

# I-SYPRES - Observational study on the diagnostic elements of Heart Failure with Preserved Systolic Function (HF-PSF) in private practice

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## General

### Identification

Detailed name Observational study on the diagnostic elements of Heart Failure with Preserved Systolic Function (HF-PSF) in private practice

Sign or acronym I-SYPRES

CNIL registration number, number and date of CPP agreement, AFSSAPS (French Health Products Safety Agency) authorisation CNIL n° 907194

### General Aspects

Medical area Cardiology

Others (details) Heart Failure with Preserved Systolic Function

Keywords metabolic and cardiological risk factors

### Scientific investigator(s) (Contact)

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Unit Sanofi Aventis

### Collaborations

### Funding

Funding status Private

Details sanofi-aventis

## Governance of the database

Sponsor(s) or organisation(s) responsible Sanofi-aventis France

Organisation status Private

## Additional contact

## Main features

### Type of database

Type of database Study databases

Study databases (details) Longitudinal study (except cohorts)

Database recruitment is carried out by an intermediary A selection of health care professionals

Database recruitment is carried out as part of an interventional study No

Additional information regarding sample selection. Consecutive inclusion of 4 eligible patients per physician

## Database objective

Main objective Describe in private cardiology practice the diagnostic elements of Heart Failure with Preserved Systolic Function (HF-PSF) for patients:  
- for whom the private cardiologist makes an initial diagnosis of HF-PSF,  
- monitored through appointments and for whom the HF-PSF diagnosis was made during hospitalization

Inclusion criteria Aged over 18; Presenting HF-PSF that has been diagnosed in the past 6 months either by a private cardiologist or during hospitalization; Agreeing to take part in the study

## Population type

Age Adulthood (19 to 24 years)  
Adulthood (25 to 44 years)  
Adulthood (45 to 64 years)  
Elderly (65 to 79 years)

Population covered Sick population

Gender	Male Woman
Geography area	National
Detail of the geography area	France
<b>Data collection</b>	
<b>Dates</b>	
Date of first collection (YYYY or MM/YYYY)	2007
Date of last collection (YYYY or MM/YYYY)	2009
<b>Size of the database</b>	
Size of the database (number of individuals)	[1000-10 000[ individuals
Details of the number of individuals	1168
<b>Data</b>	
Database activity	Data collection completed
Type of data collected	Clinical data Biological data
Clinical data (detail)	Direct physical measures
Biological data (detail)	creatinine levels, blood sodium levels, blood glucose, hemoglobin levels
Presence of a biobank	No
Health parameters studied	Health event/morbidity Health event/mortality
<b>Procedures</b>	
Participant monitoring	Yes
Details on monitoring of participants	6 months
Links to administrative sources	No
<b>Promotion and access</b>	

## Promotion

### Access

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

methods for accessing the database are currently being defined

Access to aggregated data

Access on specific project only

Access to individual data

Access on specific project only