



European
Reference
Network

Hematological Diseases
(ERN EuroBloodNet)

World Sickle Cell Awareness Day

19th June 2021

ERN-EuroBloodNet's multi level actions in the field of Sickle Cell Disease (SCD) aims at wide distribution of knowledge and cross border patients support using different approaches:

Webinars for health professionals and patients



Online educational courses targeting SCD

Visit EuroBloodNet's EDU channel on [YouTube](#) to watch videos

Preceptorships for health professionals



Short stays on SCD in expert Health Care Providers, members of the ENR-EuroBloodNet, accredited by European Board of Accreditation in Hematology (EBAH) Continuing Medical Education points

SCD patient priority blog



A [blog](#) written by people living with SCD

Training sessions for patients and caregivers



SCD Educational Sessions for people living with SCD at ASCAT [2019](#) and [2020](#)

Repository of Educational Material for Health professionals and patients



International existing documents for best practices, patient therapeutic education and development of (unique) International guidelines on clinical management

European patients' registries

ENROL – European Rare Blood Disorders Platform



RADeep – The Rare Anemia Disorders European Platform

Visit www.eurobloodnet.eu and www.radeepnetwork.eu

Supporting the Establishment of the European Network of SCD Patients Organizations



A bottom-up umbrella network of national and local organizations of SCD patients' advocates

coordination@eurobloodnet.eu
www.eurobloodnet.eu
www.eurobloodnet.eu/newsletter/



Covering more than 400 rare and complex hematological diseases, ERN-EuroBloodNet offers its online facilities, services and expertise to support orphan blood disorders.



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for rare or low prevalence
complex diseases

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