

# SOHO-France - Cohort of Schizophrenic Patients Receiving Antipsychotic Drugs

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## General

### Identification

Detailed name	Cohort of Schizophrenic Patients Receiving Antipsychotic Drugs
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Sign or acronym	SOHO-France
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CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation	CNIL
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### General Aspects

Medical area	Disability/handicap Psychology and psychiatry
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Pathology (details)	Schizophrenia
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Health determinants	Genetic Iatrogenic Lifestyle and behavior Medicine
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Keywords	antipsychotic, remission, schizophrenia, relapse, treatment
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### Scientific investigator(s) (Contact)

Name of the director	Gasquet
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Surname	Isabelle
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Organization	Paris Public Hospitals
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### Collaborations

Funding	
Funding status	Private
Details	Lilly France pharmaceutical laboratory
Governance of the database	
Sponsor(s) or organisation(s) responsible	Assistance publique?hôpitaux de Paris
Organisation status	Public
Additional contact	
Main features	
Type of database	
Type of database	Study databases
Study databases (details)	Cohort study
Database recruitment is carried out by an intermediary	A selection of health care professionals
Database recruitment is 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.	Patients were recruited by psychiatrists drawn from the LOGIMED (IMS Health) and TVF (CEGEDIM) sampling frame with layering according to activity type (private versus hospital practice or mixed) and area of practice. Each psychiatrist must recruit four patients consecutively: two patients receiving olanzapine and two patients receiving an anti-psychotic prescribed at the discretion of the treating psychiatrist from available treatment (risperidone, amisulpride, clozapine, typical oral antipsychotics and intramuscular, long-acting medication).
Database objective	
Main objective	To provide information on patients monitored as outpatients, regardless of antipsychotic treatment received and to provide answers, especially regarding treatment adherence and remission rates, as well as relapse rates over a three-year

monitoring period.

Inclusion criteria                      Schizophrenic adult patients monitored as outpatients and randomly recruited by psychiatrists.

## 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                      France

## Data collection

### Dates

Date of first collection (YYYY or MM/YYYY)                      2001

Date of last collection (YYYY or MM/YYYY)                      2005

### Size of the database

Size of the database (number of individuals)                      [500-1000[ individuals

Details of the number of individuals                      964

## Data

Database activity                      Data collection completed

Type of data collected                      Clinical data  
Declarative data

Clinical data (detail)                      Direct physical measures

Details of collected clinical data                      History of illness, previous and ongoing treatments, symptomatology (Clinical Global Impression ?

Schizophrenia [CGI-SCH] scale) and a Visual Analogue Scale [VAS]). Clinical severity was measured by the CGI-SCH scale.

Declarative data (detail)

Face to face interview

Details of collected declarative data

Demographic data, quality of life (EuroQol scale of five items [EQ-5D]; addictive comorbidities in the form of simple questions on past and current use of illicit substances and alcohol; social status (accommodation type, type of work activity, number of social activities, etc.)

Presence of a biobank

No

Health parameters studied

Health care consumption and services

Care consumption (detail)

Medicines consumption

## Procedures

Participant monitoring

Yes

Details on monitoring of participants

Follow-up at three and six months then every six months for three years.

Links to administrative sources

No

## Promotion and access

### Promotion

Link to the document

<http://www.ncbi.nlm.nih.gov/pubmed/19162420>

### 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