ATHMOS - Validation of the ACT questionnaire in general practice

Responsable(s) :Pribil Céline, Laboratoire GSK

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Général

Identification

Nom détaillé	Validation of the ACT questionnaire in general practice
Sigle ou acronyme	ATHMOS
Numéro d'enregistrement (ID-RCB ou EUDRACT, CNIL, CPP, etc.)	CNIL n°90 60 78 (12/05/2006)

Thématiques générales

Domaine médical	Pneumology
Autres, précisions	Asthma
Mots-clés	asthma, control, "Asthma Control Test ©" (ACT)

Responsable(s) scientifique(s)

Nom du responsable	Pribil
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Laboratoire	Laboratoire GSK

Collaborations

Financements

Financements	Private
Précisions	GSK laboratory

Gouvernance de la base de données

Organisation(s) responsable(s)	Laboratoire GSK

ou promoteur	
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 An administrative base or a register
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 GP: A pre-selection via random drawing will be established using a CEGEDIM file of general practitioners. The general practitioners pre-selected through the random drawing will receive a mailing which will briefly present the objectives of the study and how it will unfold. It will include a postage-paid response card to be returned to the logistics center indicating whether or not the practitioner wants to participate in the study A telephone call to confirm participation will be made to the first 1000 general practitioners who have agreed Selection of pneumologists: A pre-selection via random drawing will be established using a CEGEDIM file of pneumologist/allergy doctors. The pre-selected pneumologists/allergy doctors will be contacted by telephone (brief presentation of the objectives and the unfolding of the study) in the order of the preselection until the number of pneumologists/allergy doctors accepting to participate reaches 500. The patients will be included after they have read the patient information letter

Objectif de la base de données

Objectif principal	Describe in actual practice the level of control of asthma using the ACT questionnaire (Asthma Control Test)
Critères d'inclusion	Male or female aged 18 years or older Patient having asthma diagnosed at least 12 months ago Patient informed of the objectives of the study and accepting the collection and the analysis of the data concerning him.
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	France
Collecte	
Dates	
Année du premier recueil	2006
Année du dernier recueil	2008
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	2362
Données	
Activité de la base	Data collection completed
Type de données recueillies	Clinical data Declarative data
Données cliniques, précisions	Direct physical measures Medical registration

Données déclaratives, précisions	Paper self-questionnaire
Existence d'une biothèque	No
Paramètres de santé étudiés	Health care consumption and services Quality of life/health perception
Consommation de soins, précisions	Hospitalization Medical/paramedical consultation Medicines consumption

Modalités

Mode de recueil des données	The physician will complete the medical questionnaire for inclusion and will give the inclusion self-questionnaire to the patient. During the 3-month inclusion period, the investigating physicians must collect in a data register patients who are not included in the cohort, that meet the eligibility criteria but who cannot or do not want to participate in the study. The patients included will be examined again at the next spontaneous consultation which will also be a follow-up visit within the framework of the study where a follow-up questionnaire will be completed by the investigator and a follow-up self-questionnaire will be given to the patient
Suivi des participants	Yes
Détail du suivi	spontaneous consultation following the inclusion during which will be completed a medical follow-up questionnaire and a follow-up self-questionnaire
Appariement avec des sources administratives	No

Valorisation et accès

Valorisation et accès

Accès

Charte d'accès aux données (convention de mise à disposition, format de données et délais de mise à disposition)	Publications in progress
Accès aux données agrégées	Access on specific project only
Accès aux données individuelles	Access on specific project only