

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

Head :Vignal Franck, Sanofi Aventis

Last update : 01/01/2019 | Version : 1 | ID : 89

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)	
Name of the director	Vignal
Surname	Franck
Phone	+33 (0)1 57 63 26 47
Email	franck.vignal@sanofi-aventis.com
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: <ul style="list-style-type: none"> <li>- for whom the private cardiologist makes an initial diagnosis of HF-PSF,</li> <li>- monitored through appointments and for whom the HF-PSF diagnosis was made during hospitalization</li> </ul>
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
Data collection	
Dates	
Date of first collection (YYYY or MM/YYYY)	2007
Date of last collection (YYYY or MM/YYYY)	2009
Size of the database	
Size of the database (number of individuals)	[1000-10 000[ individuals
Details of the number of individuals	1168
Data	
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
Procedures	
Participant monitoring	Yes
Details on monitoring of participants	6 months
Links to administrative sources	No
Promotion and access	



## 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