

ATHMOS - Validation of the ACT questionnaire in general practice

Head :Pribil Céline, Laboratoire GSK

Last update : 01/01/2018 | Version : 1 | ID : 166

General	
Identification	
Detailed name	Validation of the ACT questionnaire in general practice
Sign or acronym	ATHMOS
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL n°90 60 78 (12/05/2006)
General Aspects	
Medical area	Pneumology
Others (details)	Asthma
Keywords	asthma, control, "Asthma Control Test ©" (ACT)
Scientific investigator(s) (Contact)	
Name of the director	Pribil
Surname	Céline
Phone	+33 (0)1 39 17 90 62
Email	celine.c.pribil@gsk.com
Unit	Laboratoire GSK
Collaborations	
Funding	
Funding status	Private
Details	GSK laboratory
Governance of the database	

Sponsor(s) or organisation(s) responsible	Laboratoire GSK
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 An administrative base or a register
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	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
Database objective	

Main objective	Describe in actual practice the level of control of asthma using the ACT questionnaire (Asthma Control Test)
Inclusion criteria	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.
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)
Population covered	Sick population
Gender	Male Woman
Geography area	National
Detail of the geography area	France
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2006
Date of last collection (YYYY or MM/YYYY)	2008
Size of the database	
Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	2362
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Declarative data

Clinical data (detail)	Direct physical measures Medical registration
Declarative data (detail)	Paper self-questionnaire
Presence of a biobank	No
Health parameters studied	Health care consumption and services Quality of life/health perception
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption
Procedures	
Data collection method	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
Participant monitoring	Yes
Details on monitoring of participants	spontaneous consultation following the inclusion during which will be completed a medical follow-up questionnaire and a follow-up self-questionnaire
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Publications in progress
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only

