ESPS - Health, health care and insurance survey

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General

Identification

Detailed name	Health, health care and insurance survey
Sign or acronym	ESPS
CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL n°1147702

General Aspects

Pathology (details)	Mandatory and supplemental health insurance. Perceived state of health. General health. Mental health. Dental health. Functional disabilities. Risk factors. Recourse and access to care. Renouncing care.
Health determinants	Addictions Geography Occupation Social and psychosocial factors
Keywords	Survey of the general population. Panel. State of health. Access to care. Renouncing care. Recourse to care. Mandatory health insurance. Additional health insurance. Evin Law. Family doctor. Medical deductibles. CMU (universal medical coverage). ACS (supp

Scientific investigator(s) (Contact)

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Organization	IRDES
Collaborations	
Funding	
Funding status	Mixed
Details	CNAMTS, RSI, MSA, DREES, la Mutualité, Invs.
Governance of the database	
Sponsor(s) or organisation(s) responsible	IRDES - Institut de Recherche et Documentation en Economie de la Santé
Organisation status	Both
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	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 ESPS sample is comprised as a sub-samples of the general sample of beneficiaries (EGB) using fixed characteristics of the NIR. An identical sample is comprised for RSI and MSA.

Database objective

Main objective	There are multiple purposes of the survey: Firstly, ESPS (health and social protection survey) makes it possible to study the existing links on an individual level between the state of health, access to health services, access to public and private insurance and the socio-economic status. This makes it possible to conduct many analyses covering equity in the health system and to enlighten public efforts by participating in the evaluation of public policies, whether directly or indirectly linked to health. ESPS is as such a reactive tool for researchers in the social sciences, making it possible to test new research hypotheses on French data. ESPS is also an information tool for public health. It makes it possible to assess in an exploratory manner the prevalences of affections in the absence of specific instruments and complete sources, and this using detailed data on morbidity. ESPS thus comprises a unique source of data on supplemental health insurance in France. Through its specific questioning pertaining to the supplemental contracts that the members of the households surveyed benefit from, it offers a panorama of the contracts that are actually in effect and as such of the distribution and of the changes in the levels of coverage in the population.
Inclusion criteria	Until 2008, a random sample of the insured with mandatory health insurance (Cnamts, RSI, MSA) stemming from the permanent Sample of social insurance beneficiaries (EPAS) and the members of their households. Starting in 2010, a random sample of the major beneficiaries of health insurance (Cnamts, RSI, MSA) and the members of their households.

Population type

Age	Newborns (birth to 28 days) Infant (28 days to 2 years) Early childhood (2 to 5 years) Childhood (6 to 13 years) Adolescence (13 to 18 years) 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	General 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)	1988
Size of the database	
Size of the database (number of individuals)	Greater than 20 000 individuals
Details of the number of individuals	Plus de 8 000 ménages et plus de 22 000 individus par enquête, soit près de 45 000 individus par coupe transversale du panel qui est sollicité par moitié tous les 4 ans. More than 8,000 households and more than 22,000 individuals per survey, which is nearly 45,000 individuals per transversal section of the panel which half is solicited every 4 years.
Data	
Database activity	Current data collection
Type of data collected	Declarative data Administrative data
Declarative data (detail)	Paper self-questionnaire Face to face interview Phone interview
Administrative data (detail)	EGB - SNIIR-AM
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services Quality of life/health perception Others
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption

Other (detail)	Renouncing care

Procedures

Data collection method	A main CAPI-CATI questionnaire is given during two contacts 15 days apart. The questionnaire is offered by telephone to all households for which the survey institution was able to find a telephone number and in person for the others. The households of which the beneficiaries sampled are beneficiaries of the CMU or 70 years of age or more (information present in the EGB - survey base), as well as households of 10 or more people are surveyed in person. Between two contacts, all of the members of the household are invited to complete the self-administered paper questionnaires, of at least one on health and another on the household's supplemental health insurance contracts.
Classifications used	ICD 10 for the diseases. Nomenclature of Socio-professional Categories and Professions (PCS).
Participant monitoring	Yes
Details on monitoring of participants	Half of the ESPS sample constructed using the EGB on the fixed characteristics of the NIR is surveyed every two years. Every four years, the sample is thus comprised of the same individuals, modulo additions and subtractions to the EGB.
Links to administrative sources	Yes
Linked administrative sources (detail)	SNIIR-AM - data on inter-system care consumption presented for reimbursement.

Promotion and access

Promotion

Link to the document	http://www.irdes.fr/EspaceRecherche/Enquetes/ESPS/EnqueteESPSBiblio.htm
Link to the document	http://www.irdes.fr/EspaceRecherche/Enquetes/ESPS/Documents/EPASBibliographie.pdf
Link to the document	http://www.cmh.greco.ens.fr/enquetes/XML/lil-0454.xml

Access

Terms of data access (charter for data provision, format of	The data from the ESPS survey is distributed by Irdes. Those financing the project have 12 months

data, availability delay)	of exclusive use of the data which is therefore accessible to the other organizations 22 months after the end of the survey field which itself extends over 10 months, from March to December. All of the information concerning the survey: its description, history, the questionnaires, results and publications are available at the address below. To access the survey databases, simply send your request to: esps@irdes.fr. We will send you an agreement project in which you will have to present your research or study project. Data from the ESPS survey can be accessed free of charge by all research or public data production organizations. The data from 2006 is distributed via the Quetelet network. Within this framework, the files can be accessed free of charge for the purposes of research for researchers, researcher-professors, PhD students and French and foreign Masters degree students (http://www.reseau-quetelet.cnrs.fr/spip/spip.php?rubrique67). Any commercial use is prohibited. The specific methods for accessing the ESPS survey are described on the site of the network Registration prior to the communication of the data files is required: http://www.reseau-quetelet.cnrs.fr/quetelet/quetelet.php).
Access to aggregated data	Free access
Access to individual data	Access on specific project only