

Sickle cell disease or anaemia (SCD) is common life threatening hereditary disease worldwide. With its occurrence, in another two decades, over twelve million people are estimated to suffer from sickle cell anaemia and over thirty million people are likely to inherit the trait.

High degree of marital consanguinity, caste/class and geographical endogamy, lack of medical facilities, psychosocial prejudices, certain irrational traditions and beliefs aggravating the nutritional status and health are the salient features that need to be tackled to put a pause to the malady of sickle cell disease.

Regular screening with a reliable diagnostic method is very critical to improve the life of person with SCD. HPLC method is considered, most reliable and Gold standard method for diagnosis of Sickle cell anaemia. But Testing with HPLC method in resource limited setting is very difficult and most of the time, impossible. So need of the hour is a simple Point-of-care rapid test with comparable result with HPLC.

Tulip introduces **SICKLECHECK**[™] an innovative Point-of-care rapid immunoassay for the detection of sickle cell disease and trait.

SICKLECHECK

Utilizes monoclonal anti HbA & anti HbS antibodies Facilitates specific detection of HbA & HbS

Facilitates classification of sickle cell trait, disease Differentiates HbA & HbS.

Competitive immunochromatographic assay No risk of Prozoning

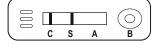
Evaluated by various Institutions; Sensitivity 99%, Specificity 99.1% compared with HPLC methods*
Reliable results

Small specimen volume & simple test procedure Suitable for Point-of-care/Field testing setup

Storage at 4°C- 40°C Suitable for most climatic conditions



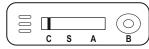
Result Interpretation



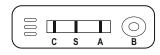




Sickle Cell Disease



Sickle Cell Trait



Other Hemoglobinopathies



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