

CALL TO CONTRIBUTE TO A SICKLE HEMOGLOBINOPATHY REGISTRY

We invite Sickle Cell Disease (SCD) researchers to expand the Sickle Hemoglobinopathy registry

ELIGIBILITY

- Attend monthly online meetings from January 2025
- Complete the REDCap registration form by 15 December 2024
- Commit a Data Manager for online meetings
- Have access to at least 1,000 SCD patient research or healthcare records
- Researchers based in the following countries are not eligible to apply: Ghana; Nigeria; Mali; Tanzania; Mali; Zimbabwe; Zambia

(These countries have been excluded because they are already contributing to a multinational registry)

BENEFITS

- Become a member of the SCD Ontology (SCDO) Work Group
- Network and collaborate with other key SCD stakeholders
- Be part of future grant applications
- Get technical support from SADaCC
- Access to ethical, legal and social implication templates
- Contribute to the expansion of SCD resources and research

**COMPLETE
REGISTRATION**

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