

ATLANTIS - Long term observational study on the therapeutic management of knee osteoarthritis by general practitioners

Head :Payen-Champenois Catherine, Bristol-Myers Squibb

Schmidely Nathalie, Bristol-Myers Squibb

Last update : 09/05/2017 | Version : 1 | ID : 121

General

Identification

Detailed name	Long term observational study on the therapeutic management of knee osteoarthritis by general practitioners
---------------	---

Sign or acronym	ATLANTIS
-----------------	----------

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL n°903352 (22/11/2003)
--	----------------------------

General Aspects

Medical area	Rheumatology
--------------	--------------

Others (details)	gonarthrosis, Arthrosis
------------------	-------------------------

Keywords	therapeutic management, paracetamol, patient reported outcomes
----------	--

Scientific investigator(s) (Contact)

Name of the director	Payen-Champenois
----------------------	------------------

Surname	Catherine
---------	-----------

Address	3, rue J. Monnier - 92500 Rueil Malmaison
---------	---

Phone	+33 (0)1 58 83 60 00
-------	----------------------

Email	catherine.payen-champenois@bms.com
-------	------------------------------------

Unit	Bristol-Myers Squibb
------	----------------------

Name of the director	Schmidely
----------------------	-----------

Surname	Nathalie
---------	----------

Address	3, rue J. Monnier - 92500 Rueil Malmaison
Phone	+33 (0)1 58 83 60 00
Email	nathalie.schmidely@bms.com
Unit	Bristol-Myers Squibb
Collaborations	
Funding	
Funding status	Private
Details	Bristol-Myers Squibb
Governance of the database	
Sponsor(s) or organisation(s) responsible	Bristol-Myers Squibb (BMS)
Organisation status	Private
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Cohort study
Database recruitment is carried out by an intermediary	A selection of health care professionals
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Random sampling in clusters
Database objective	
Main objective	Evaluate, in general practice, the change in the first-line therapeutic strategy in patients suffering from knee osteoarthritis, over one year, specially: - Evaluate the percentage of patients treated with each therapeutic modality at the end of the one year observation period. - Describe the modifications implemented in the

therapeutic management and identify the reasons justifying these modifications.

Inclusion criteria	1. Males or females 50 years or older suffering from symptomatic osteoarthritis of the knee 2. Not treated over the last 6 months for this pathology (non-pharmaceutical treatments and self-medication by analgesics or NSAIDs at low doses will be allowed). 3. For whom the physician will have decided to initiate a first-line treatment in line with the recommendations of EULAR 2000-2003 (paracetamol - systematically taken at a dose of 3 to 4g/d - as pharmacological monotherapy in combination with non-pharmaceutical treatments).
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	International
Detail of the geography area	France and Spain
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2003
Date of last collection (YYYY or MM/YYYY)	2006
Size of the database	
Size of the database (number of individuals)	[1000-10 000[individuals
Details of the number of individuals	1114 (586 in France, 528 in Spain)
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)	Face to face interview
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	paper case report form
Participant monitoring	Yes
Details on monitoring of participants	M3, M6, M9, M12
Links to administrative sources	No
Promotion and access	
Promotion	
Access	
Terms of data access (charter for data provision, format of data, availability delay)	Publications
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only