INTERMEDE - Cross-Sectional Survey on Patient-Physician Interaction and Social Health Inequalities

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General

Identification

Detailed name	Cross-Sectional Survey on Patient-Physician Interaction and Social Health Inequalities
Sign or acronym	INTERMEDE
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	Accord CNIL

General Aspects

Medical area	Disability/handicap Psychology and psychiatry
Health determinants	Social and psychosocial factors
Keywords	Physician-patient relationship, social health inequalities, health care sought, treatment.

Scientific investigator(s) (Contact)

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Unit	Unité Inserm 558, Faculté de Médecine de Toulouse
Organization	INSERM

Collaborations

Funding

Funding status	Public
Details	Institut de Recherche en Santé Publique (IReSP), CHU Toulouse, Assurance Maladie

Governance of the database

Sponsor(s) or organisation(s) responsible	INSERM
Organisation status	Public
Sponsor(s) or organisation(s) responsible	Université Paul Sabatier
Organisation status	Public

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 population file
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	The sample of patients was established in the waiting room of general practices. Recruitment of general practitioners was done in partnership with SMFG. A call for applications was sent to members of the SMFG network. Participating physicians were included on a voluntary basis with a target of 10 physicians per site.

Database objective

Main objective	To clarify elements of physician-patient interaction that may explain, beyond social differences in health care sought by patients, differentiated treatment that may ultimately contribute to differences in health. - Consultation observation and semi-structured post-consultation interviews with patients and doctors independently. - Mirrored-

	questionnaires where patients and physicians answered the same questions post-consultation.
Inclusion criteria	Patient inclusion: - at least 18 years old - not attending a general practitioner for emergency purposes - who are not attending their first consultation - not pregnant or have given birth within the last six months - who have signed the informed consent agreement

Population type

Age	Adulthood (19 to 24 years) Adulthood (25 to 44 years) Adulthood (45 to 64 years)
Population covered	Sick population
Gender	Male Woman
Geography area	National
Detail of the geography area	Ile de France, Midi Pyrénées and Pays de la Loire

Data collection

Dates

Date of first collection (YYYY or MM/YYYY)	09/2006
Date of last collection (YYYY or MM/YYYY)	10/2007

Size of the database

Size of the database (number of individuals)	[500-1000[individuals
Details of the number of individuals	phase I:11 médecins généralistes/MP, 249 patients phase II:27 médecins généralistes/MP, 585 patients

Data

Database activity	Data collection completed
Type of data collected	Declarative data
Declarative data (detail)	Paper self-questionnaire 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)	Medical/paramedical consultation

Procedures

Participant monitoring	No
Links to administrative sources	No

Promotion and access

Promotion

Link to the document	http://www.biomedcentral.com/1472-6963/9/66
Link to the document	http://www.sciencedirect.com/science/article/pii/S0738399112004600
Link to the document	http://tinyurl.com/PUBMED-INTERMEDE
Link to the document	http://tinyurl.com/HAL-INTERMEDE

Access

Terms of data access (charter for data provision, format of data, availability delay)	Contact the scientist in charge
Access to aggregated data	Access on specific project only
Access to individual data	Access on specific project only