

# LOCOX - Adaptation of locomotor activity in patients suffering from hip osteoarthritis

Head :Bonithon-Kopp Claire, CHU DIJON / INSERM CIE 1 UNIVERSITÉ DE BOURGOGNE  
Maillefert , CHU DIJON

Last update : 07/30/2014 | Version : 1 | ID : 60148

## General

### Identification

Detailed name Adaptation of locomotor activity in patients suffering from hip osteoarthritis

Sign or acronym LOCOX

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL : 28/04/2008

### General Aspects

Medical area Rheumatology

Others (details) Hip osteoarthritis

Keywords total hip replacement, Health episodes, indication

### Scientific investigator(s) (Contact)

Name of the director Bonithon-Kopp

Surname Claire

Address 21000 DIJON

Email bonithon@u-bourgogne.fr

Unit CHU DIJON / INSERM CIE 1 UNIVERSITÉ DE BOURGOGNE

Organization CHU

Name of the director Maillefert

Address 21000 DIJON

Phone + 33 (0)3 80 29 37 45

Unit	CHU DIJON
Organization	CHU
<b>Collaborations</b>	
<b>Funding</b>	
Funding status	Public
Details	PHRC NATIONAL 2004 ET CHU DE DIJON
<b>Governance of the database</b>	
Sponsor(s) or organisation(s) responsible	CHU Dijon
Organisation status	Public
<b>Additional contact</b>	
<b>Main features</b>	
<b>Type of database</b>	
Type of database	Study databases
Study databases (details)	Cohort study
Database recruitment is carried out by an intermediary	A selection of health institutions and services
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Prospective Inclusion cut-off date: 01/10/2009
<b>Database objective</b>	
Main objective	General objective: to assess the prognostic value of a characteristic evaluation of gait in persons suffering from hip osteoarthritis for subsequent indications for total hip replacement. Analysis of the adaptation of locomotor activity as a prognostic criterion for future disease progression.
Inclusion criteria	Men and women aged between 45 and 75 years - suffering from primary hip osteoarthritis defined according to ACR criteria; - presented pain in the hip for at least one month in the preceding 3

months; - radiological stage II, III or IV hip osteoarthritis according to the Kellgren and Lawrence classification; - able to understand simple instructions, packaging instructions and give their informed consent. Exclusion criteria - indication of total hip replacement at the time of inclusion; - pregnant or breast-feeding women; - Alzheimer's disease; - chronic respiratory insufficiency with clinical manifestations; - Parkinson's disease; - motor neuron disease; - major musculo-skeletal disorder (other than hip osteoarthritis); - severe non-stabilised diabetes; - non-stabilised hypertension; - hip osteoarthritis inflammation flare; - rapidly destructive hip osteoarthritis; - radiography to evaluate structural evolution of the disease done more than six months previously; - Presence of osteoarthritis in another joint, or another osteoarticular or periarticular disease of the lower limbs with, in the opinion of the patient, disability due to this other disease that is greater than the disability caused by the hip osteoarthritis; - Spinal disease causing gait disturbance (radiculalgia, lumbar canal stenosis,? with, in the opinion of the patient, disability due to this other disease that is greater than the disability caused by the hip osteoarthritis.).

## Population type

Age  
Adulthood (45 to 64 years)  
Elderly (65 to 79 years)

Population covered  
Sick population

Gender  
Male  
Woman

Geography area  
Local

French regions covered by the database  
Bourgogne Franche-Comté

Detail of the geography area  
Dijon

## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY)  
02/2008

Date of last collection (YYYY or MM/YYYY)  
10/2014

## Size of the database

Size of the database (number of individuals) < 500 individuals

Details of the number of individuals 35 patients

## Data

Database activity Current data collection

Type of data collected  
Clinical data  
Declarative data  
Paraclinical data

Clinical data (detail)  
Direct physical measures  
Medical registration

Declarative data (detail)  
Paper self-questionnaire

Paraclinical data (detail)  
Measurement of locomotor activity on specific platform

Presence of a biobank No

Health parameters studied Health event/morbidity

## Procedures

Data collection method  
Self-administered questionnaire: from paper questionnaire (manual input) and double data entry  
Interview: from paper questionnaire (manual input) with double data entry  
Clinical Examinations: handwritten (manual input) with double data entry

Participant monitoring Yes

Details on monitoring of participants  
Follow-up duration: 5 years

Links to administrative sources No

## Promotion and access

### Promotion

### Access

Terms of data access (charter for data provision, format of data, availability delay)  
To be decided if data may be used by academic teams  
To be decided if data may be used by industrial teams

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only