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Titolo	Sickle Cell Disease in Clinical Practice // by Jo Howard, Paul Telfer
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ISBN	1-4471-2473-1
Edizione	[1st ed. 2015.]
Descrizione fisica	1 online resource (300 p.)
Collana	In Clinical Practice, , 2199-6652
Disciplina	610 615.39 616.07 616.15
Soggetti	Hematology Pathology Blood - Transfusion Blood Transfusion Medicine
Lingua di pubblicazione	Inglese
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Livello bibliografico	Monografia
Note generali	Description based upon print version of record.
Nota di bibliografia	Includes bibliographical references and index.
Nota di contenuto	Overview of Sickle Cell Disease -- Laboratory Tests Used in Diagnosis and Monitoring of Sickle Cell Disease -- Organization of Care for Sickle Cell Disease -- Overview and general principles -- Pain in Sickle Cell Disease -- Respiratory and Cardiac Complications in Sickle Cell Disease -- Neurological Complications of Sickle Cell Disease -- Renal and Urological Complications in Sickle Cell Disease -- Bone and Joint Complications in Sickle Cell Disease -- Ophthalmological complications in Sickle Cell Disease -- The Spleen in Sickle Cell Disease -- Infection and Infection Prophylaxis in Sickle Cell Disease -- Gastroenterological Complications in Sickle Cell Disease -- Anemia and Sickle Cell Disease -- Leg Ulceration in Sickle Cell Disease -- Management of Pregnancy in Sickle Cell Disease -- Surgical Management of Patients with Sickle Cell Disease -- Sickle Cell Disease Treatment Modalities -- Out-Patient Management of Sickle Cell Disease Patients.
Sommario/riassunto	This practical clinical handbook reveals that sickle cell disease (SCD) is an increasingly common condition to manage in Europe and North America. SCD demands clinical expertise and experience as well as sensitivity to its social and cultural context. This book is designed to

broaden readers' knowledge in this challenging condition by describing the acute and long-term complications unique to SCD and that affect nearly every system of the body. Critically, it also details the significant recent advances in understanding the pathophysiology of SCD that are leading to novel treatment modalities. Sickle Cell Disease in Clinical Practice promotes higher quality care by outlining the clinical problems as they arise, and covering essential background information, including up-to-date research, and useful points to guide management. As such, the intended target audience is broad and includes general physicians, general practitioners, hematologists, pediatricians, emergency medicine physicians, surgeons, medical students, nurse specialists and commissioners.
