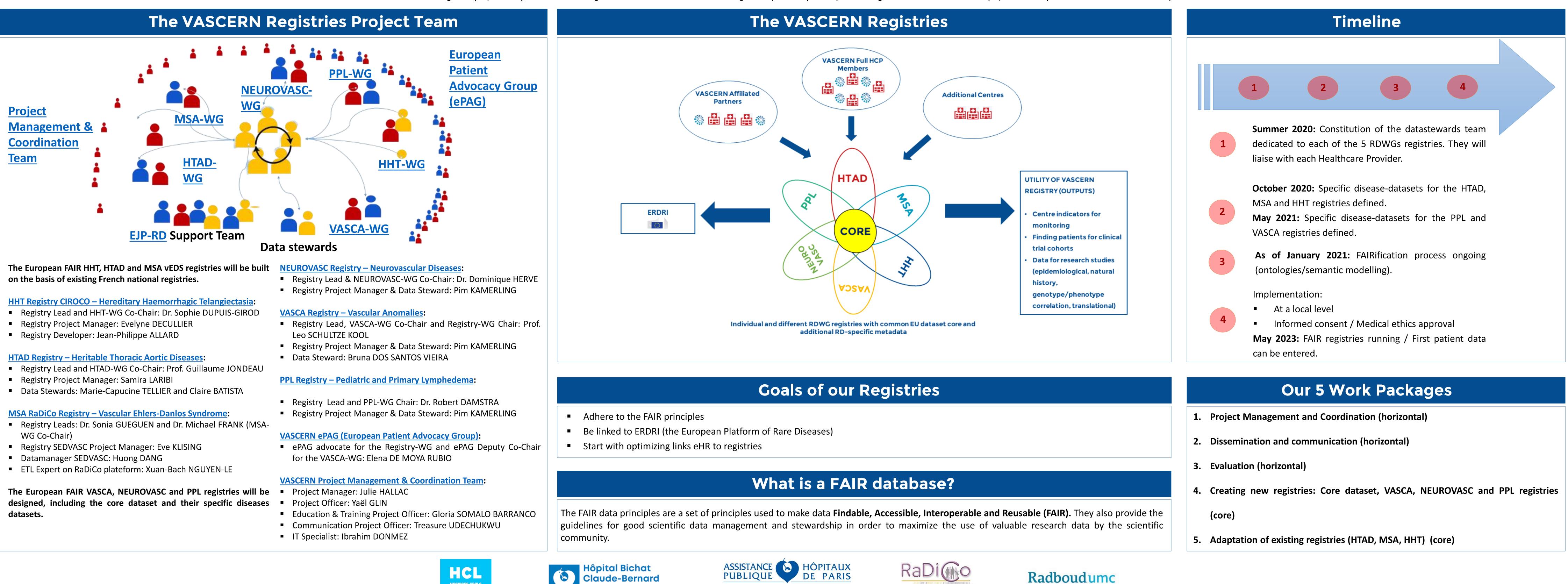




As rare diseases are rare and patient numbers are therefore often limited, sharing of data is absolutely crucial. A European-wide registry is universally recognised as beneficial for epidemiological studies, and hopefully trials of new drugs, or of repurposed drugs, in the treatment of rare diseases. With this VASCERN Registries project, we aim to reach the objective of developing our rare disease registry, with the core data elements required for all rare vascular diseases. Our project can be viewed as either a unique registry, with the core data elements required for all rare vascular disease patients and the specific disease elements varying for each 5 Rare Disease Working Groups (RDWGs), or 6 different registries with a common core making interoperability and epidemiological studies on the whole population of patients seen within the ERN possible.



VASCERN Registries EU Grant Project (2020-2023)

https://vascern.eu/what-we-do/patient-registries/



AP-HP





Vascular Diseases



F Y



Radboudumc university medical center

Gathering the best expertise in Europe to provide accessible cross-border healthcare to patients with rare vascular diseases

