

PERFECTA - Intravenous analgesics for post-operative pain: patterns of use in high-risk populations

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Last update : 01/01/2018 | Version : 1 | ID : 126

General

Identification

Detailed name Intravenous analgesics for post-operative pain: patterns of use in high-risk populations

Sign or acronym PERFECTA

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation --

General Aspects

Health determinants Iatrogenic

Others (details) Pain

Keywords Post-chirurgical, treatment for pain, IV pain killers

Scientific investigator(s) (Contact)

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Collaborations

Funding

Funding status	Private
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Details	Bristol-Myers Squibb
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Governance of the database

Sponsor(s) or organisation(s) responsible	Bristol-Myers Squibb France (BMS)
Organisation status	Private

Additional contact

Main features

Type of database

Type of database	Study databases
Study databases (details)	Not-repeated cross-sectional studies (except case control studies)
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.	random sampling in clusters

Database objective

Main objective	Describe the post-surgical drug treatment for pain after a surgical intervention, in target groups of patients having specific risks of developing complications during the use of certain classes of analgesics
Inclusion criteria	M/F patients, ≥ 18 years, have an ASA score of level 3 or 4 and having undergone a surgical

intervention, including coeloscopy, excluding cardiac surgery and neurosurgery, requiring antalgic treatment through IV in order to relieve postoperative pain.
Patients having at least one of the following risk factors: aged \geq 65 years, OCPD including sleep apnea, high blood pressure, kidney, heart or liver failure.

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 National

Detail of the geography area Metropolitan France

Data collection

Dates

Date of first collection (YYYY or MM/YYYY) 2007

Date of last collection (YYYY or MM/YYYY) 2008

Size of the database

Size of the database (number of individuals) [1000-10 000[individuals

Details of the number of individuals 1829

Data

Database activity Data collection completed

Type of data collected Clinical data

Clinical data (detail) Direct physical measures

Presence of a biobank No

Health parameters studied Health event/morbidity
Health event/mortality

Procedures

Data collection method paper CRF

Participant monitoring No

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