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Sickle Cell Disease IkeOluwa Lagunju,Manoelle Kossorotoff,Fenella Jane Kirkham,Andria Ford,2024-03-05 Sickle cell disease SCD is the most common cause of ischemic and hemorrhagic stroke in children and without treatment has a very high rate of recurrence Acute neurological symptoms and signs are common in SCD at any age and as well as stroke include transient ischaemic attack headaches seizures and coma An altered mental status with or without reduced level of consciousness headache seizures visual loss or focal signs can occur spontaneously but also in numerous contexts such as infection acute chest syndrome ACS and acute anaemia Nearly 50 years ago it was clear from conventional cerebral angiography studies that the large extracranial and intracranial vessels were abnormal in patients with SCD and neurologic deficits although the pathophysiology is not fully understood from human studies or laboratory models Transcranial Doppler ultrasound has been used successfully to predict stroke risk in children although those with abnormal time averaged maximum mean velocity 200 cm sec may have very high blood flow or arterial stenosis Although there are relatively few studies with matched controls silent cerebral infarction on MRI is found in a high proportion of patients with SCD without clinical symptoms sometimes starting as early as the 6th month of life steadily accumulating with age and predicting overt stroke Meta analyses have shown that typically siblings children and adults with SCD have cognitive difficulties over a wide range of domains compared with matched controls This includes processing speed and executive function

iPad for Beginners ,2013

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