

Hemophilias and von Willebrand's Disease

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REFERENCES

1. Fijnvandraat K, Cnossen MH, Leebeek FW, Peters M: Diagnosis and management of haemophilia. *BMJ* 344: e2707, 2012. [PMID: 22551712]
2. Srivastava A, Brewer AK, Mauser-Bunschoten EP, et al: Guidelines for the management of hemophilia. *Haemophilia* 19: e1, 2013. [PMID: 22776238]
3. Konkle BA, Huston H, Nakaya Fletcher S. Hemophilia A: In: Adam MP, Ardinger HH, Pagon RA, et al (eds): GeneReviews® [Internet]. Seattle, WA: University of Washington, Seattle; 1993-2018. [updated June 22, 2017]. [PMID: 20301578]
4. Goodeve A, James P. von Willebrand disease. In: Adam MP, Ardinger HH, Pagon RA, et al (eds): GeneReviews® [Internet]. Seattle, WA: University of Washington, Seattle; 1993-2018. [updated October 5, 2017]. [PMID: 20301765]
5. Di Michele DM, Gibb C, Lefkowitz JM, Ni Q, Gerber LM, Ganguly A: Severe and moderate haemophilia A and B in US females. *Haemophilia* 20: e136, 2014. [PMID: 24533955]
6. <https://www.hemophilia.org/node/3652>. (National Hemophilia Foundation: Guidelines for emergency department management of individuals with hemophilia and other bleeding disorders, September 17, 2017.) Accessed September 1, 2018.
7. Singleton T, Kruse-Jarres R, Leissinger C: Emergency department care for patients with hemophilia and von Willebrand disease. *J Emerg Med* 39: 158, 2010. [PMID: 18757163]
8. Vanderhave KL, Caird MS, Hake M, et al: Musculoskeletal care of the hemophiliac patient. *J Am Acad Orthop Surg* 20: 553, 2012. [PMID: 22941798]
9. <http://www1.wfh.org/publications/files/pdf-1271.pdf>. (Farrugia A, World Federation of Hemophilia: *Guide for the Assessment of Clotting Factor Concentrates*, 3rd ed, 2017. World Federation of Hemophilia.) Accessed September 1, 2018
10. <http://elearning.wfh.org/resource/online-cfc-registry/>. (World Federation of Hemophilia: On-line Registry of Clotting Factor Concentrates.) Accessed September 1, 2018.
11. <https://www.hemophilia.org/node/3675>. (National Hemophilia Foundation, Medical and Scientific Advisory Council: MASAC recommendations concerning products licensed for the treatment of hemophilia and other bleeding disorders; revised April 23, 2018.) Accessed September 1, 2018.
12. Mannucci PM: Plasma-derived versus recombinant factor VIII concentrates for the treatment of haemophilia A: plasma-derived is better. *Blood Transfus* 8: 288, 2010. [PMID: 20967171]
13. Franchini M: Plasma-derived versus recombinant factor VIII concentrates for the treatment of haemophilia A: recombinant is better. *Blood Transfus* 8: 292, 2010. [PMID: 20967172]
14. Berntorp E, Bjorkman S: The pharmacokinetics of clotting factor therapy. *Haemophilia* 9: 353, 2003. [PMID: 12828668]
15. Forsyth AL, Zourikian N, Valentino LA, Rivard GE: The effect of cooling on coagulation and haemostasis: should "Ice" be part of treatment of acute haemarthrosis in haemophilia? *Haemophilia* 18: 843, 2012. [PMID: 22913708]
16. Benson G, Auerswald G, Dolan G, et al: Diagnosis and care of patients with mild haemophilia: practical recommendations for clinical management. *Blood Transfus* 14: 1, 2017. [PMID: 29328905]
17. Franchini M, Zaffanello M, Lippi G: The use of desmopressin in mild hemophilia A. *Blood Coagul Fibrinolysis* 21: 615, 2010. [PMID: 20829683]
18. <http://elearning.wfh.org/resource/desmopressin-ddavp-in-the-treatment-of-bleeding-disorders/>. (Mannucci PM: Desmopressin (DDAVP) in the treatment of bleeding disorders. treatment of hemophilia. revised edition. No 11. Montreal, Quebec, Canada: World Federation of Hemophilia, 2012.) Accessed September 1, 2018.
19. Quon DV, Konkle BA: How we treat: haematuria in adults with haemophilia. *Haemophilia* 16: 683, 2010. [PMID: 20041958]
20. Astermark J: Overview of inhibitors. *Semin Hematol* 43(suppl 4): S3, 2006. [PMID: 16690373]
21. Rocino A, Franchini M, Coppola A: Treatment and prevention of bleeds in haemophilia patients with inhibitors to Factor VIII/IX. *J Clin Med* 6: E46, 2017. [PMID: 28420167]
22. Rodgers GM: Prothrombin complex concentrates in emergency bleeding disorders. *Am J Hematol* 87: 898, 2012. [PMID: 22648513]
23. Ljung RCR: How I manage patients with inherited haemophilia A and B and factor inhibitors. *Br J Haematol* 180: 501, 2018. [PMID: 29270992]
24. Weber KE: Acquired hemophilia A. *Semin Thromb Hemost* 38: 735, 2012. [PMID: 22941793]
25. Kruse-Jarres R, Kempton CL, Baudo F, et al: Acquired hemophilia A: updated review of evidence and treatment guidance. *Am J Hematol* 92: 695, 2017. [PMID: 28470674]
26. Seethala S, Gaur S, Enderton E, Corral J: Postpartum acquired hemophilia: a rare cause of postpartum hemorrhage. *Case Rep Hematol* 2013: 735715, 2013. [PMID: 23533849]
27. Barg AA, Livnat T, Kenet G: An extra X does not prevent acquired hemophilia: pregnancy-associated acquired hemophilia A. *Thromb Res* 151(suppl 1): S82, 2017. [PMID: 28262242]
28. Hassan MI, Saxena A, Ahmad F: Structure and function of von Willebrand factor. *Blood Coagul Fibrinolysis* 23: 11, 2012. [PMID: 22089939]
29. Rao ES, Ng CJ: Current approaches to diagnostic testing in von Willebrand disease. *Transfus Apher Sci* 57: 463, 2018. [PMID: 30064913]
30. Nichols WL, Hultin MB, James AH, et al: von Willebrand disease (VWD): evidence-based diagnosis and management guidelines, the National Heart, Lung, and Blood Institute (NHLBI) Expert Panel report (USA). *Haemophilia* 14: 171, 2008. [PMID: 18315614]
31. Castaman G: Treatment of von Willebrand disease with FVIII/VWF concentrates. *Blood Transfus* 9(suppl 2): s9, 2011. [PMID: 21839030]
32. Mannucci PM, Federici AB, James AH, Kessler CM: von Willebrand disease in the 21st century: current approaches and new challenges. *Haemophilia* 15: 1154, 2008. [PMID: 19624761]