

SYMBIOSE - Descriptive pharmaco-epidemiological study on the use of Symbicort® Turbuhaler® in the treatment of asthma in France and impact of a new treatment strategy on compliance and asthma control

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General

Identification

Detailed name Descriptive pharmaco-epidemiological study on the use of Symbicort® Turbuhaler® in the treatment of asthma in France and impact of a new treatment strategy on compliance and asthma control

Sign or acronym SYMBIOSE

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL : 908418 - 28/01/2009

General Aspects

Medical area General practice
Pneumology

Keywords Symbicort® Turbuhaler®, cohort

Scientific investigator(s) (Contact)

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Unit	AstraZeneca

Collaborations

Funding

Funding status	Private
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Details	AstraZeneca
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Governance of the database

Sponsor(s) or organisation(s) responsible	ASTRAZENECA
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 institutions and services
Database recruitment is made on the basis of:	Medication(s) taken
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	The physicians and the pulmonologists participating in the study recruit the subjects to be included among the patients that they see in consultation and according to the inclusion criteria

Database objective

Main objective	Describe the characteristics of asthmatic patients treated with Symbicort Turbuhaler
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Compare the characteristics of patients treated according to the therapeutic strategy
Evaluate the level of asthma control and compliance with treatment of patients treated with Symbicort Turbuhaler

Inclusion criteria	Patients diagnosed as asthmatic by the physician and treated for this pathology with Symbicort Turbuhaler
Population type	
Age	Childhood (6 to 13 years) 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)
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)	2009
Date of last collection (YYYY or MM/YYYY)	2011
Size of the database	
Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	1844
Data	
Database activity	Current data collection
Type of data collected	Clinical data Declarative data

Clinical data (detail)	Medical registration
Declarative data (detail)	Phone interview

Presence of a biobank	No
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Health parameters studied	Health event/morbidity Quality of life/health perception
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Procedures

Participant monitoring	Yes
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Details on monitoring of participants	Follow-up duration of 12 months
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Links to administrative sources	No
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Promotion and access

Promotion

Access

Terms of data access (charter for data provision, format of data, availability delay)	Methods for accessing the database are currently being defined
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Access to aggregated data	Access on specific project only
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Access to individual data	Access on specific project only
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