Sickle Cell-Hemoglobin D Disease

National Cancer Institute

Source

National Cancer Institute. Sickle Cell-Hemoglobin D Disease. NCI Thesaurus. Code C155310.

A variant of sickle cell disease due to heterozygosity for hemoglobin S and hemoglobin D mutations. Patients present with the symptoms of sickle cell disease but the symptoms are less frequent and severe compared to patients with hemoglobin SS disease.