

# Entred 2007-2010 - National Representative Sample of people with diabetes 2007-2010

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## General

### Identification

Detailed name National Representative Sample of people with diabetes 2007-2010

Sign or acronym Entred 2007-2010

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

907172

### General Aspects

Medical area Endocrinology and metabolism

Health determinants Lifestyle and behavior

Keywords type 1 diabetes, vascular risk, socioeconomic level, socioeconomic status, educational measures, quality of care, cost, type 2 diabetes, characteristics, complication, hospitalisation, mortality

### Scientific investigator(s) (Contact)

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Organization INVS - Institut de Veille

### Collaborations

### Funding

Funding status	Public
Details	InVS, CnamTS, RSI, HAS, INPES.
Governance of the database	
Sponsor(s) or organisation(s) responsible	INVS - Institut de Veille Sanitaire
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	An administrative base or a register
Database recruitment is carried out as part of an interventional study	No
Additional information regarding sample selection.	Sample is based on a random selection of personal keys in the Local Health Insurance databases. The sample consists of individuals with these keys who have received at least three reimbursements for oral antidiabetic medications and/or insulin over the last twelve months.
Database objective	
Main objective	To describe the characteristics of people pharmacologically treated for diabetes, their health with regards to diabetes, the quality of care received, treatment plan and self-management education; their quality of life, experiences and requirements in terms of education and information, as well as the cost of diabetes.
Inclusion criteria	Beneficiaries of the French general scheme of health insurance (CNAMTS) (excluding local mutualist sections) and the Independent Scheme for employees (RSI); residing in mainland France or overseas départements (DOM); and who have received at least three reimbursements for oral

antidiabetic medications and/or insulin over the last twelve months.

## 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)
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Population covered	Sick population
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Gender	Male Woman
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Geography area	National
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Detail of the geography area	France
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## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY)	08/2006
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Date of last collection (YYYY or MM/YYYY)	07/2009
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### Size of the database

Size of the database (number of individuals)	[10 000-20 000[ individuals
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Details of the number of individuals	8,000 people in metropolitan France, 800 people from overseas départements and 850 children.
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## Data

Database activity	Data collection completed
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Type of data collected	Clinical data Declarative data Biological data Administrative data
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Clinical data (detail)	Direct physical measures
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Details of collected clinical data	Medical questionnaire sent to attending physicians.
Declarative data (detail)	Paper self-questionnaire Phone interview
Details of collected declarative data	Self-administered patient questionnaire and phone questionnaire administered by National Health Insurance medical officers.
Biological data (detail)	Medical questionnaire sent to physicians.
Administrative data (detail)	Medical consumption, hospitalisation data and mortality data.
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality Health care consumption and services Quality of life/health perception
Care consumption (detail)	Hospitalization Medical/paramedical consultation Medicines consumption
Procedures	
Data collection method	Data collected by self-administered questionnaire, phone questionnaire and physician questionnaire, combined with passive medical and administrative data collection (medical consumption and hospitalisation) and mortality data (vital status and cause of death).
Classifications used	CIM 10.
Participant monitoring	Yes
Details on monitoring of participants	Passive medical and administrative data collection (medical consumption and hospitalisation) for two years and mortality data (vital status and cause of death).
Links to administrative sources	Yes
Linked administrative sources (detail)	Local medical consumption data, PMSI, RNIPP and CépiDC.
Promotion and access	
Promotion	

Link to the document

[principales\\_publications.docx](#)

## Access

Terms of data access (charter for data provision, format of data, availability delay)

The results are published as reports, summaries and articles. Complete bibliography on the ENTRED website..

Data are available to external teams after project is submitted to an assessment committee.