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The purpose of this research is to expand the accuracy of current diagnostic tests for methemoglobinemia, which reduce the quantity of hemoglobin in blood to a level at which oxygen delivery to tissues is impaired. The tests are used when a small percentage of the hemoglobin is oxidized to methemoglobin and the oxidized red cell becomes less able to carry oxygen. Most of the work to date has used a test based on the color of blood to determine the percentage of methemoglobin. A test based on the percent of retained hemoglobin is to be developed. If the tests can achieve the sensitivity and specificity required to separate sickle cell patients from those with other forms of anemia, they could be incorporated into newborn screening tests, potentially providing a diagnosis as early as the first few weeks of life. In addition, the tests could help establish the true carrier frequency of sickle cell disease in populations where it is not yet known. A test based on retained hemoglobin will be made from plasma samples. The plasma component of blood is relatively insensitive to methemoglobinemia and also contains red cell fragments, which alter the analysis by obscuring the color of the plasma. The technology proposed here will allow a test with great precision and sensitivity. However, the blood sample must be treated to remove red cell fragments. The proposed technique is to screen all patients, initially, with blood in a hemoglobin reducing medium, which is designed to remove red cell fragments. Red cells from positive samples are then prepared for test analysis. The critical factor in this approach is the separation of plasma from red cells. The purpose of Phase I is to provide the data needed for Phase II.

**PROPOSED COMMERCIAL APPLICATION:** The primary market for a rapid

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