégées Access on specific project only

Accès aux données individuelles Access on specific project only