

# How health centers organize to offer better, more accessible care

<b>Submission date</b>	<b>Recruitment status</b>	<input type="checkbox"/> Prospectively registered
06/08/2025	No longer recruiting	<input type="checkbox"/> Protocol
<b>Registration date</b>	<b>Overall study status</b>	<input type="checkbox"/> Statistical analysis plan
02/09/2025	Ongoing	<input type="checkbox"/> Results
<b>Last Edited</b>	<b>Condition category</b>	<input type="checkbox"/> Individual participant data
08/08/2025	Other	<input checked="" type="checkbox"/> Record updated in last year

## Plain English summary of protocol

### Background and study aims

In recent years, the way doctors and other healthcare professionals work together in France has been changing. Instead of working alone, many now work as teams in health centers. These teams include general practitioners (GPs), nurses, medical assistants, and even social workers. The goal is to offer patients more complete care that is easier to access and better organized. However, it's not yet clear which ways of organizing these teams work best, especially for patients who are more vulnerable because of age, poverty, or social situations. This study, called EPIDAURE-2, aims to understand how different health centers organize care, how patients and professionals experience it, and whether this improves access and quality of care.

The study is especially focused on the role of the primary care doctor (known in France as the *médecin traitant*), who is meant to coordinate each patient's care, and how this role fits into increasingly team-based care models.

### Who can participate?

Patients of all ages (adults and accompanied children) who have named a GP at one of the participating health centers, and all professionals working in the selected health centers during the study period, including doctors, nurses, administrative staff, and others.

### What does the study involve?

This is an observational study, which means no treatment or care will be changed. Instead, the study will collect information through:

- Questionnaires filled out by patients about their experience and satisfaction with care.
- Questionnaires filled out by healthcare professionals about how their teams are organized and how they feel about their work.
- Social and demographic data, including a measure of patient vulnerability using the EPICES score (a tool used in France to assess social disadvantage).

### What are the possible benefits and risks of participating?

Benefits include helping researchers and policymakers better understand what makes care effective and accessible, especially for vulnerable patients. Participating centers may also benefit from feedback on how to improve teamwork and patient satisfaction.

There are no medical risks involved, as the study only collects survey responses and demographic information. All data will be collected anonymously and handled securely.

Where is the study run from?

The study is coordinated by the Institut Jean-François Rey (IJFR), in collaboration with the French Society of General Practice (SFMG) and academic partners such as the École des Hautes Études en Santé Publique (EHESP, Rennes) and the INSERM CEPIA research team.

When is the study starting and how long will it run?

November 2024 to June 2026

Who is funding the study?

The study is funded by the French Ministry of Health, through the Direction Générale de l'Offre de Soins (DGOS), as part of a grant awarded via the RESPIR (Regional Support for Research in Primary Care) for proposals managed by the GIRCI Île-de-France (Interregional Clinical Research and Innovation Group).

Who is the main contact?

Prof. Julien Le Breton, julien.lebreton@u-pec.fr

## Contact information

**Type(s)**

Public, Scientific, Principal investigator

**Contact name**

Prof Julien Le Breton

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**Contact details**

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## Additional identifiers

**Clinical Trials Information System (CTIS)**

Nil known

**ClinicalTrials.gov (NCT)**

Nil known

**Protocol serial number**

RESPIR-21-003

# Study information

## Scientific Title

Analysis of coordinated care organization in health centers: accessibility and quality of curative and preventive services

## Acronym

EPIDAURE-2

## Study objectives

### Primary Objective

To assess the relevance of different organizational models in health centers — including modes of interprofessional cooperation, coordination, and regulation within teams, and the role of general practitioners — in delivering coordinated healthcare.

### Secondary Objectives

- To evaluate the ability of these organizational models to provide care for socially vulnerable populations and to identify the factors associated with this capacity
- To assess the perceptions (experience, satisfaction) of healthcare professionals involved in these organizational models and identify associated factors
- To assess the perceptions (experience, satisfaction) of patients (those listed with a primary care physician) using these organizational models and identify associated factors

## Ethics approval required

Ethics approval required

## Ethics approval(s)

approved 18/03/2025, Comité Ethique du CNGE (155 rue de Charonne, Paris, 75011, France; +33 (0)175622290; comite-ethique@cnge.fr), ref: 740

## Study design

Exploratory observational study using descriptive epidemiology to analyze organizational models in health centers

## Primary study design

Observational

## Study type(s)

Other

## Health condition(s) or problem(s) studied

Access to and quality of coordinated primary care, including care for socially vulnerable populations

## Interventions

This is an observational, descriptive epidemiological study conducted in nine community health centers in the Île-de-France region, France. The study involves two participant populations:

1. Healthcare professionals (clinical and non-clinical staff) working in the centers.
2. Patients who consult with a general practitioner (GP) within these centers and have declared a GP at the center.

### **Participant involvement:**

- Healthcare professionals will complete the SAPHORA-JOB questionnaire, which collects information on their professional characteristics, experience of coordination, and job satisfaction.
- Patients will complete the EUROPEP questionnaire, providing data on their healthcare experience and satisfaction, as well as sociodemographic information and social vulnerability (EPICES score).

No intervention is delivered as part of this study. All data are collected via self-administered or assisted questionnaires during routine care.

Participation does not alter clinical care pathways.

### **Duration of participation:**

- For patients: A single-point questionnaire during the consultation period.
- For professionals: One-time data collection during the study period.

Total observation period: March 2025 – March 2026 (12 months total)

Total duration of follow-up for each participant: No follow-up; single timepoint only.

## **Intervention Type**

Other

### **Primary outcome(s)**

The relevance of organizational models in health centers will be assessed using the following tools:

1. Perceived job organization and coordination measured using the SAPHORA-JOB questionnaire among healthcare professionals at a single timepoint during the study period
2. Patient experience and satisfaction measured using the EUROPEP questionnaire among patients at a single timepoint during the study period

### **Key secondary outcome(s)**

1. Perception and experience of healthcare professionals measured using the SAPHORA-JOB questionnaire at a single timepoint.
2. Perception and experience of patients measured using the EUROPEP questionnaire at a single timepoint.
3. Patients' social vulnerability measured using the EPICES score at a single timepoint.

## **Completion date**

30/06/2026

## **Eligibility**

### **Key inclusion criteria**

#### **Health Centers:**

The study will include a total of 9 health centers in the Île-de-France region, with variation in organizational and remuneration models:

1. 2 centers with activity-based remuneration
2. 7 centers with fixed-salary remuneration

#### **Patients:**

1. Prospectively and randomly included during consultations.
2. Eligible patients are adults who have declared a primary care physician within the participating health center.

**Healthcare Professionals:**

All staff working in the participating centers during the study period will be included, whether medical or non-medical, clinical or non-clinical.

**Participant type(s)**

Patient, Health professional

**Healthy volunteers allowed**

No

**Age group**

Adult

**Lower age limit**

18 years

**Upper age limit**

120 years

**Sex**

All

**Key exclusion criteria****Patients:**

Patients who have not declared a primary care physician within the participating health center

**Healthcare Professionals:**

1. Professionals not working at the participating health centers during the study period
2. Temporary staff or visiting professionals with no regular activity in the center

**Date of first enrolment**

27/03/2025

**Date of final enrolment**

31/12/2025

## Locations

**Countries of recruitment**

France

**Study participating centre**

CMS La Courneuve

2 mail de l'égalité

La Courneuve

France

93120

**Study participating centre**

**CMS Nanterre**

8 Rue Jean-Baptiste Lebon  
Nanterre  
France  
92000

**Study participating centre**

**CMS Malakoff**

74 Av. Pierre Larousse  
Malakoff  
France  
92240

**Study participating centre**

**CMS Arcueil**

13 avenue du Chaperon Vert  
Arcueil  
France  
94110

**Study participating centre**

**CDS Richerand**

4 Av. Richerand  
Paris  
France  
75010

**Study participating centre**

**CMS Champigny**

5, rue de l'Abreuvoir  
Champigny-sur-Marne  
France  
94500

**Study participating centre**

**CMS Saint-Denis**

14, rue Henri-Barbusse

Saint-Denis  
France  
93200

**Study participating centre**

**CDS Edisson**  
44 Rue Charles Moureu  
Paris  
France  
75013

**Study participating centre**

**CDS Yvonne Pouzin**  
14 Rue Volta  
Paris  
France  
75003

## **Sponsor information**

**Organisation**

Institut Jean-François Rey

## **Funder(s)**

**Funder type**

Government

**Funder Name**

Direction Générale de l'offre de Soins

**Alternative Name(s)**

DGOS

**Funding Body Type**

Government organisation

**Funding Body Subtype**

National government

**Location**

France

## Results and Publications

**Individual participant data (IPD) sharing plan**

The datasets generated during and/or analysed during the current study are/will be available upon request from Prof. Julien Le Breton, julien.lebreton@u-pec.fr.

Type of data that will be shared:

De-identified individual-level data from questionnaires completed by patients and healthcare professionals. This includes:

- Patient responses on care experience and satisfaction (EUROPEP questionnaire)
- Professional responses on team coordination and work experience (SAPHORA-JOB questionnaire)
- Social vulnerability data (EPICES score)
- Basic demographic characteristics

When the data will become available:

Within 6 months after publication of the main study results.

For how long the data will be available:

For a period of 2 years after publication.

Access criteria:

Data will be shared with qualified researchers affiliated with academic or healthcare institutions, upon submission and approval of a data access request outlining the proposed secondary analysis.

Mechanism for data access:

Requests should be submitted by email to the study contact. A data-sharing agreement (DSA) will be required to ensure data use complies with ethical, legal, and confidentiality obligations.

Consent and anonymisation:

All identifiable information will be fully anonymised before sharing. Participants have been informed of potential secondary data use during the consent process.

Ethical or legal restrictions:

Data sharing is subject to compliance with GDPR and French data protection laws. Any qualitative data will be reviewed to ensure full de-identification prior to release.

Additional comments:

The data will not be deposited in a public repository due to confidentiality considerations, but access can be granted on a case-by-case basis through controlled procedures.

**IPD sharing plan summary**

Available on request

**Study outputs**

Output type	Details	Date created	Date added	Peer reviewed?	Patient-facing?
<a href="#"><u>Participant information sheet</u></a>	Participant information sheet	11/11/2025	11/11/2025	No	Yes