

I-NORM - Management by French General Practitioners of uncontrolled hypertensive patients treated by dual therapy

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General

Identification

Detailed name	Management by French General Practitioners of uncontrolled hypertensive patients treated by dual therapy
Sign or acronym	I-NORM
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL N° 833542 version 16

General Aspects

Medical area	Cardiology
Others (details)	high blood pressure
Keywords	High blood pressure, uncontrolled hypertension, dual therapy, therapeutic strategy, care, hypertension

Scientific investigator(s) (Contact)

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Unit	Bristol-Myers Squibb

Collaborations

Funding

Funding status	Private
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Details Bristol-Myers Squibb France / Sanofi-Aventis France

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 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.	random sampling in clusters

Database objective

Main objective Describe medical strategies used by general practitioners for uncontrolled hypertension despite dual-therapy: the I-NORM survey

Inclusion criteria	Register: Patient ≥ 18 years, seen in routine consultation, monitored for HBP, known for more than 3 months, treated via fixed or free dual therapy for his HBP, dual therapy initiated by the participating doctor or the corresponding specialist. Study: pt from the register, uncontrolled HBP (SBP ≥ 140 mmhg and /or DBP ≥ 90 mmhg all comor pt or SBP ≥ 130 mmhg and /or DBP ≥ 80 mmhg diabetic pt or with kidney failure), not participating into a clinical trial
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	25211 : - Registre: 21208 - Etude: 4003
Data	
Database activity	Data collection completed
Type of data collected	Clinical data Administrative data

Clinical data (detail)	Direct physical measures
Declarative data (detail)	Phone interview
Administrative data (detail)	questionnaire

Presence of a biobank	No
Health parameters studied	Health care consumption and services

Care consumption (detail)	Medical/paramedical consultation Medicines consumption
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Procedures

Data collection method	Paper CRF
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Participant monitoring	No
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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)	Publications
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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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