

OCSIGEN - Follow-up cohort of asthmatic patient treated with inhaled corticosteroid

Responsable(s) :Pribil Céline, Laboratoire GSK

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Général

Identification

Nom détaillé Follow-up cohort of asthmatic patient treated with inhaled corticosteroid

Sigle ou acronyme OCSIGEN

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

Thématiques générales

Domaine médical Pneumology

Déterminants de santé Medicine

Mots-clés inhaled steroids, cohort

Responsable(s) scientifique(s)

Nom du responsable Pribil

Prénom Céline

Téléphone +33 (0)1 39 17 90 62

Email celine.c.pribil@gsk.com

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

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

GP section:
The investigating physicians will be the Thalès physicians accepting to participate in the study

Pneumologist section:
The investigating centers for the study will be obtained by random drawing stratified over the region. The minimum survey rate allowing for representation of each of the regions is about 4%, which corresponds to a sample of 101 pneumologists in France

Objectif de la base de données

Objectif principal

Describe the conditions of use of the fluticasone and other inhaled corticosteroids in a pragmatic situation in general practice and in specialized pneumology, evaluate the severity of the asthma of patients cared for and measure the adequacy of the care with the national recommendations

Critères d'inclusion

GP section:
The eligible population is defined by all of the asthmatic patients over the age of 15 years who have consulted at least twice in the previous year and in whom an inhaled corticotherapy is in progress (at least one prescription in the last six

months). Patients defined as such who come to consult spontaneously for asthma during the period of inclusion will also be included. Refusal to participate and the association of the asthma with an OCPD are criteria for non-inclusion.

Pneumologist section:

Asthmatic persons cared for in liberal pneumology present in the practice of the investigators of the study and meeting the inclusion criteria

Type de population

Age	Adolescence (13 to 18 years) 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 concernée	Sick population
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Sexe	Male Woman
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Champ géographique	National
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Détail du champ géographique	France
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Collecte

Dates

Année du premier recueil	2004
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Année du dernier recueil	2009
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Taille de la base de données

Taille de la base de données (en nombre d'individus)	[1000-10 000[individuals
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Détail du nombre d'individus	1691
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Données

Activité de la base	Data collection completed
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Type de données recueillies	Clinical data Declarative data
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Données cliniques, précisions	Direct physical measures Medical registration
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Données déclaratives, précisions	Paper self-questionnaire
Existence d'une bibliothèque	No
Paramètres de santé étudiés	Health care consumption and services
Consommation de soins, précisions	Hospitalization Medical/paramedical consultation Medicines consumption
Modalités	
Mode de recueil des données	<p>GP section: The characteristics of the patients will be collected in historical data present in the Thalès database, then at the inclusion visit and during all of the visits occurring during the follow-up period. At each visit, the stage of the severity of the patients according to GINA clinical and therapeutic classifications will be evaluated. During the inclusion period, at each consultation of an eligible patient, the study will be offered to him. Where applicable, the computerized inclusion questionnaire is filled out and the patient is included in the cohort. At each consultation carried out during the follow-up period, a "pop-up" screen containing the follow-up questionnaire will appear on the investigator's computer screen.</p> <p>Pneumologist section: The conditions for the use of inhaled steroids according to the stages of severity will be described through three questionnaires: a historical questionnaire (data coming from the medical dossier of the patient), an inclusion questionnaire and a follow-up questionnaire which will be implemented in computerized format. A fourth questionnaire, self-administered pertaining to the control of the asthma, will be completed by the patient at inclusion and at each visit during the follow-up. The collection of data will be carried out on a computerized support via a secure internet site devoted to the study. The physician will directly enter the information desired into the on-line forms</p>
Suivi des participants	Yes
Détail du suivi	<p>GP section: Follow-up for 24 months, Pneumologist section: Follow-up for 24 months, at each new consultation carried out during the follow-up period, the pneumologist fills out the computerized on-line follow-up questionnaire</p>
Appariement avec des sources	No

administratives

Valorisation et accès

Valorisation et accès

Lien vers le document

<http://tinyurl.com/pubmed-ocsigen>

Description

Liste des publications dans Pubmed

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