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Titolo	Congenital Bleeding Disorders : Diagnosis and Management / / edited by Akbar Dorgalaleh
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Descrizione fisica	1 online resource (XVI, 463 p. 102 illus., 95 illus. in color.)
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Nota di bibliografia	Includes bibliographical references.
Nota di contenuto	An Overview of Hemostasis -- Congenital Bleeding Disorders: Diagnosis and Management -- von Willebrand disease: an update on diagnosis and treatment -- Hemophilia A, Diagnosis and Management -- Hemophilia B, Diagnosis and Management -- Congenital fibrinogen disorders, diagnosis, and management -- Congenital Prothrombin Deficiency: Diagnosis and Management -- Congenital Factor V Deficiency -- Combined Factor V and Factor VIII Deficiency, Diagnosis and Management -- Congenital Factor VII Deficiency, Diagnosis and Management -- Vitamin K-dependent Coagulation Factors Deficiency, Diagnosis and Management -- Congenital Factor X deficiency, Diagnosis and Management -- Congenital Factor XI Deficiency -- Congenital Factor XIII Deficiency, Diagnosis and Management -- Glanzmann Thrombasthenia -- Bernard-Soulier Syndrome, Diagnosis and Management -- Gray Platelet Syndrome (GPS). .
Sommario/riassunto	This significantly updated new edition describes in detail the clinical presentations, diagnosis, and management of a wide range of congenital bleeding disorders. It will assist readers in overcoming the significant challenges involved in clinical and laboratory diagnosis and in providing effective clinical care that makes optimal use of new products, including recombinant factor concentrate. The coverage ranges from hemophilia A and B and von Willebrand disease to rare

bleeding disorders such as congenital factor V, factor X, factor XI, and factor XIII deficiency and inherited platelet function disorders. The exceptional attention to rarer conditions is of particular importance given the considerable risk of overlooking them during diagnosis, with potential consequences for disease-related morbidity and mortality. The authors are acknowledged specialists in the field from across the world who have particular expertise in the disorder that they discuss. The book will be of value tohematologists, oncologists, pediatricians, laboratory specialists and technicians, general physicians, and trainees.