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Haemophiliacs; DDAVP for Treatment of Mild Haemophilia during Surgery; Haemophilia and Haemodialysis; Haemophilia and Hepatitis C Treatment; Haemophilia and Physical Therapy; Haemophilia and Renal Bleeds; Haemophilia and Scuba Diving; Haemophilia and Ventricular Septal Defect Repair
Haemophilia with Hepatitis C and Recurrent Bleeding; Isotretinoin in Haemophilia; Laser Eye Surgery in a Haemophiliac; Managing Haemophilic Pseudotumours; Continuous NovoSeven: Pros and Cons; Complications of Treatment; Haemophilia B and Immune Tolerance with Anaphylaxis; rFVIIa (NovoSeven); Thrombosis in PCCs vs. APCCs; 2 Von Willebrand Disease; Epidurals and VWD; Anticoagulation for a Cardiac Valve in a Patient with VWD Type 1; VWD Type 2A and Pregnancy; VWD Type 2B and Pregnancy; VWD Type 2B vs. Platelet Type; Prophylaxis in VWD Type 3; Platelet Type VWD; 3 Factor Deficiencies
Combined Factor V and VIII Deficiency; DVT Prophylaxis in FVII Deficiency; Menorrhagia in Factor VII Deficiency; Anticoagulation for Atrial Fibrillation in FX Deficiency; Management of Factor X Deficiency; Anticoagulation for Atrial Fibrillation in a Patient with Factor XI Deficiency; Factor XI Deficiency and Surgery; Prophylaxis for Patients with Factor XIII Deficiency and Intracranial Bleeding; 4 Rare Platelet and Coagulation Disorders; Coagulation Disorders; Afibrinogenaemia; Dysfibrinogenaemia; Hypofibrinogenaemia; Gardner-Diamond Syndrome; Hereditary Haemorrhagic Telangiectasia
Hereditary Vitamin-K-Dependent Coagulation Factors Deficiency and Pregnancy; Platelet Disorders; Glanzmann's Thrombasthaenia and Gastrointestinal Angiodysplasia; Glanzmann's Thrombasthaenia and Pregnancy; rVIIa (NovoSeven®) and Wiskott-Aldrich Syndrome; 5 Acquired Bleeding Diatheses; Acquired Haemophilia and Second Pregnancy; Developing a Factor IX Inhibitor; End-Stage Liver Disease and Surgery; Treatment for Acute DIC; 6 Miscellaneous Questions; Bleeding Time vs. PFA-100; Cocaine and DDAVP; Hyponatraemia and DDAVP; The Use of the INR
Selective Serotonin Reuptake Inhibitors and Clotting Disturbances

Sommario/riassunto

This unique book covers the major cases one might encounter in diagnosing, managing and treating haemophilia. It will provide a practical and informative guide to the broad range of topics concerning both bleeding and clotting disorders. Haemophilia and Haemostasis: A Case-based Approach to Management is divided into major chapter sections by disorder. Each chapter contains questions and cases that were originally submitted to the online Haemostasis-forum (previously Haemophilia-forum). Cases and questions associated with each disorder are presented alongside practical answers from a wide
