The Sickle Cell Disease Coalition (SCDC) is an alliance of diverse stakeholders from around the world uniting to conquer sickle cell disease. Established in 2016 by the American Society of Hematology, the SCDC aims to raise awareness, improve education, expand research, and advance care for those living with sickle cell disease globally.

**SCDC Goals**

- **Broaden, amplify, and harmonize voices within the sickle cell stakeholder community**
- **Unite diverse stakeholders to lead coordinated and collaborative SCD initiatives**
- **Promote evidence-based interventions and shared decision-making related to sickle cell care**

**SCDC Committees & Current Priorities**

- **Access to Care Working Group**
  Facilitating dissemination & implementation of sickle cell materials
- **Blood Donor Diversity Task Force**
  Spreading awareness about SCD & need for more diverse blood donors
- **Global Issues Working Group**
  Documenting SCD stakeholders working locally in regions most impacted by sickle cell
- **Research & Clinical Trials Working Group**
  Helping investigators write & disseminate plain-language summaries on sickle cell research
- **Sickle Cell Trait (SCT) Task Force**
  Promoting SCT awareness, screening, documentation & high-quality genetic counseling

**SCDC Membership** = 100+ Organizations

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To learn more about SCDC and opportunities to collaborate:

- **Email**: coordinator@scdcoalition.org
- **Twitter**: @ConquerSCD
- **Website**: scdcoalition.org

Sign up for our Monthly E-Newsletter: **SCDC Update**

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**SNAPSHOT**

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