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Sickle Cell Disease: Translating clinical care to low-resource countries

through international research collaborations

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Abstract

The vast majority of the world's population of children and adults with sickle cell disease (SCD) are born in low-resource settings, particularly in sub-Saharan Africa, the Caribbean, the Middle East, and India. As a result numerous well-established, cost-effective, and evidence-based strategies for managing SCD such as

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