



European **Rare Diseases**
Research Alliance



We are a European partnership
uniting over 170 public and private
organisations across 37 countries
around a single goal:

Advancing research in prevention, diagnosis and treatment for the **30 million people** living with a rare disease in Europe

Unify

knowledge,
resources
and expertise

Spur

innovation
and EU
competitiveness

Boost

clinical research
with and for
patients



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ERDERA provides a series of integrated services to drive rare disease research forward. The patient voice is at the heart of our mission, guiding all ERDERA services, activities and governance.

Fd

Funding

E:DERA

Cr

Clinical Research

E:DERA

Funding

ERDERA unites funding resources from public and private entities to help bridge knowledge gaps and advance rare disease research.

Clinical Research

A pan-European Clinical Research Network linking ERNs, diagnostic centres, and undiagnosed disease programmes to share data, accelerate trials and work towards ATMP readiness.

Data Hub

Connecting FAIR data sources, methods, and digital tools to maximise rare disease research through guidance, technical support, and advanced analysis.

Expertise Services

Supporting researchers with tailored services at every step of their journey; including mentoring, regulatory advice, and methodological expertise.

Dh

Data Hub

E:DERA

Es

Expertise Services

E:DERA

Te

Training & Education

E:DERA

Ea

ERDERA Accelerator

E:DERA

Ia

International Alignment

E:DERA

Training & Education

Covering lab skills, policy, leadership, clinical research, omics, data science, and entrepreneurship through its learning hub.

Accelerator

Our Public-Private Collaboration Accelerator turns ideas into investor-ready projects, offering expert support to de-risk and advance pre-clinical research.

International Alignment

We strengthen collaboration through National Mirror Groups, supporting underrepresented countries, and building global alliances.