Human Research and Clinical Study Support Platform (SHREC)

What are the platform's objectives ?

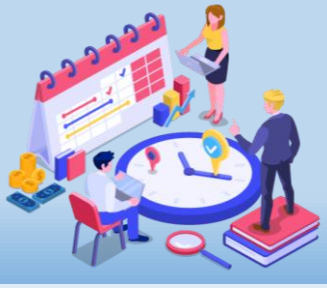
To support CNRS researchers, their academic or private partners in the development of research requiring a clinical study with:

- ✓ Evaluation of the project's feasibility : from lab to clinic
- ✓ Identification of the type of clinical study concerned
- ✓ Personalized support in the methodology (definition of the main and secondary objective, workforce, etc.), regulatory procedures and assistance in drafting the necessary documents; generally in collaboration with hospital methodologists.
- ✓ Assistance with the processing of personal data in connection with regulations (European regulation GDPR2) and the Data Protection Act (CNIL)
- ✓ Help with filing various documents with regulatory authorities (ANSM, CNIL, CPP, etc.)
- ✓ Referral to specialized contacts and service providers to continue the study after validation by the authorities
- ✓ Training and education of researchers towards clinical research
- ✓ Interfacing labs with the Research and Innovation Department of Montpellier University Hospital

Key steps:

You want to set up a study on human beings:
SHREC teams assess the feasibility of your project.

We can help you on your human clinical research studies:
Definition of primary and secondary objectives, population size, etc.



Assistance with regulatory procedures and the drafting of your documents.
Your documents are reviewed and submitted to the authorities within the allotted time.

You will be supported until the regulatory authorities have accepted your application.

SHREC Team :

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