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

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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)

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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