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

Responsable(s) :Crochard Anne

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Général

Identification

Nom détaillé	DEPRESSION ? ANXIETY AND DISABIILTY 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

Thématiques générales

Domaine médical	General practice Psychology and psychiatry
Mots-clés	depression, anxiety, functional disability, Sheehan, general practice, antidepressant

Responsable(s) scientifique(s)

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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
Champ géographique	National
Détail du champ géographique	Metropolitan France
Collecte	
Dates	
Année du premier recueil	2009
Année du dernier recueil	2010
Taille de la base de données	
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' disbility 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