SPONSOR: Walton Gray

This bill establishes within the Department of Health and Senior Services the Sickle Cell Standing Committee as a subcommittee of the Missouri Genetic Disease Advisory Committee to assess the impact of sickle cell disease on urban communities, examine the existing services and resources to address the needs of people with the disease, and develop recommendations for educational services to schools on the traits of the disease and their effects.

The committee will consist one member who is a licensed physician with experience in the diagnosis and treatment of sickle cell disease and who must serve as chair of the committee, one member who has sickle cell disease or is a family member of someone with sickle cell disease, one member with expertise in sickle cell disease organization, one member from a leading sickle cell disease organization, one member with expertise in minority health, and one member from each of the hemoglobinopathy centers which contract with the department.

The committee must meet in at least three urban regions of the state to seek public input and to examine trends in the state sickle cell disease populations and their needs, existing services and resources, needed state policies or responses, and the replacement of the genetic testing and counseling program which has been eliminated due to lack of funding. A report of the findings and any recommendations by the committee must be submitted to the Governor and General Assembly by December 31, 2015. The committee must continue to meet at the request of the chair and at least once each year after that date and submit an annual supplemental report.