

von Willebrand disease (VWD): evidence-based diagnosis  
National Heart, Lung, and Blood Institute (NHLBI) Expert

Haemophilia

14, 171-232

DOI: [10.1111/j.1365-2516.2007.01643.x](https://doi.org/10.1111/j.1365-2516.2007.01643.x)

Citation Report

#	ARTICLE	IF	CITATIONS
1	A highly-sensitive plasma von Willebrand factor ristocetin cofactor (VWF:RCo) activity assay by flow cytometry. <i>Journal of Thrombosis and Haemostasis</i> , 2008, 6, 323-330.	1.9	42
2	Acquired von Willebrand syndrome. <i>Haemophilia</i> , 2008, 14, 856-858.	1.0	2
3	Consideration of platelet function disorders in patients with reduced VWF levels. <i>Haemophilia</i> , 2008, 14, 1131-1132.	1.0	1
4	A phase II prospective open-label escalating dose trial of recombinant interleukin-11 in mild von Willebrand disease. <i>Haemophilia</i> , 2008, 14, 968-977.	1.0	31
7	Bleeding symptom assessment and hemostasis evaluation of menorrhagia. <i>Current Opinion in Hematology</i> , 2008, 15, 465-472.	1.2	17
8	A consensus statement on the management of pregnancy and delivery in women who are carriers of or have bleeding disorders. <i>Medical Journal of Australia</i> , 2009, 191, 460-463.	0.8	26
9	Advocacy for Women with Bleeding Disorders. , 0, , 176-183.		1
11	Molecular Basis of Hemostatic and Thrombotic Diseases. , 2009, , 247-264.		0
12	An Adolescent With a History of Menorrhagia. <i>Laboratory Medicine</i> , 2009, 40, 271-273.	0.8	0
13	Low von Willebrand factor: sometimes a risk factor and sometimes a disease. <i>Hematology American Society of Hematology Education Program</i> , 2009, 2009, 106-112.	0.9	86
14	Dominant von Willebrand Disease Type 2A Groups I and II due to Missense Mutations in the A2 Domain of the von Willebrand Factor Gene: Diagnosis and Management. <i>Acta Haematologica</i> , 2009, 121, 154-166.	0.7	7
15	Managing Patients with von Willebrand Disease Type 1, 2 and 3 with Desmopressin and von Willebrand Factor-Factor VIII Concentrate in Surgical Settings. <i>Acta Haematologica</i> , 2009, 121, 167-176.	0.7	22
16	The safety of plasma-derived von Willebrand/factor VIII concentrates in the management of inherited von Willebrand disease. <i>Expert Opinion on Drug Safety</i> , 2009, 8, 203-210.	1.0	16
17	Optimizing treatment of von Willebrand disease by using phenotypic and molecular data. <i>Hematology American Society of Hematology Education Program</i> , 2009, 2009, 113-123.	0.9	17
18	Toward a New Paradigm for the Identification and Functional Characterization of von Willebrand Disease. <i>Seminars in Thrombosis and Hemostasis</i> , 2009, 35, 060-075.	1.5	35
19	Factor VIII-von Willebrand Factor Complex Inhibits Osteoclastogenesis and Controls Cell Survival. <i>Journal of Biological Chemistry</i> , 2009, 284, 31704-31713.	1.6	58
20	Protein Replacement Therapy and Gene Transfer in Canine Models of Hemophilia A, Hemophilia B, von Willebrand Disease, and Factor VII Deficiency. <i>ILAR Journal</i> , 2009, 50, 144-167.	1.8	71
21	INHERITED BLEEDING DISORDERS IN PREGNANCY. <i>Fetal and Maternal Medicine Review</i> , 2009, 20, 205-227.	0.3	1

#	ARTICLE	IF	CITATIONS
22	Von Willebrand disease and other bleeding disorders in women: consensus on diagnosis and management from an international expert panel. American Journal of Obstetrics and Gynecology, 2009, 201, 12.e1-12.e8.	0.7	130
23	Clinical and laboratory diagnosis of von Willebrand disease: A synopsis of the 2008 NHLBI/NIH guidelines. American Journal of Hematology, 2009, 84, 366-370.	2.0	88
24	Considerations for epidural anesthesia in a patient with type 1 von Willebrand disease. Journal of Anesthesia, 2009, 23, 597-600.	0.7	8
25	Applying diagnostic criteria for type 1 von Willebrand disease to a pediatric population. Pediatric Blood and Cancer, 2009, 52, 102-107.	0.8	13
26	Diagnosis of type 1 VWD: Can the clinical history trump laboratory findings?. Pediatric Blood and Cancer, 2009, 52, 7-8.	0.8	1
27	Transient neonatal acquired von Willebrand syndrome due to transplacental transfer of maternal monoclonal antibodies. Pediatric Blood and Cancer, 2009, 53, 655-657.	0.8	13
28	von Willebrand factor content in Alphanate <sup>®</sup> (Laurence J. Logan) â€“ reply. Haemophilia, 2009, 15, 370-371.	1.0	0
29	Registry of inherited coagulopathies in Brazil: first report. Haemophilia, 2009, 15, 142-149.	1.0	30
30	von Willebrand disease in the 21st century: current approaches and new challenges. Haemophilia, 2009, 15, 1154-1158.	1.0	11
31	Bleeding symptoms and laboratory correlation in patients with severe von Willebrand disease. Haemophilia, 2009, 15, 918-925.	1.0	30
32	Multisite management study of menorrhagia with abnormal laboratory haemostasis: a prospective crossover study of intranasal desmopressin and oral tranexamic acid. British Journal of Haematology, 2009, 145, 212-220.	1.2	117
33	Comparison of coagulation factor XIII content and concentration in cryoprecipitate and freshâ€“frozen plasma. Transfusion, 2009, 49, 765-770.	0.8	58
34	Limitations of the ristocetin cofactor assay in measurement of von Willebrand factor function. Journal of Thrombosis and Haemostasis, 2009, 7, 1832-1839.	1.9	60
35	Desmopressin therapy to assist the functional identification and characterisation of von Willebrand disease: Differential utility from combining two (VWF:CB and VWF:RCo) von Willebrand factor activity assays?. Thrombosis Research, 2009, 123, 862-868.	0.8	36
36	Guidelines for bleeding disorders in women. Thrombosis Research, 2009, 123, S124-S128.	0.8	11
37	Classification of inherited von Willebrand disease and implications in clinical practice. Thrombosis Research, 2009, 124, S2-S6.	0.8	26
38	Laboratory Testing for Von Willebrand Disease: Toward a Mechanism-Based Classification. Clinics in Laboratory Medicine, 2009, 29, 193-228.	0.7	8
39	Alergia de las vÃas respiratorias. Medicine, 2009, 10, 2264-2270.	0.0	0

#	ARTICLE	IF	CITATIONS
40	Protocolo terapéutico del asma extrínseca. <i>Medicine</i> , 2009, 10, 2301-2304.	0.0	0
41	Global Hemostasis Testing Thromboelastography: Old Technology, New Applications. <i>Clinics in Laboratory Medicine</i> , 2009, 29, 391-407.	0.7	130
42	Potential supplementary utility of combined PFA-100 and functional von Willebrand factor testing for the laboratory assessment of desmopressin and factor concentrate therapy in von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2009, 20, 475-483.	0.5	23
43	Perioperative Management of a Child with von Willebrand Disease Undergoing Surgical Repair of Craniostynosis: Looking at Unusual Targets. <i>Anesthesia and Analgesia</i> , 2009, 109, 720-724.	1.1	3
44	Diagnosis of Von Willebrand Disease in Children. <i>Current Pediatric Reviews</i> , 2009, 5, 169-175.	0.4	1
45	von Willebrand factor to the rescue. <i>Blood</i> , 2009, 113, 5049-5057.	0.6	138
46	How I treat von Willebrand disease. <i>Blood</i> , 2009, 114, 1158-1165.	0.6	79
47	Von Willebrand factor/factor VIII concentrates in the treatment of von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2009, 20, 89-100.	0.5	35
48	Common VWF exon 28 polymorphisms in African Americans affecting the VWF activity assay by ristocetin cofactor. <i>Blood</i> , 2010, 116, 280-286.	0.6	148
49	The dominant-negative von Willebrand factor gene deletion p.P1127_C1948delinsR: molecular mechanism and modulation. <i>Blood</i> , 2010, 116, 5371-5376.	0.6	23
53	Biological and clinical response to desmopressin (DDAVP) in a retrospective cohort study of children with low von Willebrand factor levels and bleeding history. <i>Thrombosis and Haemostasis</i> , 2010, 104, 984-989.	1.8	20
55	Enzymes that hydrolyze adenine nucleotides in platelets and polymorphisms in the $\beta 2$ gene of integrin $\alpha 2$ in patients with von Willebrand disease. <i>Molecular and Cellular Biochemistry</i> , 2010, 340, 249-256.	1.4	1
56	von Willebrand disease and pregnancy: a practical approach for the diagnosis and treatment. <i>American Journal of Obstetrics and Gynecology</i> , 2010, 203, 194-200.	0.7	206
57	The genetic basis of von Willebrand disease. <i>Blood Reviews</i> , 2010, 24, 123-134.	2.8	145
58	Genetic testing for von Willebrand disease: the case for. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 13-16.	1.9	26
59	Absent collagen binding in a VWF A3 domain mutant: utility of the VWF:CB in diagnosis of VWD. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 1431-1433.	1.9	74
60	von Willebrand factor variant p.Arg924Gln marks an allele associated with reduced von Willebrand factor and factor VIII levels. <i>Journal of Thrombosis and Haemostasis</i> , 2010, 8, 1986-1993.	1.9	19
61	Clinical efficacy and safety of the factor VIII/von Willebrand factor concentrate BIOSTATE <sup>®</sup> in patients with von Willebrand disease: a prospective multicentre study. <i>Haemophilia</i> , 2010, 16, 615-624.	1.0	37

#	ARTICLE	IF	CITATIONS
62	Thrombin time and fibrinogen as initial screening tests for people with inherited bleeding disorders. <i>Haemophilia</i> , 2010, 16, 700-701.	1.0	1
63	Building our global family – achieving treatment for all. <i>Haemophilia</i> , 2010, 16, 1-10.	1.0	11
64	Women and bleeding disorders. <i>Haemophilia</i> , 2010, 16, 160-167.	1.0	73
65	Molecular Basis of Hemostatic and Thrombotic Diseases. , 2010, , 175-188.		0
67	Prospective study of low-dose ristocetin-induced platelet aggregation to identify type 2B von Willebrand disease (VWD) and platelet-type VWD in children. <i>Thrombosis and Haemostasis</i> , 2010, 104, 1158-1165.	1.8	20
68	Porcine and Canine von Willebrand Factor and von Willebrand Disease: Hemostasis, Thrombosis, and Atherosclerosis Studies. <i>Thrombosis</i> , 2010, 2010, 1-11.	1.4	22
69	Role of glycoprotein Ib mobility in platelet function. <i>Thrombosis and Haemostasis</i> , 2010, 103, 1033-1043.	1.8	17
70	A randomised pilot trial of the anti-von Willebrand factor aptamer ARC1779 in patients with type 2b von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2010, 104, 563-570.	1.8	71
71	Targeting von Willebrand factor and platelet glycoprotein Ib receptor. <i>Expert Review of Cardiovascular Therapy</i> , 2010, 8, 1689-1701.	0.6	68
72	Bleeding disorders in the tribe: result of consanguineous in breeding. <i>Orphanet Journal of Rare Diseases</i> , 2010, 5, 23.	1.2	43
73	Obstetric Management of Adolescents with Bleeding Disorders. <i>Journal of Pediatric and Adolescent Gynecology</i> , 2010, 23, S31-S37.	0.3	12
74	von Willebrand Disease in the Pediatric and Adolescent Population. <i>Journal of Pediatric and Adolescent Gynecology</i> , 2010, 23, S3-S10.	0.3	28
75	Clinical practice guidelines on menorrhagia: management of abnormal uterine bleeding before menopause. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2010, 152, 133-137.	0.5	142
76	Plasma von Willebrand factor multimer quantitative analysis by in-gel immunostaining and infrared fluorescent imaging. <i>Thrombosis Research</i> , 2010, 126, 543-549.	0.8	27
78	Comprehensive Treatment of Periodontitis in Patients With von Willebrand Disease. <i>Journal of Periodontology</i> , 2010, 81, 1432-1440.	1.7	11
79	Gynaecological and obstetric bleeding in moderate and severe von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2011, 106, 885-892.	1.8	64
80	Hypertrophic Obstructive Cardiomyopathy, Acquired von Willebrand Syndrome, and Gastrointestinal Bleeding. <i>Mayo Clinic Proceedings</i> , 2011, 86, 181-182.	1.4	11
81	Hypertrophic Obstructive Cardiomyopathy, Bleeding History, and Acquired von Willebrand Syndrome: Response to Septal Myectomy. <i>Mayo Clinic Proceedings</i> , 2011, 86, 219-224.	1.4	42

#	ARTICLE	IF	CITATIONS
82	von Willebrand disorder. Paediatrics and Child Health (United Kingdom), 2011, 21, 348-352.	0.2	1
83	Inherited disorders of coagulation. , 2011, , 547-564.		0
84	Von Willebrand Disease in Pregnancy. Hematology/Oncology Clinics of North America, 2011, 25, 335-358.	0.9	13
85	Evaluation and management of acute menorrhagia in women with and without underlying bleeding disorders: consensus from an international expert panel. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2011, 158, 124-134.	0.5	108
86	von Willebrand disease. Genetics in Medicine, 2011, 13, 365-376.	1.1	87
87	General anesthesia for cesarean delivery at a tertiary care hospital from 2000 to 2005: a retrospective analysis and 10-year update. International Journal of Obstetric Anesthesia, 2011, 20, 10-16.	0.2	97
88	The Adolescent with Menorrhagia: Why, Who, and How to Evaluate for a Bleeding Disorder. Journal of Pediatric and Adolescent Gynecology, 2011, 24, 228-230.	0.3	10
89	Hematologic Disease. , 2011, , 261-280.		0
90	Consensus statement by hospital based dentists providing dental treatment for patients with inherited bleeding disorders*. Australian Dental Journal, 2011, 56, 221-226.	0.6	30
91	Perioperative Management of Acquired von Willebrand Disease in Cardiac Surgery: Type 2B or Not 2B?. Journal of Cardiothoracic and Vascular Anesthesia, 2011, 25, 131-133.	0.6	9
92	ARFI Ultrasound Monitoring of Hemorrhage and Hemostasis In Vivo in Canine Von Willebrand Disease and Hemophilia. Ultrasound in Medicine and Biology, 2011, 37, 2126-2132.	0.7	13
94	Hemostatic Disorders. , 2011, , 378-418.		0
95	Phase II prospective open-label trial of recombinant interleukin-11 in women with mild von Willebrand disease and refractory menorrhagia. Thrombosis and Haemostasis, 2011, 106, 641-645.	1.8	24
96	Treatment of patients with von Willebrand disease. Journal of Blood Medicine, 2011, 2, 49.	0.7	10
97	Efficacy and safety of a new generation von Willebrand factor/factor VIII concentrate (Wilate®) in the management of perioperative haemostasis in von Willebrand disease patients undergoing surgery. Thrombosis and Haemostasis, 2011, 105, 1072-1079.	1.8	46
98	Management of the Medically Compromised Patient: Hematologic Disorders, Cancer, Hepatitis, and AIDS. , 2011, , 487-509.		1
99	Long-term secondary prophylaxis in children, adolescents and young adults with von Willebrand disease. Thrombosis and Haemostasis, 2011, 105, 597-604.	1.8	38
100	von Willebrand disease. Hematologie, 2011, 17, 278-288.	0.0	4

#	ARTICLE	IF	CITATIONS
102	Secondary postpartum hemorrhage due to uterine artery pseudoaneurysm rupture in von Willebrand disease. <i>Journal of Obstetrics and Gynaecology Research</i> , 2011, 37, 1887-1890.	0.6	10
103	Acquired von Willebrand Syndrome in congenital heart disease: does it promote an increased bleeding risk?. <i>British Journal of Haematology</i> , 2011, 155, 622-624.	1.2	17
104	Haemophilia: provision of factors and novel therapies: World Federation of Hemophilia goals and achievements. <i>British Journal of Haematology</i> , 2011, 154, 704-714.	1.2	28
105	The spectrum of haemostatic characteristics of women with unexplained menorrhagia. <i>Haemophilia</i> , 2011, 17, e223-9.	1.0	41
106	Oral health in adult patients with congenital coagulation disorders - a case control study. <i>Haemophilia</i> , 2011, 17, 527-531.	1.0	20
107	von Willebrand factor/factor VIII concentrate (Humate-P) for management of elective surgery in adults and children with von Willebrand disease. <i>Haemophilia</i> , 2011, 17, 895-905.	1.0	61
108	Diagnosis and management of women with bleeding disorders – international guidelines and consensus from an international expert panel. <i>Haemophilia</i> , 2011, 17, 3-5.	1.0	15
109	Repeated infusions of VWF/FVIII concentrate: impact of VWF:FVIII ratio on FVIII trough and peak levels in a rabbit model. <i>Haemophilia</i> , 2011, 17, 808-814.	1.0	7
110	Laboratory diagnosis of von Willebrand disease: results from a prospective and blind study in 32 laboratories worldwide using lyophilized plasmas. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 220-222.	1.9	8
111	Screening for von Willebrand disease in children: a case-control study. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 1086-1089.	1.9	5
112	Validation of an automated latex particle-enhanced immunoturbidimetric von Willebrand factor activity assay. <i>Journal of Thrombosis and Haemostasis</i> , 2011, 9, 1993-2002.	1.9	42
115	von Willebrand disease: a clinical and laboratory study of sixty-five patients. <i>Annals of Hematology</i> , 2011, 90, 1183-1190.	0.8	15
116	von Willebrand Disease in Children: Diagnosis and Management of a Pediatric Cohort in One Single Center in Argentina. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 560-567.	1.5	4
117	Clinical and Laboratory Assessment of the Bleeding Pediatric Patient. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 756-762.	1.5	13
118	Diagnosis and Management of von Willebrand Disease in Spain. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 503-510.	1.5	15
119	von Willebrand Disease in the United States: A Perspective from Wisconsin. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 528-534.	1.5	26
120	Management of Inherited von Willebrand Disease in Italy: Results from the Retrospective Study on 1234 Patients. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 511-521.	1.5	48
121	Laboratory Diagnosis and Management of von Willebrand Disease in Turkey: Izmir Experience. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 581-586.	1.5	7

#	ARTICLE	IF	CITATIONS
122	Diagnosis and Management of von Willebrand Disease: A Developing Country Perspective. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 587-594.	1.5	13
123	Diagnosis and Management of von Willebrand Disease in a Single Institution of Argentina. <i>Seminars in Thrombosis and Hemostasis</i> , 2011, 37, 568-575.	1.5	8
124	Acquired type 2A von Willebrand syndrome caused by aortic valve disease corrects during valve surgery. <i>British Journal of Anaesthesia</i> , 2011, 106, 494-500.	1.5	32
125	Management of Labor and Delivery of a Patient With Von Willebrand Disease Type 2A. <i>International Anesthesiology Clinics</i> , 2011, 49, 74-80.	0.3	10
126	Congenital Bleeding Disorders in Karachi, Pakistan. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2011, 17, E131-E137.	0.7	11
127	An Evaluation of the DDAVP Infusion Test With PFA-100 and vWF Activity Assays to Distinguish vWD Types in Children. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2011, 17, 441-448.	0.7	10
128	Clinical Features and Types of Von Willebrand Disease in Karachi. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2011, 17, E102-E105.	0.7	5
129	The pharmacokinetic diversity of two von Willebrand factor (VWF)/ factor VIII (FVIII) concentrates in subjects with congenital von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2011, 106, 279-288.	1.8	30
130	Advances in the diagnosis and management of type 1 von Willebrand disease. <i>Expert Review of Hematology</i> , 2011, 4, 95-106.	1.0	8
132	Impact of loss of high-molecular-weight von Willebrand factor multimers on blood loss after aortic valve replacement. <i>British Journal of Anaesthesia</i> , 2012, 108, 754-762.	1.5	26
133	Approach to the Diagnosis and Management of Common Bleeding Disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2012, 38, 711-719.	1.5	18
134	Laboratory Investigations for Bleeding Disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2012, 38, 742-752.	1.5	41
135	Flexible Ureteroscopy in Children with von Willebrand Disease. <i>Journal of Endourology</i> , 2012, 26, 783-786.	1.1	1
136	The Influence of the ABO Blood Type on the Distribution of von Willebrand Factor in Healthy Children With no Bleeding Symptoms. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2012, 18, 316-319.	0.7	10
137	Current Management of Patients with Severe von Willebrand Disease Type 3: A 2012 Update. <i>Acta Haematologica</i> , 2012, 128, 88-99.	0.7	22
138	Newer agents in antiplatelet therapy: a review. <i>Journal of Blood Medicine</i> , 2012, 3, 33.	0.7	42
139	Unexpected Bleeding in the Operating Room. <i>Anesthesia and Analgesia</i> , 2012, 114, 73-81.	1.1	14
140	Laboratory diagnostic approach of the parentsâ€“children relationship in differentiating low-level von Willebrand factor from mild type 1 von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2012, 23, 351-353.	0.5	4



#	ARTICLE	IF	CITATIONS
142	New developments in the diagnosis and treatment of von Willebrand disease. <i>Clinical Investigation</i> , 2012, 2, 781-795.	0.0	2
143	Contribution of platelet vs. endothelial VWF to platelet adhesion and hemostasis. <i>Journal of Thrombosis and Haemostasis</i> , 2012, 10, 1646-1652.	1.9	108
145	Postpartum Hemorrhage. <i>Primary Care - Clinics in Office Practice</i> , 2012, 39, 167-187.	0.7	18
146	Biological therapies for von Willebrand disease. <i>Expert Opinion on Biological Therapy</i> , 2012, 12, 551-564.	1.4	33
147	Management of children with inherited mild bleeding disorders undergoing adenotonsillar procedures. <i>International Journal of Pediatric Otorhinolaryngology</i> , 2012, 76, 291-294.	0.4	12
148	Management of a child with von Willebrand disease (type 2A) and extensive burns: A case report. <i>Burns</i> , 2012, 38, e1-e4.	1.1	0
149	Why Is My Patient Bleeding Or Bruising?. <i>Hematology/Oncology Clinics of North America</i> , 2012, 26, 321-344.	0.9	14
150	Targeting von Willebrand factor as a novel anti-platelet therapy; Application of ARC1779, an Anti-vWF aptamer, against thrombotic risk. <i>Archives of Pharmacal Research</i> , 2012, 35, 1693-1699.	2.7	36
151	Evaluation of bleeding disorders in women with menorrhagia: a survey of obstetrician-gynecologists. <i>American Journal of Obstetrics and Gynecology</i> , 2012, 207, 269.e1-269.e5.	0.7	18
152	Coagulation management in patients undergoing mechanical circulatory support. <i>Bailliere's Best Practice and Research in Clinical Anaesthesiology</i> , 2012, 26, 179-198.	1.7	77
153	Haematological Problems in Pregnancy. , 2012, , 151-172.		1
155	Determinants of bleeding phenotype in adult patients with moderate or severe von Willebrand disease. <i>Thrombosis and Haemostasis</i> , 2012, 108, 683-692.	1.8	87
156	12 Angeborenes von-Willebrand-Syndrom. , 2012, , .		0
157	27 Einzelfaktoren und Inhibitoren der plasmatischen Gerinnung. , 2012, , .		0
158	Hemorrhagic Disorders. , 2012, , e34-e43.		4
159	Hemorrhagic Disorders. , 2012, , 1137-1145.		1
160	A novel flow cytometry single tube bead assay for quantitation of von Willebrand factor antigen and collagen-binding. <i>Thrombosis and Haemostasis</i> , 2012, 108, 999-1005.	1.8	14
161	The bleeding edge of symptom assessment. <i>Pediatric Blood and Cancer</i> , 2012, 58, 657-658.	0.8	3

#	ARTICLE	IF	CITATIONS
162	Haemostasis prophylaxis using single dose desmopressin acetate and extended use epsilon aminocaproic acid for adenotonsillectomy in patients with type 1 von Willebrand disease. Haemophilia, 2012, 18, 200-204.	1.0	8
163	Effect of fibrinolysis on bleeding phenotype in moderate and severe von Willebrand disease. Haemophilia, 2012, 18, 444-451.	1.0	10
164	Challenges in the evaluation for possible abuse: Presentations of congenital bleeding disorders in childhood. Child Abuse and Neglect, 2012, 36, 127-134.	1.3	27
165	Treatment of Hodgkin's lymphoma in a patient with type <sc>III</sc> von Willebrand's disease. Haemophilia, 2012, 18, e378-9.	1.0	0
166	Critical von Willebrand factor A1 domain residues influence type VI collagen binding. Journal of Thrombosis and Haemostasis, 2012, 10, 1417-1424.	1.9	54
167	Comparison of type I, type III and type VI collagen binding assays in diagnosis of von Willebrand disease. Journal of Thrombosis and Haemostasis, 2012, 10, 1425-1432.	1.9	48
168	Development of an ELISA method for testing VWF ristocetin cofactor activity with improved sensitivity and reliability in the diagnosis of von Willebrand disease. European Journal of Haematology, 2012, 88, 439-445.	1.1	9
169	Communication and education about triggers and environmental control strategies during pediatric asthma visits. Patient Education and Counseling, 2012, 86, 63-69.	1.0	17
170	Approaches to investigating common bleeding disorders: An evaluation of North American coagulation laboratory practices. American Journal of Hematology, 2012, 87, S45-50.	2.0	33
171	von Willebrand disease: Clinical and laboratory lessons learned from the large von Willebrand disease studies. American Journal of Hematology, 2012, 87, S4-11.	2.0	40
172	US Hemophilia Treatment Center population trends 1990â€“2010: patient diagnoses, demographics, health services utilization. Haemophilia, 2013, 19, 21-26.	1.0	40
173	Hemophilias and Other Disorders of Hemostasis. , 2013, , 1-33.		12
174	Animal Models of Hemophilia and Related Bleeding Disorders. Seminars in Hematology, 2013, 50, 175-184.	1.8	34
175	Patients with Disorders of Thrombosis and Hemostasis. Medical Clinics of North America, 2013, 97, 1161-1180.	1.1	12
176	Can von Willebrand Disease Be Investigated on Combined Hormonal Contraceptives?. Journal of Pediatric and Adolescent Gynecology, 2013, 26, 138-141.	0.3	18
177	Management of bleeding and coagulopathy following major trauma: an updated European guideline. Critical Care, 2013, 17, R76.	2.5	780
178	How I manage heavy menstrual bleeding. British Journal of Haematology, 2013, 162, 721-729.	1.2	16
179	Case studies in the management of refractory bleeding in patients with haemophilia A and inhibitors. Haemophilia, 2013, 19, e151-66.	1.0	6

#	ARTICLE	IF	CITATIONS
180	Pitfalls in special coagulation testing: three illustrative case studies. <i>International Journal of Laboratory Hematology</i> , 2013, 35, 334-338.	0.7	4
181	Bleeding disorders, menorrhagia and iron deficiency: impacts on health-related quality of life. <i>Haemophilia</i> , 2013, 19, 385-391.	1.0	45
182	Inherited Disorders of Platelet Function. <i>Pediatric Clinics of North America</i> , 2013, 60, 1475-1488.	0.9	7
183	No inhibitor development after continuous infusion of factor concentrates in subjects with bleeding disorders undergoing surgery: a prospective study. <i>Haemophilia</i> , 2013, 19, 438-444.	1.0	20
184	Inherited Abnormalities of Coagulation. <i>Pediatric Clinics of North America</i> , 2013, 60, 1419-1441.	0.9	34
185	Indexes of von Willebrand Factor as Biomarkers of Aortic Stenosis Severity (from the Biomarkers of) Tj ETQq1 1 0.784314 rgBT /Overl	0.7	45
186	Xenon ventilation computed tomography rules: new technology may open up further understanding in asthma. <i>Annals of Allergy, Asthma and Immunology</i> , 2013, 111, 81.	0.5	1
187	Treating symptomatic coronary artery disease in patients with Von Willebrand disease. <i>Hematology/Oncology and Stem Cell Therapy</i> , 2013, 6, 101-104.	0.6	11
188	The molecular characterization of von <scp>W</scp>illebrand disease: good in parts. <i>British Journal of Haematology</i> , 2013, 161, 166-176.	1.2	26
189	Von Willebrand disease: Pathogenesis and management. <i>Thrombosis Research</i> , 2013, 131, S47-S50.	0.8	13
190	Principles of care for the diagnosis and treatment of von Willebrand disease. <i>Haematologica</i> , 2013, 98, 667-674.	1.7	175
191	Approach to a Child with Bleeding in the Emergency Room. <i>Indian Journal of Pediatrics</i> , 2013, 80, 411-420.	0.3	8
192	Poquito a Poquito: How Latino Families With Children Who Have Asthma Make Changes in Their Home. <i>Journal of Pediatric Health Care</i> , 2013, 27, e1-e11.	0.6	5
193	Rituximab effectiveness in a patient with juvenile systemic lupus erythematosus complicated with acquired Von Willebrand syndrome. <i>Lupus</i> , 2013, 22, 1514-1517.	0.8	11
194	A novel use of thromboelastography in type 2B von Willebrand disease. <i>International Journal of Laboratory Hematology</i> , 2013, 35, e11-4.	0.7	5
195	Desmopressin acetate (DDAVP) for preventing and treating acute bleeds during pregnancy in women with congenital bleeding disorders. , 2013, , CD009824.		3
196	Arginine and Asthma. <i>Nestle Nutrition Institute Workshop Series</i> , 2013, 77, 1-15.	1.5	12
197	Regulation in Hemostasis and Thrombosis: Part Iâ€”In Vitro Diagnostics. <i>Seminars in Thrombosis and Hemostasis</i> , 2013, 39, 235-249.	1.5	34

#	ARTICLE	IF	CITATIONS
198	A Common VWF Exon 28 Haplotype in the Turkish Population. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2013, 19, 550-556.	0.7	4
199	Evaluation for Bleeding Disorders in Suspected Child Abuse. <i>Pediatrics</i> , 2013, 131, e1314-e1322.	1.0	82
200	Evaluating for Suspected Child Abuse: Conditions That Predispose to Bleeding. <i>Pediatrics</i> , 2013, 131, e1357-e1373.	1.0	55
201	Linkage analysis identifies a locus for plasma von Willebrand factor undetected by genome-wide association. <i>Proceedings of the National Academy of Sciences of the United States of America</i> , 2013, 110, 588-593.	3.3	85
202	Validation of a new panel of automated chemiluminescence assays for von Willebrand factor antigen and activity in the screening for von Willebrand disease. <i>International Journal of Laboratory Hematology</i> , 2013, 35, 555-565.	0.7	28
203	Quantification of perioperative changes in von Willebrand factor and factor VIII during elective orthopaedic surgery in normal individuals. <i>Haemophilia</i> , 2013, 19, 758-764.	1.0	20
204	Translational medicine advances in von Willebrand disease. <i>Journal of Thrombosis and Haemostasis</i> , 2013, 11, 75-83.	1.9	14
205	Improving blood disorder diagnosis: reflections on the challenges. <i>International Journal of Laboratory Hematology</i> , 2013, 35, 244-253.	0.7	3
206	Variability in platelet- and collagen-binding defects in type 2M von Willebrand disease. <i>Haemophilia</i> , 2013, 19, 590-594.	1.0	24
207	Similarity in joint function limitation in Type 3 von Willebrand's disease and moderate haemophilia A. <i>Haemophilia</i> , 2013, 19, 595-601.	1.0	12
208	Utility of platelet function analyzer as a screening tool for the diagnosis of Von Willebrand disease in adolescents with menorrhagia. <i>Pediatric Blood and Cancer</i> , 2013, 60, 1184-1187.	0.8	9
209	von Willebrand Disease in Children. , 2013, , 79-89.		1
210	The C-type lectin receptor CLEC4M binds, internalizes, and clears von Willebrand factor and contributes to the variation in plasma von Willebrand factor levels. <i>Blood</i> , 2013, 121, 5228-5237.	0.6	102
211	No increase in bleeding identified in type 1 VWD subjects with D1472H sequence variation. <i>Blood</i> , 2013, 121, 3742-3744.	0.6	28
212	Management of severe perioperative bleeding. <i>European Journal of Anaesthesiology</i> , 2013, 30, 270-382.	0.7	740
213	Phase II prospective open-label trial of recombinant interleukin-11 in desmopressin-unresponsive von Willebrand disease and mild or moderate haemophilia A. <i>Thrombosis and Haemostasis</i> , 2013, 109, 248-254.	1.8	25
214	Platelet interaction with von Willebrand factor is enhanced by shear-induced clustering of glycoprotein IbA. <i>Haematologica</i> , 2013, 98, 1810-1818.	1.7	24
215	Intra-Operative Hemorrhage: A Review of Literature. <i>Journal of Medical Diagnostic Methods</i> , 2013, 02, .	0.0	11

#	ARTICLE	IF	CITATIONS
216	Acquired Coagulation Disorders Caused by Inhibitors. , 2013, , 79-89.		0
217	Clinical, biological and molecular diagnosis. Hematologie, 2014, 20, 30-49.	0.0	3
218	Cost&ndash;consequence analysis of long-term prophylaxis in the treatment of von Willebrand disease in the Italian context. ClinicoEconomics and Outcomes Research, 2015, 7, 17.	0.7	6
219	Clinical features of gastrointestinal bleeding in patients with congenital bleeding disorders (hemophilia and von Willebrand disease) and three cases with congenital bleeding disorders diagnosed incidentally at colonoscopy. Japanese Journal of Thrombosis and Hemostasis, 2014, 25, 504-511.	0.1	0
220	Bortezomib in the Treatment of Acquired von Willebrand Disease Secondary to Monoclonal Gammopathy of Undetermined Significance. Kansas Journal of Medicine, 2014, 7, 167-170.	0.1	1
226	Allosteric activation of ADAMTS13 by von Willebrand factor. Proceedings of the National Academy of Sciences of the United States of America, 2014, 111, 18584-18589.	3.3	123
227	Hemophilias and Other Disorders of Hemostasis. , 2014, , .		2
228	Providing Young Women with Credible Health Information about Bleeding Disorders. American Journal of Preventive Medicine, 2014, 47, 674-680.	1.6	13
229	Perils, Problems, and Progress in Laboratory Diagnosis of von Willebrand Disease. Seminars in Thrombosis and Hemostasis, 2014, 40, 041-048.	1.5	32
230	Von Willebrand disease. Hematology, 2014, 19, 370-371.	0.7	1
231	Clinical and laboratory diagnosis of VWD. Hematology American Society of Hematology Education Program, 2014, 2014, 524-530.	0.9	23
232	Management of VWD. Hematology American Society of Hematology Education Program, 2014, 2014, 536-541.	0.9	16
233	A Systematic Approach to the Preoperative Medical Evaluation of Adults. Hospital Practice (1995), 2014, 42, 52-64.	0.5	2
234	Orthopaedic surgery in patients with von Willebrand disease. Haemophilia, 2014, 20, 133-140.	1.0	25
235	Diagnostic challenges in patients with bleeding phenotype and von Willebrand exon 28 polymorphism p.D1472H. Haemophilia, 2014, 20, e211-4.	1.0	4
236	Hyponatraemic seizure following arginine vasopressin for von Willebrand disease: pernicious, predictable and preventable. Internal Medicine Journal, 2014, 44, 521-522.	0.5	2
237	Quantitative impact of using different criteria for the laboratory diagnosis of type 1 von Willebrand disease. Journal of Thrombosis and Haemostasis, 2014, 12, 1238-1243.	1.9	22
238	Females with FVIII and FIX deficiency have reduced joint range of motion. American Journal of Hematology, 2014, 89, 831-836.	2.0	43

#	ARTICLE	IF	CITATIONS
239	Acquired von Willebrand syndrome in patients with overt hypothyroidism: a prospective cohort study. <i>Haemophilia</i> , 2014, 20, 326-332.	1.0	28
240	Identification and Basic Management of Bleeding Disorders in Adults. <i>Journal of the American Board of Family Medicine</i> , 2014, 27, 549-564.	0.8	20
241	Performance evaluation and multicentre study of a von Willebrand factor activity assay based on GPIb binding in the absence of ristocetin. <i>Blood Coagulation and Fibrinolysis</i> , 2014, 25, 860-870.	0.5	58
242	Current management of von Willebrand disease and von Willebrand syndrome. <i>Current Opinion in Anaesthesiology</i> , 2014, 27, 353-358.	0.9	13
243	Diagnosing von Willebrand Disease: A Short History of Laboratory Milestones and Innovations, Plus Current Status, Challenges, and Solutions. <i>Seminars in Thrombosis and Hemostasis</i> , 2014, 40, 551-570.	1.5	44
244	Acquired von Willebrand Syndrome in a Child Following Berlin Heart EXCOR Pediatric Ventricular Assist Device Implantation. <i>World Journal for Pediatric &amp; Congenital Heart Surgery</i> , 2014, 5, 592-598.	0.3	13
245	Diagnosing type 1 von Willebrand disease: good for patient's health or for doctor's prestige?: comment. <i>Journal of Thrombosis and Haemostasis</i> , 2014, 12, 2131-2134.	1.9	1
246	Diagnosing type 1 von Willebrand disease: good for patient's health or for doctor's prestige?: reply. <i>Journal of Thrombosis and Haemostasis</i> , 2014, 12, 2134-2136.	1.9	0
247	Desmopressin responsiveness at a capped dose of 15 µg in type 1 von Willebrand disease and mild hemophilia A. <i>Blood Coagulation and Fibrinolysis</i> , 2014, 25, 820-823.	0.5	25
248	Acquired von Willebrand syndrome in patients with Gaucher disease. <i>Blood Cells, Molecules, and Diseases</i> , 2014, 52, 205-207.	0.6	7
250	Von Willebrand Disease: Range of the Disease, and Management. <i>Current Pediatrics Reports</i> , 2014, 2, 60-70.	1.7	1
251	Towards improved diagnosis of von Willebrand disease: Comparative evaluations of several automated von Willebrand factor antigen and activity assays. <i>Thrombosis Research</i> , 2014, 134, 1292-1300.	0.8	57
252	Diagnosing type 1 von Willebrand disease: good for patient's health or for doctor's prestige?. <i>Journal of Thrombosis and Haemostasis</i> , 2014, 12, 1234-1237.	1.9	4
254	FVIII/VWF ratio is not a reliable predictor of VWD in children. <i>Pediatric Blood and Cancer</i> , 2014, 61, 936-939.	0.8	5
255	Anesthesia implications of coagulation and anticoagulation during pregnancy. <i>Seminars in Perinatology</i> , 2014, 38, 370-377.	1.1	10
256	Von Willebrand Disease. <i>Pediatrics in Review</i> , 2014, 35, 136-137.	0.2	0
257	Is gingival bleeding a symptom of patients with type 1 von Willebrand disease? A case-control study. <i>Journal of Clinical Periodontology</i> , 2014, 41, 766-771.	2.3	9
258	Technological advances in diagnostic testing for von Willebrand disease: new approaches and challenges. <i>International Journal of Laboratory Hematology</i> , 2014, 36, 334-340.	0.7	12

#	ARTICLE	IF	CITATIONS
259	Facteur von Willebrand et maladie de Willebrand : nouvelles approches. Revue Francophone Des Laboratoires, 2014, 2014, 53-63.	0.0	1
260	A two-center retrospective review of the hematologic evaluation and laboratory abnormalities in suspected victims of non-accidental injury. Child Abuse and Neglect, 2014, 38, 1794-1800.	1.3	12
261	Evaluating errors in the laboratory identification of von Willebrand disease in the real world. Thrombosis Research, 2014, 134, 393-403.	0.8	68
262	Analysis of the role of von Willebrand factor, platelet glycoprotein VI-, and Î±2Î²1-mediated collagen binding in thrombus formation. Blood, 2014, 124, 1799-1807.	0.6	26
263	The bleeding score predicts clinical outcomes and replacement therapy in adults with von Willebrand disease. Blood, 2014, 123, 4037-4044.	0.6	123
265	A 14-year-old girl with anemia. , 0, , 237-240.		0
266	Thromboembolic incidence with transiently elevated levels of coagulation factors in patients with von Willebrand disease treated with VWF:FVIII concentrate during surgery. Haemophilia, 2014, 20, e404-e406.	1.0	20
267	Haemophilia A carriers demonstrate pathological and radiological evidence of structural joint changes. Haemophilia, 2014, 20, e426-9.	1.0	28
268	rVWF: treatment finally reaches the modern age. Blood, 2015, 126, 1975-1976.	0.6	2
269	Diagnostic approach to von Willebrand disease. Blood, 2015, 125, 2029-2037.	0.6	148
270	Interference from lupus anticoagulant on von Willebrand factor measurement in splenic marginal zone lymphoma. Blood Coagulation and Fibrinolysis, 2015, 26, 454-457.	0.5	2
271	von Willebrand disease, molecular biology and diagnosis. CirugÃa Y Cirujanos (English Edition), 2015, 83, 255-264.	0.0	1
272	Interaction between VWF and FVIII in treating VWD. European Journal of Haematology, 2015, 95, 449-454.	1.1	7
273	The Platelet Function Analyser (<sc>PFA</sc>)â€100 and von Willebrand disease: a story well over 16Âyears in the making. Haemophilia, 2015, 21, 642-645.	1.0	23
274	Common skin and bleeding disorders that can potentially masquerade as child abuse. American Journal of Medical Genetics, Part C: Seminars in Medical Genetics, 2015, 169, 328-336.	0.7	18
275	Bleeding spectrum in children with moderate or severe von <sc>W</sc>illebrand disease: <sc>R</sc>elevance of pediatricâ€specific bleeding. American Journal of Hematology, 2015, 90, 1142-1148.	2.0	46
276	Preventing postpartum haemorrhageâ€when guidelines fall short. Haemophilia, 2015, 21, 502-504.	1.0	11
277	Changes in von Willebrand factor level and von Willebrand activity with age in type 1 von Willebrand disease. Haemophilia, 2015, 21, 636-641.	1.0	58

#	ARTICLE	IF	CITATIONS
278	Asthma and school functioning in children: Still more work to do. The Brown University Child and Adolescent Behavior Letter, 2015, 31, 1-6.	0.0	0
279	Commentary. Clinical Chemistry, 2015, 61, 912-912.	1.5	0
280	Platelet function analyser (<scp>PFA</scp>â€100) results and von Willebrand factor deficiency: a 16â€year â€realâ€worldâ€™ experience. Haemophilia, 2015, 21, 646-652.	1.0	37
281	Utility of a Paediatric Bleeding Questionnaire as a screening tool for von Willebrand disease in apparently healthy children. Haemophilia, 2015, 21, 806-811.	1.0	22
282	Qualitative and quantitative modifications of von Willebrand factor in patients with essential thrombocythemia and controlled platelet count. Journal of Thrombosis and Haemostasis, 2015, 13, 1226-1237.	1.9	48
283	Utility of a high VWF. Blood Coagulation and Fibrinolysis, 2015, 26, 515-521.	0.5	3
284	â€Bleeding in the jungleâ€. American Journal of Hematology, 2015, 90, 843-846.	2.0	3
285	An update on the management of bleeding disorders during pregnancy. Current Opinion in Hematology, 2015, 22, 397-405.	1.2	13
286	Hemostasis. , 2015, , .		0
287	Gene therapy for hemophilia. Frontiers in Bioscience - Landmark, 2015, 20, 556-603.	3.0	51
289	Evaluation and Management of Adolescents with Abnormal Uterine Bleeding. Pediatric Annals, 2015, 44, e218-22.	0.3	27
290	Contribution of the collagen binding activity (VWF:CB) in the range of tests for the diagnosis and classification of von Willebrand disease. Annales De Biologie Clinique, 2015, 73, 461-468.	0.2	0
291	Successful Aortic Aneurysm Repair in a Woman with Severe von Willebrand (Type 3) Disease. Case Reports in Hematology, 2015, 2015, 1-8.	0.3	1
294	A Case Report of Aortic Valve Replacement for a Patient with von Willebrand Disease. Japanese Journal of Cardiovascular Surgery, 2015, 44, 292-295.	0.0	1
295	Peripartum management of patient with a rare combination of two bleeding diatheses: Recognizing active role of anesthesiologists during preparatory preemptive prepartum multi-disciplinary conference. Journal of Obstetric Anaesthesia and Critical Care, 2015, 5, 30.	0.0	0
296	Disorders of Hemostasis. , 2015, , 127-134.		0
297	Desmopressin acetate (DDAVP) for preventing and treating acute bleeds during pregnancy in women with congenital bleeding disorders. The Cochrane Library, 2015, , CD009824.	1.5	9
298	Plateletâ€dependent von Willebrand factor activity. Nomenclature and methodology: communication from the SSC of the ISTH. Journal of Thrombosis and Haemostasis, 2015, 13, 1345-1350.	1.9	119



#	ARTICLE	IF	CITATIONS
299	Treatment of the acquired von Willebrand syndrome. <i>Expert Review of Hematology</i> , 2015, 8, 799-818.	1.0	19
300	Recent advances in laboratory-aided diagnosis of von Willebrand disease. <i>Expert Opinion on Orphan Drugs</i> , 2015, 3, 975-995.	0.5	11
301	Disorders of coagulation in pregnancy. <i>British Journal of Anaesthesia</i> , 2015, 115, ii75-ii88.	1.5	219
302	Clinical use of Haemate <sup>®</sup> P in von Willebrand disease: A 25-year retrospective observational study. <i>Thrombosis Research</i> , 2015, 135, 479-484.	0.8	19
303	Microvascular abnormalities in patients with early systemic sclerosis: less severe morphological changes than in patients with definite disease. <i>Scandinavian Journal of Rheumatology</i> , 2015, 44, 48-55.	0.6	13
304	Continuous-flow Left Ventricular Assist Device, Valvular Disease, and Acquired von Willebrand Syndrome. <i>Clinical Gastroenterology and Hepatology</i> , 2015, 13, 1376-1377.	2.4	0
305	Reply. <i>Clinical Gastroenterology and Hepatology</i> , 2015, 13, 1375-1376.	2.4	0
306	Practice Guideline: Epistaxis in Children. <i>Journal of Pediatric Health Care</i> , 2015, 29, 484-488.	0.6	12
308	Current controversies in the diagnosis and management of von Willebrand disease. <i>Therapeutic Advances in Hematology</i> , 2015, 6, 209-216.	1.1	16
309	Von Willebrand Disease: diagnosis and management. <i>Paediatrics and Child Health (United Kingdom)</i> , 2015, 25, 354-359.	0.2	3
311	Haemophilia A carriers experience reduced health-related quality of life. <i>Haemophilia</i> , 2015, 21, 761-765.	1.0	29
312	A cross-sectional study of bleeding phenotype in haemophilia A carriers. <i>British Journal of Haematology</i> , 2015, 170, 223-228.	1.2	75
313	Complex haemostatic abnormalities as a cause of bleeding after neurosurgery in a patient with Gaucher disease. <i>Platelets</i> , 2015, 26, 260-262.	1.1	2
314	Platelet Disorders. , 2015, , 572-575.		0
315	Caring for the Patient With von Willebrand Disease Who Requires Ambulatory Surgery. <i>Journal of Perianesthesia Nursing</i> , 2015, 30, 423-426.	0.3	0
316	Elderly Female with a Personal and Family History of a Bleeding Disorder. <i>Clinical Chemistry</i> , 2015, 61, 909-912.	1.5	1
317	Laboratory diagnosis of von Willebrand disease. <i>International Journal of Laboratory Hematology</i> , 2015, 37, 11-17.	0.7	35
318	New Treatments for Left Ventricular Assist Device-Associated Bleeding? —. <i>JACC: Heart Failure</i> , 2015, 3, 870-872.	1.9	0

#	ARTICLE	IF	CITATIONS
319	Predictors of von Willebrand disease diagnosis in individuals with borderline von Willebrand factor plasma levels. <i>Journal of Thrombosis and Haemostasis</i> , 2015, 13, 228-236.	1.9	18
320	Combined oral contraceptives do not influence von Willebrand factor related biomarkers despite an induced acute phase response. <i>Thrombosis Research</i> , 2015, 135, 208-211.	0.8	5
322	Management of the Medically Compromised Patient. , 2016, , 540-562.		2
323	Prolonged Bleeding in a 34-Year-Old Man following Oral Surgery. <i>Clinical Chemistry</i> , 2016, 62, 1676-1677.	1.5	0
324	Recent progress in von Willebrand disease type 2B. <i>Hematologie</i> , 2016, 22, 259-270.	0.0	0
325	Human plasma-derived FVIII/VWD concentrate (Biostat): a review of experimental and clinical pharmacokinetic, efficacy and safety data. <i>Drugs in Context</i> , 2016, 5, 1-10.	1.0	6
326	Disorders of Coagulation. , 2016, , 279-333.		2
327	Unnecessary Hysterectomy due to Menorrhagia and Disorders of Hemostasis: An Example of Overuse and Excessive Demand for Medical Services. <i>Frontiers in Pharmacology</i> , 2016, 7, 507.	1.6	2
328	Biological diagnosis of von Willebrand disease: analytical characteristics of Innovance vWF:Ac assay kit on STA-R Evolution Expert series analyzer (Stago). <i>Annales De Biologie Clinique</i> , 2016, 74, 355-364.	0.2	0
329	Practical aspects of factor concentrate use in patients with von Willebrand disease undergoing invasive procedures: a European survey. <i>Haemophilia</i> , 2016, 22, 739-751.	1.0	16
330	Surgery in patients with von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2016, 27, 812-816.	0.5	21
331	A diagnostic approach to mild bleeding disorders. <i>Journal of Thrombosis and Haemostasis</i> , 2016, 14, 1507-1516.	1.9	63
332	Developing a multidisciplinary Young Women's Blood Disorders Program: a single-centre approach with guidance for other centres. <i>Haemophilia</i> , 2016, 22, 199-207.	1.0	38
333	Practical aspects of <sc>DDAVP</sc> use in patients with von Willebrand Disease undergoing invasive procedures: a European survey. <i>Haemophilia</i> , 2016, 22, 110-120.	1.0	24
334	Von Willebrand factor for menorrhagia: a survey and literature review. <i>Haemophilia</i> , 2016, 22, 397-402.	1.0	37
335	Baseline factor <sc>VIII</sc> plasma levels and age at first bleeding in patients with severe forms of von Willebrand disease. <i>Haemophilia</i> , 2016, 22, 564-569.	1.0	4
336	Pediatric Coagulation Disorders. <i>Pediatrics in Review</i> , 2016, 37, 279-291.	0.2	13
337	Congenital and acquired bleeding disorders in pregnancy. <i>Hematology American Society of Hematology Education Program</i> , 2016, 2016, 232-235.	0.9	10

#	ARTICLE	IF	CITATIONS
338	What have we learned from large population studies of von Willebrand disease?. Hematology American Society of Hematology Education Program, 2016, 2016, 670-677.	0.9	10
339	Diagnosing von Willebrand disease: genetic analysis. Hematology American Society of Hematology Education Program, 2016, 2016, 678-682.	0.9	31
340	New treatment approaches to von Willebrand disease. Hematology American Society of Hematology Education Program, 2016, 2016, 683-689.	0.9	27
341	Challenges of diagnosing and managing the adolescent with heavy menstrual bleeding. Thrombosis Research, 2016, 143, 91-100.	0.8	30
342	Common gynecological challenges in adolescents with sickle cell disease. Expert Review of Hematology, 2016, 9, 187-196.	1.0	13
343	The European guideline on management of major bleeding and coagulopathy following trauma: fourth edition. Critical Care, 2016, 20, 100.	2.5	1,014
344	Laboratory tests used to help diagnose von Willebrand disease: an update. Pathology, 2016, 48, 303-318.	0.3	81
345	Why Do Patients Bleed?. The Surgery Journal, 2016, 02, e29-e43.	0.3	16
346	Update on inherited disorders of haemostasis and pregnancy. Obstetric Medicine, 2016, 9, 64-72.	0.5	9
347	Type 2M and Type 2A von Willebrand Disease: Similar but Different. Seminars in Thrombosis and Hemostasis, 2016, 42, 483-497.	1.5	35
348	Rapid discrimination of the phenotypic variants of von Willebrand disease. Blood, 2016, 127, 2472-2480.	0.6	19
349	How I treat patients with inherited bleeding disorders who need anticoagulant therapy. Blood, 2016, 128, 178-184.	0.6	80
350	Defining von Willebrand disease. Blood, 2016, 127, 2373-2374.	0.6	3
351	Pediatric Acquired von Willebrand Disease With Berlin Heart Excor Ventricular Assist Device Support. World Journal for Pediatric & Congenital Heart Surgery, 2016, 7, 614-618.	0.3	9
352	Coagulation factor concentrates for inherited bleeding disorders. , 2016, , 328-343.		0
353	Von Willebrand Disease and Pregnancy: A Review of Evidence and Expert Opinion. Seminars in Thrombosis and Hemostasis, 2016, 42, 717-723.	1.5	34
354	Treatment Options for the Adolescent Patient Experiencing Abnormal Uterine Bleeding. Current Treatment Options in Pediatrics, 2016, 2, 184-195.	0.2	1
355	State of the art: von Willebrand disease. Haemophilia, 2016, 22, 54-59.	1.0	31

#	ARTICLE	IF	CITATIONS
356	Current and emerging approaches for assessing von Willebrand disease in 2016. <i>International Journal of Laboratory Hematology</i> , 2016, 38, 41-49.	0.7	14
357	A single-centre study of haemostatic outcomes of joint replacement in von Willebrand disease and control patients and an analysis of the literature. <i>Haemophilia</i> , 2016, 22, 934-942.	1.0	6
358	Von Willebrand's Disease. <i>New England Journal of Medicine</i> , 2016, 375, 2067-2080.	13.9	389
359	von Willebrand Disease: Prevention of Complications and Management of the Disease. , 2016, , 295-311.		0
360	Clinical and laboratory variability in a cohort of patients diagnosed with type 1 VWD in the United States. <i>Blood</i> , 2016, 127, 2481-2488.	0.6	96
361	Management of Bleeding Patients. , 2016, , .		3
362	Present day management of inherited bleeding disorders in pregnancy. <i>Expert Review of Hematology</i> , 2016, 9, 987-995.	1.0	5
363	A medical elaboration on von Willebrand disease with its dental management. <i>Journal of Oral and Maxillofacial Surgery, Medicine, and Pathology</i> , 2016, 28, 442-445.	0.2	2
364	Hemostatic Agents Used to Stop Bleeding. , 2016, , 321-336.		1
366	A prospective study of von Willebrand factor levels and bleeding in pregnant women with type 1 von Willebrand disease. <i>Haemophilia</i> , 2016, 22, e562-e564.	1.0	18
367	Heavy Menstrual Bleeding. , 2016, , 199-206.		1
369	Joint surgery in von Willebrand disease: a multicentre cross-sectional study. <i>Haemophilia</i> , 2016, 22, 256-262.	1.0	6
370	Von Willebrand disease in the emergency department. <i>Haemophilia</i> , 2016, 22, 263-267.	1.0	4
371	Type 2M von Willebrand disease " more often misidentified than correctly identified. <i>Haemophilia</i> , 2016, 22, e145-55.	1.0	38
372	Prophylaxis in von Willebrand Disease: Coming of Age?. <i>Seminars in Thrombosis and Hemostasis</i> , 2016, 42, 498-506.	1.5	21
373	Review of patients studied for coagulopathy in a haematology/oncology unit. <i>Anales De Pediatr�a (English Edition)</i> , 2016, 84, 85-91.	0.1	0
374	Evaluation of a von Willebrand factor three test panel and chemiluminescent-based assay system for identification of, and therapy monitoring in, von Willebrand disease. <i>Thrombosis Research</i> , 2016, 141, 202-211.	0.8	57
375	Use of a Pediatric Bleeding Questionnaire in the Screening of Von Willebrand Disease in Young Females at Menarche in the Primary Care Setting. <i>Journal of Pediatric Health Care</i> , 2016, 30, 408-413.	0.6	6

#	ARTICLE	IF	CITATIONS
376	Towards a more automatic and rapid laboratory diagnosis of von Willebrand disease. <i>Thrombosis Research</i> , 2016, 141, 198-201.	0.8	7
377	Treatment of von Willebrand Disease. <i>Seminars in Thrombosis and Hemostasis</i> , 2016, 42, 133-146.	1.5	59
379	Genome-wide association studies identify genetic loci for low von Willebrand factor levels. <i>European Journal of Human Genetics</i> , 2016, 24, 1035-1040.	1.4	45
380	Validation of automated latex von Willebrand factor activity assay in a new prospective study including post-treatment patient samples. <i>International Journal of Laboratory Hematology</i> , 2017, 39, 286-292.	0.7	1
381	Laboratory monitoring of replacement therapy for major surgery in von Willebrand disease. <i>Haemophilia</i> , 2017, 23, 182-187.	1.0	18
382	Molecular diagnosis of von Willebrand disease. <i>Haemophilia</i> , 2017, 23, 188-197.	1.0	32
383	von Willebrand Disease: A Concise Review and Update for the Practicing Physician. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 900-910.	0.7	36
384	Utilization and Effectiveness of Desmopressin Acetate After Cardiac Surgery Supplemented With Point-of-Care Hemostatic Testing: A Propensity-Score Matched Analysis. <i>Journal of Cardiothoracic and Vascular Anesthesia</i> , 2017, 31, 883-895.	0.6	5
385	Screening of female family members of von Willebrand disease patients: utility of a modified screening tool in a high-risk population. <i>Haemophilia</i> , 2017, 23, 736-742.	1.0	3
386	Pediatric Acquired von Willebrand Syndrome in Cardiopulmonary Disorders: Do Laboratory Abnormalities Predict Bleeding Risk?. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, 121-125.	0.3	8
387	Management of severe perioperative bleeding. <i>European Journal of Anaesthesiology</i> , 2017, 34, 332-395.	0.7	650
388	Management of Inherited Bleeding Disorders in Pregnancy. <i>BJOG: an International Journal of Obstetrics and Gynaecology</i> , 2017, 124, e193-e263.	1.1	74
389	Controversies in the diagnosis of Type 1 von Willebrand disease. <i>International Journal of Laboratory Hematology</i> , 2017, 39, 61-68.	0.7	16
390	A Microfluidic Model of Hemostasis Sensitive to Platelet Function and Coagulation. <i>Cellular and Molecular Bioengineering</i> , 2017, 10, 3-15.	1.0	45
391	von Willebrand factor disruption and continuous-flow circulatory devices. <i>Journal of Heart and Lung Transplantation</i> , 2017, 36, 1155-1163.	0.3	39
392	Feasibility of the Von Willebrand disease PREVENT trial. <i>Thrombosis Research</i> , 2017, 156, 8-13.	0.8	9
393	Enhanced Local Disorder in a Clinically Elusive von Willebrand Factor Provokes High-Affinity Platelet Clumping. <i>Journal of Molecular Biology</i> , 2017, 429, 2161-2177.	2.0	36
394	Risk factors and prognosis of ovarian vein thrombosis. <i>Blood Coagulation and Fibrinolysis</i> , 2017, 28, 468-474.	0.5	15

#	ARTICLE	IF	CITATIONS
395	Type 2B von Willebrand Disease: An Unusual Cause of Severe Neonatal Thrombocytopenia. <i>Journal of Pediatric Hematology/Oncology</i> , 2017, 39, 473-475.	0.3	1
396	Clinical and laboratory phenotype variability in type 2M von Willebrand disease. <i>Journal of Thrombosis and Haemostasis</i> , 2017, 15, 1559-1566.	1.9	13
397	Pregnancy loss in women with von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2017, 28, 393-397.	0.5	16
398	von Willebrand disease Outreach into Integrated Care Education (VOICE): a call to action. <i>Haemophilia</i> , 2017, 23, e370-e373.	1.0	3
399	The use of antifibrinolytics in pediatric patients with hypoproliferative thrombocytopenia. <i>Pediatric Blood and Cancer</i> , 2017, 64, e26641.	0.8	1
400	Efficacy and safety of a <sup>VWF</sup>/<sup>FVIII</sup> concentrate (wilate<sup>®</sup>) in inherited von Willebrand disease patients undergoing surgical procedures. <i>Haemophilia</i> , 2017, 23, 264-272.	1.0	29
401	Laboratory Testing for von Willebrand Disease: The Past, Present, and Future State of Play for von Willebrand Factor Assays that Measure Platelet Binding Activity, with or without Ristocetin. <i>Seminars in Thrombosis and Hemostasis</i> , 2017, 43, 075-091.	1.5	22
402	Managing and Supporting Surgery in Patients with Bleeding Disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2017, 43, 653-671.	1.5	5
403	Novel insights into the clinical phenotype and pathophysiology underlying low VWF levels. <i>Blood</i> , 2017, 130, 2344-2353.	0.6	98
404	Recombinant von Willebrand factor for severe gastrointestinal bleeding unresponsive to other treatments in a patient with type 2A von Willebrand disease. <i>Blood Coagulation and Fibrinolysis</i> , 2017, 28, 570-575.	0.5	8
405	Haemate P <sup>®</sup> zastosowanie w profilaktyce i leczeniu krwawieÅ, w chorobie von Willebranda oraz indukcji immunotolerancji w hemofilii A powikÅ,anej inhibitorem. <i>Acta Haematologica Polonica</i> , 2017, 48, 125-129.	0.1	0
406	Impact of diagnosis of von Willebrand disease on patient outcomes: Analysis of medical insurance claims data. <i>Haemophilia</i> , 2017, 23, 743-749.	1.0	22
407	Current and Emerging Options for the Management of Inherited von Willebrand Disease. <i>Drugs</i> , 2017, 77, 1531-1547.	4.9	28
408	Laboratory Testing for von Willebrand Factor Antigen (VWF:Ag). <i>Methods in Molecular Biology</i> , 2017, 1646, 403-416.	0.4	15
409	Laboratory Testing for von Willebrand Factor Collagen Binding (VWF:CB). <i>Methods in Molecular Biology</i> , 2017, 1646, 417-433.	0.4	19
410	Laboratory Testing for von Willebrand Factor Ristocetin Cofactor (VWF:RCo). <i>Methods in Molecular Biology</i> , 2017, 1646, 435-451.	0.4	19
411	Laboratory Testing for von Willebrand Factor: Factor VIII Binding (for 2N VWD). <i>Methods in Molecular Biology</i> , 2017, 1646, 461-472.	0.4	14
412	Low VWF: an established mild bleeding disorder?. <i>Blood</i> , 2017, 130, 2241-2242.	0.6	5

#	ARTICLE	IF	CITATIONS
413	Advances in the diagnosis and treatment of Von Willebrand disease. <i>Blood</i> , 2017, 130, 2386-2391.	0.6	64
414	Can you grow out of von Willebrand disease?. <i>Haemophilia</i> , 2017, 23, 807-809.	1.0	7
415	Monitoring Therapy during Treatment of von Willebrand Disease. <i>Seminars in Thrombosis and Hemostasis</i> , 2017, 43, 338-354.	1.5	25
416	von Willebrand disease type1/type 2N compound heterozygotes: diagnostic and management challenges. <i>British Journal of Haematology</i> , 2017, 176, 994-997.	1.2	4
417	Outcomes in Patients With Hemophilia and von Willebrand Disease Undergoing Invasive or Surgical Procedures. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2017, 23, 148-154.	0.7	17
418	Von Willebrand disease – the D <sup>+</sup> and D <sup>-</sup> on'ts™ in surgery. <i>European Journal of Haematology</i> , 2017, 98, 121-127.	1.1	30
419	The Impact of Antithrombin Deficiency on Women's Reproductive Health Experiences and Healthcare Decision-Making. <i>Journal of Women's Health</i> , 2017, 26, 1350-1355.	1.5	0
420	Blood volume–based von Willebrand factor to prevent postpartum hemorrhage in von Willebrand disease. <i>Blood Advances</i> , 2017, 1, 703-706.	2.5	11
421	Diagnosis and Treatment of von Willebrand Disease and Rare Bleeding Disorders. <i>Journal of Clinical Medicine</i> , 2017, 6, 45.	1.0	59
422	Semantic prioritization of novel causative genomic variants. <i>PLoS Computational Biology</i> , 2017, 13, e1005500.	1.5	28
423	Investigation of Haemostasis. , 2017, , 366-409.		17
424	Advances in the diagnosis and treatment of Von Willebrand disease. <i>Hematology American Society of Hematology Education Program</i> , 2017, 2017, 379-384.	0.9	17
425	3rd GUIDELINE FOR PERIOPERATIVE CARDIOVASCULAR EVALUATION OF THE BRAZILIAN SOCIETY OF CARDIOLOGY. <i>Arquivos Brasileiros De Cardiologia</i> , 2017, 109, 1-104.	0.3	21
426	A case report on the successful perioperative management of hepatectomy for hepatocellular carcinoma in a patient with von Willebrand disease. <i>International Journal of Surgery Case Reports</i> , 2018, 44, 131-134.	0.2	1
427	Current issues in diagnosis and treatment of von Willebrand disease. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2018, 2, 34-41.	1.0	19
429	The course of acquired von Willebrand syndrome during pregnancy among patients with essential thrombocytosis. <i>Journal of Thrombosis and Thrombolysis</i> , 2018, 46, 304-309.	1.0	7
430	Desmopressin stimulation testing: Response to intravenous and intranasal forms. <i>Haemophilia</i> , 2018, 24, e194-e198.	1.0	2
431	Analysis of current perioperative management with Haemate <sup>®</sup> P/Humate P <sup>®</sup> in von Willebrand disease: Identifying the need for personalized treatment. <i>Haemophilia</i> , 2018, 24, 460-470.	1.0	28

#	ARTICLE	IF	CITATIONS
432	Advantage of recombinant von Willebrand factor for perioperative management in paediatric acquired von Willebrand syndrome. <i>Haemophilia</i> , 2018, 24, e120-e121.	1.0	5
433	How I treat type 2B von Willebrand disease. <i>Blood</i> , 2018, 131, 1292-1300.	0.6	40
434	Comprehensive reevaluation of historical von Willebrand disease diagnosis in association with whole blood platelet aggregation and function. <i>International Journal of Laboratory Hematology</i> , 2018, 40, 304-311.	0.7	8
435	Preanalytical issues that may cause misdiagnosis in haemophilia and von Willebrand disease. <i>Haemophilia</i> , 2018, 24, 198-210.	1.0	20
436	Von Willebrand factor multimer quantitation for assessment of cardiac lesion severity and bleeding risk. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2018, 2, 155-161.	1.0	3
437	Differential sensitivity of von Willebrand factor activity assays to reduced VWF molecular weight forms: A large international cross-laboratory study. <i>Thrombosis Research</i> , 2018, 166, 96-105.	0.8	23
438	Relevance of Abusive Head Trauma to Intracranial Hemorrhages and Bleeding Disorders. <i>Pediatrics</i> , 2018, 141, e20173485.	1.0	15
439	The primary haemostasis is more preserved in thrombocytopenic patients with liver cirrhosis than cancer. <i>Blood Coagulation and Fibrinolysis</i> , 2018, 29, 307-313.	0.5	4
441	von Willebrand Disease. , 2018, , 348-362.		0
442	Evaluation of a microfluidic flow assay to screen for von Willebrand disease and low von Willebrand factor levels. <i>Journal of Thrombosis and Haemostasis</i> , 2018, 16, 104-115.	1.9	16
443	Abnormal Uterine Bleeding including coagulopathies and other menstrual disorders. <i>Best Practice and Research in Clinical Obstetrics and Gynaecology</i> , 2018, 48, 51-61.	1.4	13
444	Outgrowing the laboratory diagnosis of type 1 von Willebrand disease: A two decade study. <i>American Journal of Hematology</i> , 2018, 93, 232-237.	2.0	17
445	Bleeding score in Type 1 von Willebrand disease patients using the ISTH-BAT questionnaire. <i>International Journal of Laboratory Hematology</i> , 2018, 40, 175-180.	0.7	19
447	Von Willebrand disease in the United States: perspective from the Zimmerman program. <i>Annals of Blood</i> , 2018, 3, 7-7.	0.4	14
448	Diagnosis and management of von Willebrand disease in Australia. <i>Annals of Blood</i> , 0, 3, 31-31.	0.4	3
449	Diagnosis and management of von Willebrand disease in Italy. <i>Annals of Blood</i> , 0, 3, 28-28.	0.4	1
450	Endothelial Cell von Willebrand Factor Secretion in Health and Cardiovascular Disease. , 2018, ,		8
451	Maladie de Willebrand et prévention du risque hémorragique en chirurgie. <i>Praticien En Anesthésie Réanimation</i> , 2018, 22, 326-334.	0.0	0



#	ARTICLE	IF	CITATIONS
452	Von Willebrand Disease. , 0, , 83-99.		1
453	Evaluation and management of heavy menstrual bleeding in adolescents: the role of the hematologist. Blood, 2018, 132, 2134-2142.	0.6	17
454	Utility of a screening tool for haemostatic defects in a multicentre cohort of adolescents with heavy menstrual bleeding. Haemophilia, 2018, 24, 957-963.	1.0	13
456	Long-Term Outcome after Joint Bleeds in Von Willebrand Disease Compared to Haemophilia A: A Post Hoc Analysis. Thrombosis and Haemostasis, 2018, 118, 1690-1700.	1.8	10
457	Genetic regulation of plasma von Willebrand factor levels in health and disease. Journal of Thrombosis and Haemostasis, 2018, 16, 2375-2390.	1.9	48
458	Pathophysiology of Coagulation and Deficiencies of Coagulation Factors in Newborn. , 2018, , 1431-1453.		1
459	Clinically relevant differences between assays for von Willebrand factor activity. Journal of Thrombosis and Haemostasis, 2018, 16, 2413-2424.	1.9	26
460	Rare forms of von Willebrand disease. Annals of Translational Medicine, 2018, 6, 345-345.	0.7	12
461	Von Willebrand Disease in the elderly: clinical perspectives. Clinical Interventions in Aging, 2018, Volume 13, 1531-1541.	1.3	13
462	Pharmacokinetics-based clinical management of acquired von Willebrand syndrome: a case report. Journal of Blood Medicine, 2018, Volume 9, 9-13.	0.7	2
463	Preoperative screening for bleeding disorders: A comprehensive laboratory assessment of clinical practice. Research and Practice in Thrombosis and Haemostasis, 2018, 2, 767-777.	1.0	30
464	Evaluation and management of heavy menstrual bleeding in adolescents: the role of the hematologist. Hematology American Society of Hematology Education Program, 2018, 2018, 390-398.	0.9	24
465	Significant gynecological bleeding in women with low von Willebrand factor levels. Blood Advances, 2018, 2, 1784-1791.	2.5	79
466	von Willebrand Disease. Pediatric Clinics of North America, 2018, 65, 527-541.	0.9	11
467	Abnormal Uterine Bleeding in Young Women with Blood Disorders. Pediatric Clinics of North America, 2018, 65, 543-560.	0.9	12
468	Von Willebrand Disease: Classification and Diagnosis. , 2018, , 149-173.		0
469	Clinical Approach to the Patient With Bleeding or Bruising. , 2018, , 1912-1921.		2
470	Laboratory Evaluation of Hemostatic and Thrombotic Disorders. , 2018, , 1922-1931.		0

#	ARTICLE	IF	CITATIONS
471	Structure, Biology, and Genetics of von Willebrand Factor. , 2018, , 2051-2063.		3
472	Current approaches to diagnostic testing in von Willebrand Disease. Transfusion and Apheresis Science, 2018, 57, 463-465.	0.5	9
473	Coagulopathies and Hypercoagulable States. , 2018, , 361-372.		0
474	How I investigate for bleeding disorders. International Journal of Laboratory Hematology, 2018, 40, 6-14.	0.7	35
475	Chemical- and Drug-Induced Asthma. , 2018, , 623-637.		0
476	Molecular Basis of Hemostatic and Thrombotic Diseases. , 2018, , 277-297.		1
477	Von Willebrand disease: Clinical conundrums. Haemophilia, 2018, 24, 37-43.	1.0	9
478	Evaluation of a new commercial method for von Willebrand factor multimeric analysis. International Journal of Laboratory Hematology, 2018, 40, 586-591.	0.7	13
479	Essentials of Hematology. , 2019, , 217-239.e8.		1
480	A diagnosis of von Willebrand disease despite normal test results for factor VIII and von Willebrand factor antigen and activity. American Journal of Hematology, 2019, 94, 1425-1432.	2.0	3
481	Women and inherited bleeding disorders – A review with a focus on key challenges for 2019. Transfusion and Apheresis Science, 2019, 58, 613-622.	0.5	13
482	Bleeding Symptoms and von Willebrand Factor Levels: 30-Year Experience in a Tertiary Care Center. Clinical and Applied Thrombosis/Hemostasis, 2019, 25, 107602961986691.	0.7	2
483	Utility of repeat testing in the evaluation for von Willebrand disease in pediatric patients. Journal of Thrombosis and Haemostasis, 2019, 17, 1838-1847.	1.9	19
484	von Willebrand Disease in Pediatrics. Hematology/Oncology Clinics of North America, 2019, 33, 425-438.	0.9	6
485	Review of von Willebrand Disease and Acquired von Willebrand Syndrome for Patients Undergoing Cardiac Surgery. Journal of Cardiothoracic and Vascular Anesthesia, 2019, 33, 3446-3457.	0.6	5
487	How I manage severe von Willebrand disease. British Journal of Haematology, 2019, 187, 418-430.	1.2	24
488	A personalized approach to the management of VWD. Transfusion and Apheresis Science, 2019, 58, 590-595.	0.5	6
489	Updated overview on von Willebrand disease: focus on the interest of genotyping. Expert Review of Hematology, 2019, 12, 1023-1036.	1.0	6

#	ARTICLE	IF	CITATIONS
490	Spinal Anesthesia in 2 Consecutive Cesarean Deliveries in a Parturient With Type 3 von Willebrand Disease: A Case Report. <i>A&amp;A Practice</i> , 2019, 12, 79-81.	0.2	2
491	Von Willebrand disease: diagnosis and management. <i>Paediatrics and Child Health (United Kingdom)</i> , 2019, 29, 339-344.	0.2	0
492	Analgesia, anaesthesia and obstetric outcome in women with inherited bleeding disorders. <i>European Journal of Obstetrics, Gynecology and Reproductive Biology</i> , 2019, 239, 60-63.	0.5	6
493	VWFâ€VIII concentrates in the treatment of inherited von Willebrand disease: A singleâ€centre retrospective study. <i>Haemophilia</i> , 2019, 25, e300-e303.	1.0	2
494	Evolution of replacement therapy for von Willebrand disease: From plasma fraction to recombinant von Willebrand factor. <i>Blood Reviews</i> , 2019, 38, 100572.	2.8	29
495	Periprocedural management of von Willebrand disease: An institutional experience. <i>Haemophilia</i> , 2019, 25, e199-e203.	1.0	3
496	Acquired Von Willebrand Syndrome (AVWS) in cardiovascular disease: a state of the art review for clinicians. <i>Journal of Thrombosis and Thrombolysis</i> , 2019, 48, 14-26.	1.0	32
497	Update on Molecular Testing in von Willebrand Disease. <i>Seminars in Thrombosis and Hemostasis</i> , 2019, 45, 708-719.	1.5	9
498	Advances in diagnosis of von Willebrand disease. <i>Expert Opinion on Orphan Drugs</i> , 2019, 7, 147-155.	0.5	1
499	Recent Advances in Mainstream Hemostasis Diagnostics and Coagulation Testing. <i>Seminars in Thrombosis and Hemostasis</i> , 2019, 45, 228-246.	1.5	17
500	Recurrent Gastrointestinal Bleeding from Dieulafoyâ€™s Lesions in a Patient with Type 1 von Willebrand Disease: A Rare Association. <i>GE Portuguese Journal of Gastroenterology</i> , 2019, 26, 202-206.	0.3	5
501	Updated Australian consensus statement on management of inherited bleeding disorders in pregnancy. <i>Medical Journal of Australia</i> , 2019, 210, 326-332.	0.8	20
502	Acquired von Willebrand Syndrome Associated with Cardiovascular Diseases. <i>Journal of Atherosclerosis and Thrombosis</i> , 2019, 26, 303-314.	0.9	69
503	Factor products. <i>JACCP Journal of the American College of Clinical Pharmacy</i> , 2019, 2, 82-86.	0.5	0
504	Increased galactose expression and enhanced clearance in patients with low von Willebrand factor. <i>Blood</i> , 2019, 133, 1585-1596.	0.6	32
505	Low endoscopy bleeding risk in patients with congenital bleeding disorders. <i>Haemophilia</i> , 2019, 25, 289-295.	1.0	6
506	Perioperative management of patients with von Willebrand disease. <i>Hematology American Society of Hematology Education Program</i> , 2019, 2019, 604-609.	0.9	14
507	New advances in the diagnosis of von Willebrand disease. <i>Hematology American Society of Hematology Education Program</i> , 2019, 2019, 596-600.	0.9	21

#	ARTICLE	IF	CITATIONS
508	New therapies for von Willebrand disease. Hematology American Society of Hematology Education Program, 2019, 2019, 590-595.	0.9	7
509	Clinical Issues in Women with Inherited Bleeding Disorders. , 2019, , .		0
510	Replacement Therapy in Patients with Von Willebrand Diseaseâ€™ Indications and Monitoring. Hamostaseologie, 2019, 39, 326-338.	0.9	2
511	New therapies for von Willebrand disease. Blood Advances, 2019, 3, 3481-3487.	2.5	29
512	The role of genetics in the pathogenesis and diagnosis of type 1 Von Willebrand disease. Current Opinion in Hematology, 2019, 26, 331-335.	1.2	10
513	Hemostatic abnormalities in adult patients with Marfan syndrome. Cardiovascular Diagnosis and Therapy, 2019, 9, S209-S220.	0.7	9
514	Case report of two siblings with type 2A von Willebrand disease involving a novel mutation within the calcium-binding site of the A2 domain of von Willebrand factor. Blood Coagulation and Fibrinolysis, 2019, 30, 161-167.	0.5	0
515	Self-reported reproductive health experiences in women with von Willebrand disease: a qualitative interview-based study. Journal of Obstetrics and Gynaecology, 2019, 39, 288-290.	0.4	6
516	How I treat low von Willebrand factor levels. Blood, 2019, 133, 795-804.	0.6	36
517	Defective collagen binding and increased bleeding in a murine model of von Willebrand disease affecting collagenâ€™ binding. Journal of Thrombosis and Haemostasis, 2019, 17, 63-71.	1.9	7
518	Diagnostic challenges of inherited mild bleeding disorders: a bait for poorly explored clinical and basic research. Journal of Thrombosis and Haemostasis, 2019, 17, 257-270.	1.9	38
519	Management of von Willebrand disease in patients undergoing total hip and knee arthroplasty. Journal of Perioperative Practice, 2019, 29, 266-269.	0.3	3
520	Laboratory Diagnosis of Inherited von Willebrand Disease. , 2019, , 799-805.		1
521	Laboratory Diagnosis of Acquired von Willebrandâ€™s Syndrome. , 2019, , 807-808.		0
522	Laboratory Assessment of Treatment of von Willebrand Disease. , 2019, , 809-810.		0
523	Acquired Coagulation Disorders Caused by Inhibitors. , 2019, , 80-92.		1
524	Surgery and Hemostasis. , 2019, , 696-720.		0
525	Sensitive and specific assessment of recombinant von Willebrand factor in platelet function analyzer. Platelets, 2019, 30, 264-270.	1.1	8

#	ARTICLE	IF	CITATIONS
526	How we make an accurate diagnosis of von Willebrand disease. <i>Thrombosis Research</i> , 2020, 196, 579-589.	0.8	18
527	Maternal and neonatal bleeding complications in relation to peripartum management in women with Von Willebrand disease: A systematic review. <i>Blood Reviews</i> , 2020, 39, 100633.	2.8	26
528	One piece of the puzzle: Population pharmacokinetics of FVIII during perioperative Haemate PÂ®/Humate PÂ® treatment in von Willebrand disease patients. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 295-305.	1.9	6
529	Higher rates of bleeding and use of treatment products among young boys compared to girls with von Willebrand disease. <i>American Journal of Hematology</i> , 2020, 95, 10-17.	2.0	12
530	An international survey to inform priorities for new guidelines on von Willebrand disease. <i>Haemophilia</i> , 2020, 26, 106-116.	1.0	32
531	Practical Recommendations for Treatment of Dental Patients with Congenital Bleeding Disorders during the Covid-19 Pandemic: A Narrative Review. <i>International Journal of Environmental Research and Public Health</i> , 2020, 17, 7245.	1.2	6
532	Low VWF: insights into pathogenesis, diagnosis, and clinical management. <i>Blood Advances</i> , 2020, 4, 3191-3199.	2.5	22
533	Classification of von Willebrand disease in the context of modern contemporary von Willebrand factor testing methodologies. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2020, 4, 952-957.	1.0	8
534	Prospective evaluation of ISTHâ€BAT as a predictor of bleeding disorder in adolescents presenting with heavy menstrual bleeding in a multidisciplinary hematology clinic. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 2542-2550.	1.9	21
535	Understanding the risks of total hip arthroplasty in patients with von Willebrandâ€™s disease. <i>Journal of Orthopaedic Surgery</i> , 2020, 28, 230949902096024.	0.4	1
536	&lt;p&gt;An Open-Label Extension Study to Assess the Long-Term Efficacy and Safety of a Plasma-Derived von Willebrand Factor (VWF)/Factor VIII (FVIII) Concentrate in Patients with von Willebrand Disease (SWIFT-VWDext Study)&lt;p&gt;. <i>Journal of Blood Medicine</i> , 2020, Volume 11, 345-356.	0.7	1
537	Molecular basis of hemostatic and thrombotic diseases. , 2020, , 229-246.		0
538	Population Pharmacokinetic Modeling of von Willebrand Factor Activity in von Willebrand Disease Patients after Desmopressin Administration. <i>Thrombosis and Haemostasis</i> , 2020, 120, 1407-1416.	1.8	3
539	Platelet Dysfunction Diseases and Conditions: Clinical Implications and Considerations. <i>Advances in Therapy</i> , 2020, 37, 3707-3722.	1.3	1
540	The relationship between ABO blood group, von Willebrand factor, and primary hemostasis. <i>Blood</i> , 2020, 136, 2864-2874.	0.6	75
541	How I treat von Willebrand disease. <i>Thrombosis Research</i> , 2020, 196, 618-625.	0.8	17
542	von Willebrand factor variant D1472H has no effect in mice with humanized VWF-platelet interactions. <i>Blood Advances</i> , 2020, 4, 4065-4068.	2.5	1
543	ADAMTSâ€13 and bleeding phenotype in von Willebrand disease. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2020, 4, 1331-1339.	1.0	3

#	ARTICLE	IF	CITATIONS
544	New developments in von Willebrand disease. <i>British Journal of Haematology</i> , 2020, 191, 329-339.	1.2	27
545	Low von Willebrand factor in pediatric patients: Retrospective analysis of 293 cases informs diagnostic and therapeutic decision making. <i>Pediatric Blood and Cancer</i> , 2020, 67, e28497.	0.8	2
546	Differentiation of Patients with Symptomatic Low von Willebrand Factor from Those with Asymptomatic Low von Willebrand Factor. <i>Thrombosis and Haemostasis</i> , 2020, 120, 793-804.	1.8	1
547	Pediatric Bleeding Disorders. , 2020, , .		1
548	Spinal Cord Stimulator Placement in Patient With von Willebrand Disease. <i>A&amp;A Practice</i> , 2020, 14, 149-151.	0.2	1
549	Evaluation of a fully automated von Willebrand factor assay panel for the diagnosis of von Willebrand disease. <i>Haemophilia</i> , 2020, 26, 298-305.	1.0	7
550	Perioperative management for patients with von Willebrand disease: Defining the optimal approach. <i>European Journal of Haematology</i> , 2020, 105, 365-377.	1.1	5
551	The impact of foetal restrictions on mode of delivery in women with inherited bleeding disorders. <i>European Journal of Haematology</i> , 2020, 105, 555-560.	1.1	1
552	Standardizing care to manage bleeding disorders in adolescents with heavy mensesâ€”A joint project from the ISTH pediatric/neonatal and women's health SSCs. <i>Journal of Thrombosis and Haemostasis</i> , 2020, 18, 2759-2774.	1.9	11
553	&lt;p&gt;Pharmacokinetics, Efficacy and Safety of a Plasma-Derived VWF/FVIII Concentrate (Formulation) Tj ETQq1 1 0.784314 rgBT / Dv Medicine, 2020, Volume 11, 213-225.	0.7	6
554	Navigating the Myriad of von Willebrand Factor Assays. <i>Hamostaseologie</i> , 2020, 40, 431-442.	0.9	19
555	Utility of the platelet function analyser (PFA-100/200) for exclusion or detection of von Willebrand disease: A study 22Âyears in the making. <i>Thrombosis Research</i> , 2020, 188, 17-24.	0.8	20
556	An integrated approach to inherited platelet disorders: results from a research collaborative, the Sydney Platelet Group. <i>Pathology</i> , 2020, 52, 243-255.	0.3	15
557	&lt;p&gt;Potential Undiagnosed VWD Or Other Mucocutaneous Bleeding Disorder Cases Estimated From Private Medical Insurance Claims&lt;p&gt;. <i>Journal of Blood Medicine</i> , 2020, Volume 11, 1-11.	0.7	11
558	Hemophilia and von Willebrand Disease: A Review of Emergency Department Management. <i>Journal of Emergency Medicine</i> , 2020, 58, 756-766.	0.3	8
559	von Willebrand factor binding to myosin assists in coagulation. <i>Blood Advances</i> , 2020, 4, 174-180.	2.5	5
560	Low VWF levels in children and lack of association with bleeding in children undergoing tonsillectomy. <i>Blood Advances</i> , 2020, 4, 100-105.	2.5	20
561	Desmopressin acetate use in von Willebrandâ€™s disease: a survey on current practices in Brazil. <i>Hematology, Transfusion and Cell Therapy</i> , 2021, 43, 43-49.	0.1	1

#	ARTICLE	IF	CITATIONS
562	Women and girls with heavy menstrual bleeding and inherited bleeding disorders: A call to action for the Haemophilia Treatment Centre Nurse. <i>Haemophilia</i> , 2021, 27, 82-86.	1.0	4
563	<scp>Postâ€Tonsillectomy</scp> Hemorrhage and the Diagnosis of Occult Pediatric Coagulopathies. <i>Laryngoscope</i> , 2021, 131, E2069-E2073.	1.1	4
564	Dental extraction in a patient with undiagnosed Von Willebrandâ€™s Disease: a case report. <i>Australian Dental Journal</i> , 2021, 66, 105-111.	0.6	2
565	Laboratory variability in the diagnosis of type 2 VWD variants. <i>Journal of Thrombosis and Haemostasis</i> , 2021, 19, 131-138.	1.9	10
566	Evaluation and Management of Coagulopathies and Thrombophilias in Pediatric Patients. <i>Clinics in Laboratory Medicine</i> , 2021, 41, 83-100.	0.7	2
567	Assessment of von Willebrand disease and pregnancy outcomes at regional Australian hospitals. <i>European Journal of Haematology</i> , 2021, 106, 456-466.	1.1	5
568	Management of elective procedures in low von Willebrand factor patients in the LoVIC study. <i>Journal of Thrombosis and Haemostasis</i> , 2021, 19, 701-710.	1.9	7
569	Thrombocytosis with acquired von Willebrand disease in an adolescent with sickle cell disease. <i>Clinical Case Reports (discontinued)</i> , 2021, 9, 457-460.	0.2	0
570	How we diagnose 2M von Willebrand disease (VWD): Use of a strategic algorithmic approach to distinguish 2M VWD from other VWD types. <i>Haemophilia</i> , 2021, 27, 137-148.	1.0	13
571	Dihydroartemisinin inhibits the expression of von Willebrand factor by downregulation of transcription factor ERG in endothelial cells. <i>Fundamental and Clinical Pharmacology</i> , 2021, 35, 321-330.	1.0	3
572	Type 3 von Willebrand Disease in Pregnancy: A Systematic Literature Review. <i>American Journal of Perinatology</i> , 2021, 38, 436-448.	0.6	6
573	Totally extraperitoneal inguinal hernia repair in patients with hemophilia and von Willebrand disease. Prospective controlled study. <i>Wideochirurgia I Inne Techniki Maloinwazyjne</i> , 2021, 16, 552-559.	0.3	0
574	Blood Disorders. , 2021, , 641-664.		0
575	Hemostatic Agents and Blood Components Used to Stop Bleeding. , 2021, , 425-443.		0
576	Successful Perioperative Management of Orthotopic Cardiac Transplantation in a Pediatric Patient With Concurrent Congenital von Willebrand Disease and Acquired von Willebrand Syndrome Using Recombinant von Willebrand Factor. <i>Journal of Cardiothoracic and Vascular Anesthesia</i> , 2022, 36, 724-727.	0.6	2
578	Criteria for low von Willebrand factor diagnosis and risk score to predict future bleeding. <i>Journal of Thrombosis and Haemostasis</i> , 2021, 19, 719-731.	1.9	5
579	Heavy Menstrual Bleeding. , 2021, , 255-265.		0
580	ASH ISTH NHF WFH 2021 guidelines on the management of von Willebrand disease. <i>Blood Advances</i> , 2021, 5, 301-325.	2.5	152

#	ARTICLE	IF	CITATIONS
581	Diseases of the Coagulation System: Hemophilia, Von Willebrands Disease, Cryoglobulinemia, and Inborn Errors of Factor Synthesis. , 2021, , 121-127.		0
582	Bleeding Associated with Connective Tissue Disorders. , 2021, , 201-209.		0
583	Thrombocytopenia: Perioperative Considerations for Patients Undergoing Cardiac Surgery. Journal of Cardiothoracic and Vascular Anesthesia, 2021, , .	0.6	2
584	The effect of age at diagnosis of type 1 von Willebrand disease on diagnostic laboratory values: A paediatric perspective. Haemophilia, 2021, 27, e412-e414.	1.0	1
585	Incidence of von Willebrand disease in Denmark, 1995â€2016: A cohort study. Haemophilia, 2021, 27, 277-282.	1.0	4
586	Type 2B von Willebrand Disease in Pregnancy: A Systematic Literature Review. Seminars in Thrombosis and Hemostasis, 2021, 47, 201-216.	1.5	9
587	Toward Personalized Treatment for Patients with Low von Willebrand Factor and Quantitative von Willebrand Disease. Seminars in Thrombosis and Hemostasis, 2021, 47, 192-200.	1.5	2
588	Current Challenges in the Peripartum Management of Women with von Willebrand Disease. Seminars in Thrombosis and Hemostasis, 2021, 47, 217-228.	1.5	12
589	Management of a hemispherotomy for Rasmussen encephalitis in a patient with mild factor 10 deficiency and low von Willebrand factor levels. Haemophilia, 2021, 27, e513-e516.	1.0	1
590	Economic burden of major bleeding events in commercially insured patients with von Willebrand disease based on claims data from the United States. Journal of Managed Care & Specialty Pharmacy, 2021, 27, 175-185.	0.5	5
591	Population pharmacokinetics of the von Willebrand factorâ€factor VIII interaction in patients with von Willebrand disease. Blood Advances, 2021, 5, 1513-1522.	2.5	5
592	Translating the success of prophylaxis in haemophilia to von Willebrand disease. Thrombosis Research, 2021, 199, 67-74.	0.8	15
593	Delivery and neuraxial technique outcomes in patients with hemophilia and in hemophilia carriers: a systematic review. Journal of Anesthesia, 2021, 35, 288-302.	0.7	4
594	Occurrence rates of von Willebrand disease among people receiving care in specialized treatment centres in the United States. Haemophilia, 2021, 27, 445-453.	1.0	13
595	Obstacles to Early Diagnosis and Treatment of Inherited von Willebrand Disease: Current Perspectives. Journal of Blood Medicine, 2021, Volume 12, 165-175.	0.7	4
597	Anesthetic Management of Von Willebrand Disease in Pregnancy: A Retrospective Analysis of a Large Case Series. Anesthesia and Analgesia, 2021, 133, 1244-1250.	1.1	7
598	von Willebrand disease: what does the future hold?. Blood, 2021, 137, 2299-2306.	0.6	13
599	Translation, validation, and usability of the International Society on Thrombosis and Haemostasis Bleeding Assessment Tool (Selfâ€STHâ€BAT). European Journal of Haematology, 2021, 107, 104-110.	1.1	5



#	ARTICLE	IF	CITATIONS
600	Characteristics, complications, and sites of bleeding among infants and toddlers less than 2 years of age with VWD. <i>Blood Advances</i> , 2021, 5, 2079-2086.	2.5	4
601	Elevated von Willebrand factor levels during heavy menstrual bleeding episodes limit the diagnostic utility for von Willebrand disease. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12513.	1.0	15
602	Traumatic splenic laceration with delayed rupture secondary to coughing in a patient with Von Willebrand disease. <i>American Journal of Emergency Medicine</i> , 2021, 50, 812.e5-812.e7.	0.7	0
603	Combined effects of plasma von Willebrand factor and platelet measures on the risk of incident venous thromboembolism. <i>Blood</i> , 2021, 138, 2269-2277.	0.6	13
604	Von Willebrand factor multimeric assay: novel diagnostics capabilities. <i>Russian Journal of Pediatric Hematology and Oncology</i> , 2021, 8, 35-41.	0.1	0
605	Inactivation of Cerebral Cavernal Malformation Genes Results in Accumulation of von Willebrand Factor and Redistribution of Weibel-Palade Bodies in Endothelial Cells. <i>Frontiers in Molecular Biosciences</i> , 2021, 8, 622547.	1.6	7
606	Genotypes of European and Iranian patients with type 3 von Willebrand disease enrolled in 3WINTERS-IPS. <i>Blood Advances</i> , 2021, 5, 2987-3001.	2.5	11
607	Economic Burden Associated with Major Surgery in Patients with von Willebrand Disease: A United States Retrospective Administrative Database Analysis. <i>Journal of Blood Medicine</i> , 2021, Volume 12, 699-708.	0.7	3
608	Management of large volume liposuction in lipedema patients with von Willebrand disease: A systematic review and treatment algorithm. <i>Clinical Hemorheology and Microcirculation</i> , 2021, 78, 311-324.	0.9	1
609	Von Willebrand Disease. <i>Hematology/Oncology Clinics of North America</i> , 2021, 35, 1085-1101.	0.9	14
610	Ex vivo human placental transfer study on recombinant Von Willebrand factor (rVWF). <i>Placenta</i> , 2021, 111, 69-75.	0.7	2
611	Diagnosis of von Willebrand disease: An assessment of the quality of testing in North American laboratories. <i>Haemophilia</i> , 2021, 27, e713-e720.	1.0	12
612	Addressing unmet needs in rare bleeding disorders: selected poster extracts of recent research in hemophilia A and von Willebrand disease presented at the 14th Annual Congress of the European Association for Haemophilia and Allied Disorders (EAHAD) (Feb 3-5, 2021; virtual congress). <i>Expert Review of Hematology</i> , 2021, 14, 1-18.	1.0	2
613	Maternal and neonatal bleeding complications in relation to peripartum management in hemophilia carriers: A systematic review. <i>Blood Reviews</i> , 2021, 49, 100826.	2.8	11
614	A Comparative Evaluation of an Automated Functional Assay for Von Willebrand Factor Activity in Type 1 Von Willebrand Disease. <i>International Journal of General Medicine</i> , 2021, Volume 14, 5167-5174.	0.8	0
615	Bleeding assessment tools in the diagnosis of VWD in adults and children: a systematic review and meta-analysis of test accuracy. <i>Blood Advances</i> , 2021, 5, 5023-5031.	2.5	6
616	Inherited Bleeding Disorders. , 0, , 825-841.		4
621	A rat model of severe VWD by elimination of the VWF gene using CRISPR/Cas9. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2020, 4, 64-71.	1.0	2

#	ARTICLE	IF	CITATIONS
622	von Willebrand Disease: Differential Diagnosis and Diagnostic Approach to Specific Subtypes. , 2016, , 285-293.		2
623	Diagnosis in General. , 2017, , 87-95.		1
624	Investigation of haemostasis. , 2012, , 393-445.		8
625	Von Willebrand Disease and Hemorrhagic Abnormalities of Platelet and Vascular Function. , 2012, , 1131-1136.		1
627	Early Resolution of Heyde's Syndrome following Transcatheter Aortic Valve Replacement. Seminars in Thrombosis and Hemostasis, 2021, 47, 102-104.	1.5	3
628	Genetic causes of haemophilia in women and girls. Haemophilia, 2021, 27, e164-e179.	1.0	28
629	Management of von Willebrand disease with a factor VIIIâ€”poor von Willebrand factor concentrate: Results from a prospective observational postâ€”marketing study. Journal of Thrombosis and Haemostasis, 2020, 18, 1922-1933.	1.9	12
630	Increasing levels of von Willebrand factor and factor VIII with age in patients affected by von Willebrand disease. Journal of Thrombosis and Haemostasis, 2021, 19, 96-106.	1.9	7
631	Major haemorrhage related to surgery in patients with type 1 and possible type 1 von Willebrand disease. Thrombosis and Haemostasis, 2008, 100, 797-802.	1.8	15
632	Is my patient a bleeder? A diagnostic framework for mild bleeding disorders. Hematology American Society of Hematology Education Program, 2012, 2012, 466-74.	0.9	23
633	Making a diagnosis of VWD. Hematology American Society of Hematology Education Program, 2012, 2012, 161-167.	0.9	18
634	Is my patient a bleeder? A diagnostic framework for mild bleeding disorders. Hematology American Society of Hematology Education Program, 2012, 2012, 466-474.	0.9	51
635	Perioperative management of a pediatric patient with suspected type 1 von Willebrand disease undergoing tonsillectomy: a case report. JA Clinical Reports, 2019, 5, 54.	0.2	1
636	Is gingival bleeding a symptom of type 2 and 3 von Willebrand disease?. PLoS ONE, 2018, 13, e0191291.	1.1	6
637	A Multicenter Study on von Willebrand Disease Realities in Yeungnam Region. Clinical Pediatric Hematology-Oncology, 2019, 26, 46-54.	0.0	1
638	Gingival bleeding and oral hygiene in women with von Willebrand Disease (VWD): a pilot study. The Journal of Haemophilia Practice, 2017, 4, 49-57.	0.2	2
639	Preface to Special Issue: diagnosis and management of von Willebrand diseaseâ€”diverse approaches to a global and common bleeding disorder. Annals of Blood, 0, 3, 43-43.	0.4	2
640	The current status of remote diagnosis of von Willebrand disease in children in Russia. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 54-60.	0.1	2

#	ARTICLE	IF	CITATIONS
641	Recommendations for the transfusion management of patients in the peri-operative period. I. The pre-operative period. Blood Transfusion, 2011, 9, 19-40.	0.3	69
642	Recommendations for the transfusion management of patients in the peri-operative period. II. The intra-operative period. Blood Transfusion, 2011, 9, 189-217.	0.3	109
643	Recommendations for the implementation of a Patient Blood Management programme. Application to elective major orthopaedic surgery in adults. Blood Transfusion, 2016, 14, 23-65.	0.3	78
644	Laboratory Diagnosis of von Willebrand Disease. Clinical Laboratory Science: Journal of the American Society for Medical Technology, 2017, 30, 65-74.	0.1	5
645	Bleeding Disorders in Orthopedic Surgery. Orthopedics, 2012, 35, 1053-1062.	0.5	11
646	Droplet-Shaped Deep Intraretinal Hemorrhage as Initial Presentation of Von Willebrand Disease Type 1. Ophthalmic Surgery Lasers and Imaging Retina, 2016, 47, 1044-1048.	0.4	3
647	Von willebrand disease: An overview. Indian Journal of Pharmaceutical Sciences, 2011, 73, 7.	1.0	32
648	Treatment and Management Patterns of Patients with Von Willebrand Disease in the United States. Journal of Blood Disorders & Transfusion, 2013, 04, .	0.1	2
649	Endovascular coil embolization of unruptured intracranial aneurysm in a patient with von Willebrand's disease: case report. Journal of Neuroendovascular Therapy, 2009, 3, 181-186.	0.1	1
651	Methodological Challenges in Observational Research: A Pharmacoepidemiological Perspective. British Journal of Pharmaceutical Research, 2013, 3, 161-175.	0.4	6
652	Screening for von Willebrand disease does not impact posttonsillectomy bleeding in a low-risk population. Pediatric Blood and Cancer, 2021, 68, e29371.	0.8	0
653	Gynecologic and obstetric management of women with von Willebrand disease: summary of 3 systematic reviews of the literature. Blood Advances, 2022, 6, 228-237.	2.5	15
654	Cardiac Surgery in Patients With Blood Disorders. Heart Lung and Circulation, 2021, , .	0.2	3
655	Hematologic Complications of Pregnancy. , 2007, , 1044-1063.		2
656	von Willebrand Disease in Childhood Chronic ITP. The Korean Journal of Hematology, 2008, 43, 232.	0.7	0
657	The patient at risk from thrombosis and bleeding disorders. Reproductive Medicine and Assisted Reproductive Techniques Series, 2008, , 121-136.	0.1	0
658	Molecular Basis of Disorders of Hemostasis and Thrombosis. Molecular Pathology Library, 2010, , 511-528.	0.1	0
660	Association of CD40 Genotyping and its Protein Expression with Airway Inflammatory Diseases. Journal of Molecular Biomarkers & Diagnosis, 2011, 02, .	0.4	3

#	ARTICLE	IF	CITATIONS
663	Inherited Bleeding Disorders in Obstetrics. , 0, , 153-160.		0
664	Pathophysiology of Coagulation and Deficiencies of Coagulation Factors in the Newborn. , 2012, , 748-762.		0
668	Bleeding Tendency of a Light Chain (AL) Amyloidosis Patient Accompanied by Asymptomatic Plasma Cell Myeloma. Laboratory Medicine Online, 2013, 3, 183.	0.0	0
669	Prolonged PTT. , 2014, , 57-70.		0
670	von Willebrand Disease. , 2014, , .		0
671	Progress in the Diagnosis and Management of Type 1 von Willebrand Disease. , 2014, 11, .		0
672	Modern Patient Blood Management in Arthroplasty. , 2015, , 3-17.		1
673	Bronchial Thermoplasty by Application of Ultrasound Energy. Journal of Pulmonary & Respiratory Medicine, 2015, 05, .	0.1	0
674	Emergency Craniotomy for Extradural Hematoma Evacuation in a Known Case of von Willebrand Disease: Anesthetic Implications. Journal of Research & Innovation in Anesthesia, 2016, 1, 58-60.	0.0	0
675	Lessons from life: the role of the advanced nurse specialist in IBDs. The Journal of Haemophilia Practice, 2016, 3, 55-61.	0.2	1
676	Congenital Coagulation Disorders. , 2016, , 93-114.		0
677	Acquired Coagulation Disorders. , 2016, , 115-128.		0
678	Known Bleeding Disorders for Surgery. , 2016, , 67-76.		0
679	Retrospective Analysis of Epsilon-Amino-Caproic Acid (EACA) Therapy in Patients with Hypoproliferative Thrombocytopenia. Hematology & Transfusion International Journal, 2017, 4, .	0.1	2
680	Pathophysiology of Coagulation and Deficiencies of Coagulation Factors in Newborn. , 2017, , 1-23.		0
681	Intraoperative Coagulopathies. , 2017, , 319-326.		0
682	Prothrombin Time (PT), Activated Partial Thromboplastin Time (APTT) and International Normalized Ratio (INR) as Predictive Factors of Coagulopathy in Newly Diagnosed Hypertensive Patients. Hematology & Transfusion International Journal, 2017, 4, .	0.1	0
683	Diagnosis and management of von willebrand disease in Spain. Annals of Blood, 0, 3, 5-5.	0.4	4

#	ARTICLE	IF	CITATIONS
684	Inherited Bleeding Disorders in Iraq and Consanguineous Marriage. International Journal of Hematology-Oncology and Stem Cell Research, 0, , .	0.3	4
685	Update in the Investigation of von Willebrand Disease. , 2019, , 327-338.		0
686	Desmopressin acetate (DDAVP) for preventing and treating acute bleeds during pregnancy in women with congenital bleeding disorders. The Cochrane Library, 2021, 2021, CD009824.	1.5	4
687	PREVALENCE OF FACTOR V LEIDEN AND PROTHROMBIN G20210A IN WOMEN WITH VON WILLEBRAND DISEASE TYPE 1. Gematologiya I Transfuziologiya, 2019, 64, 60-65.	0.1	1
688	Enfermedad de Von Willebrand tipo III en una paciente obstétrica. Revista Universitas Medica, 2019, 60, 1-9.	0.0	0
689	Evidence-Based Minireview: Perioperative management of the VWD patient at elevated thrombotic risk. Hematology American Society of Hematology Education Program, 2019, 2019, 601-603.	0.9	0
690	Presentation and Management of Type 2 von Willebrand Disease. , 2020, , 99-112.		0
691	Evaluation of the Adolescent with Heavy Menstrual Bleeding. , 2020, , 3-11.		0
692	Heavy Menstrual Bleeding in Adolescent Girls. Pediatric Annals, 2020, 49, e163-e169.	0.3	6
693	von Willebrand Disease. , 2021, , 233-245.		0
695	Commentary on the ASH ISTH NHF WFH 2021 guidelines on the diagnosis of VWD: reflections based on recent contemporary test data. Blood Advances, 2022, 6, 416-419.	2.5	21
696	Selected Disorders of the Blood and Hematopoietic System. , 2020, , 1-15.		0
697	Results of extended phenotyping of von Willebrand disease in the remote diagnostic program in children. Pediatric Hematology/Oncology and Immunopathology, 2020, 19, 102-106.	0.1	0
698	Evaluation of Abnormal Bleeding in Children. Pediatric Annals, 2020, 49, e36-e42.	0.3	3
699	Clinical Approach to Type 3 von Willebrand Disease. , 2020, , 113-125.		0
700	Bleeding Disorders in Pregnancy. , 2020, , 319-322.		0
701	Classification and Management of Type 1 von Willebrand Disease. , 2020, , 83-98.		0
702	Pathophysiology and Management of Acquired von Willebrand Syndrome. , 2020, , 127-137.		0

#	ARTICLE	IF	CITATIONS
703	Hemophilia A and von Willebrand deficiency: therapeutic implications. <i>Blood Coagulation and Fibrinolysis</i> , 2020, 31, 397-401.	0.5	1
704	Postpartum Hemorrhage in Patients with Type 1 von Willebrand Disease: A Systematic Review. <i>Seminars in Thrombosis and Hemostasis</i> , 2021, 48, .	1.5	1
705	Heavy Menstrual Bleeding in Adolescent: Normal or a Sign of an Underlying Disease?. <i>Seminars in Reproductive Medicine</i> , 2021, , .	0.5	1
706	Efficacy and safety evaluation of Fanhdi<sup>®</sup>, a plasma-derived factor VIII/ von Willebrand factor concentrate, in Von Willebrand's disease patients undergoing surgery or invasive procedures: A prospective clinical study. <i>Haemophilia</i> , 2022, 28, .	1.0	1
707	A highly-sensitive plasma von Willebrand factor ristocetin cofactor (VWF:RCo) activity assay by flow cytometry. <i>Journal of Thrombosis and Haemostasis</i> , 2008, 6, 323-330.	1.9	13
708	Anesthetic Management of Cardiac Surgery Patients with Uncommon Diseases. , 2021, , 209-223.		0
709	Clinical features and types of von Willebrand disease in women with menorrhagia referred to hematology clinic of kermanshah. <i>International Journal of Hematology-Oncology and Stem Cell Research</i> , 2013, 7, 1-5.	0.3	2
710	Current therapy in children and adolescents with von Willebrand disease. <i>Journal of Medicine and Life</i> , 2014, 7, 264-9.	0.4	1
712	Chitosan scaffold enhances growth factor release in wound healing in von Willebrand disease. <i>International Journal of Clinical and Experimental Medicine</i> , 2015, 8, 15611-20.	1.3	6
713	Towards personalised therapy for von Willebrand disease: a future role for recombinant products. <i>Blood Transfusion</i> , 2016, 14, 262-76.	0.3	15
714	Making a diagnosis of VWD. <i>Hematology American Society of Hematology Education Program</i> , 2012, 2012, 161-7.	0.9	6
715	Diagnosis and management of patients with von Willebrand's disease in Italy: an Expert Meeting Report. <i>Blood Transfusion</i> , 2018, 16, 326-328.	0.3	0
716	Current challenges in the diagnosis and management of patients with inherited von Willebrand's disease in Italy: an Expert Meeting Report on the diagnosis and surgical and secondary long-term prophylaxis. <i>Blood Transfusion</i> , 2018, 16, 371-381.	0.3	8
717	Inherited Bleeding Disorders in Iraq and Consanguineous Marriage. <i>International Journal of Hematology-Oncology and Stem Cell Research</i> , 2018, 12, 273-281.	0.3	2
719	Effectiveness and safety of hFVIII/VWF concentrate (Voncento) in patients with inherited von Willebrand disease requiring surgical procedures: the OPALE multicentre observational study. <i>Blood Transfusion</i> , 2021, 19, 152-157.	0.3	1
720	Disorders of coagulation. , 2022, , 287-340.		1
721	Bleeding patterns in patients before and after diagnosis of von Willebrand disease: Analysis of a US medical claims database. <i>Haemophilia</i> , 2021, , .	1.0	5
722	Percutaneous left atrial appendage closure in patients with primary hemostasis disorders and atrial fibrillation. <i>Journal of Interventional Cardiac Electrophysiology</i> , 2022, 64, 497-509.	0.6	4

#	ARTICLE	IF	CITATIONS
723	Multimer Analysis of Von Willebrand Factor in Von Willebrand Disease with a Hydrasys Semi-Automatic Analyzer—Single-Center Experience. <i>Diagnostics</i> , 2021, 11, 2153.	1.3	7
724	Von Willebrand Disease. <i>Pediatrics in Review</i> , 2014, 35, 136-137.	0.2	0
725	Efficacy of emicizumab in von Willebrand disease (VWD) patients with and without alloantibodies to von Willebrand factor (VWF): Report of two cases and review of literature. <i>Haemophilia</i> , 2022, 28, 286-291.	1.0	6
726	Bleeding Disorders. , 2021, , .		0
727	Gastrointestinal Bleeding in Congenital Bleeding Disorders. <i>Seminars in Thrombosis and Hemostasis</i> , 2022, 48, 529-541.	1.5	2
728	The predictive value of bleeding score on the diagnosis of Von Willebrand disease in children applied to the hematologic clinic with epistaxis. <i>Journal of Health Sciences and Medicine</i> , 2022, 5, 22-25.	0.0	0
729	The 2021 von Willebrand disease guidelines: Clarity and controversy. <i>Haemophilia</i> , 2022, 28, 1-3.	1.0	15
730	A novel mouse model of type 2N VWD was developed by CRISPR/Cas9 gene editing and recapitulates human type 2N VWD. <i>Blood Advances</i> , 2022, 6, 2778-2790.	2.5	1
731	Selected Disorders of the Blood and Hematopoietic System. , 2022, , 1831-1845.		0
732	The impact of aberrant von Willebrand factor-GPIb $\beta$ interaction on megakaryopoiesis and platelets in humanized type 2B von Willebrand disease model mouse. <i>Haematologica</i> , 2022, , .	1.7	1
734	Laboratory assays of VWF activity and use of desmopressin trials in the diagnosis of VWD: a systematic review and meta-analysis. <i>Blood Advances</i> , 2022, 6, 3735-3745.	2.5	3
736	Integrative genetic and immune cell analysis of plasma proteins in healthy donors identifies novel associations involving primary immune deficiency genes. <i>Genome Medicine</i> , 2022, 14, 28.	3.6	8
737	2B von Willebrand disease diagnosis: Considerations reflecting on 2021 multisociety guidelines. <i>Research and Practice in Thrombosis and Haemostasis</i> , 2021, 5, e12635.	1.0	8
738	Abnormal uterine bleeding. Hemostatic therapy: hematologist point of view. <i>Gynecology</i> , 2022, 24, 157-162.	0.1	2
739	Haematological Problems in Pregnancy. , 0, , 270-281.		0
740	Recommended indications for the administration of polyclonal immunoglobulin preparations. <i>Acta Clinica Belgica</i> , 2011, 66, 346-60.	0.5	5
746	Non-factor therapies for bleeding disorders: A primer for the general haematologist. <i>EJHaem</i> , 2022, 3, 584-595.	0.4	5
747	VWF $\Delta$ Gly2752Ser, a novel non-cysteine substitution variant in the CK domain, exhibits severe secretory impairment by hampering C-terminal dimer formation. <i>Journal of Thrombosis and Haemostasis</i> , 2022, , .	1.9	1

#	ARTICLE	IF	CITATIONS
748	Postpartum Hemorrhage in Women with von Willebrand Disease: Consider Other Etiologies. Journal of Obstetrics and Gynaecology Canada, 2022, 44, 972-977.	0.3	2
749	Hematologic Complications of Pregnancy. , 2017, , 947-964.e2.		0
750	Efficacy and safety of von Willebrand factor concentrate almost devoid of factor VIII (Wilfactin) in paediatric patients under 6 years of age with severe von Willebrand disease.. Blood Transfusion, 2022, , .	0.3	0
751	Case Report: An Unusual Case of Nutcracker Syndrome and Literature Review. Frontiers in Urology, 2022, 2, .	0.2	0
752	Genetic Alterations, DNA Methylation, Alloantibodies and Phenotypic Heterogeneity in Type III von Willebrand Disease. Genes, 2022, 13, 971.	1.0	0
753	The p. <sc>P1127S</sc> pathogenic variant lowers von Willebrand factor levels through higher affinity for the macrophagic scavenger receptor <sc>LRP1</sc> : clinical phenotype and pathogenic mechanisms. Journal of Thrombosis and Haemostasis, 0, , .	1.9	2
754	<sc>JTH</sc> in Clinic â€•Obstetric bleeding: <sc>VWD</sc> and other inherited bleeding disorders. Journal of Thrombosis and Haemostasis, 0, , .	1.9	1
755	Phenotypic and Genotypic Signatures of VWF Exon 18 in Eastern Saudi Patients Previously Diagnosed with Type 1 von Willebrand Disease. International Journal of General Medicine, 0, Volume 15, 5385-5394.	0.8	1
756	Mass Spectrometry Approaches Empowering Neuropeptide Discovery and Therapeutics. Pharmacological Reviews, 2022, 74, 662-679.	7.1	5
757	Recent advances in therapeutic options for rare hemostatic disorders: selected poster extracts of recent research in hemophilia A, congenital hemophilia with inhibitors, von Willebrand disease, and thrombotic thrombocytopenic purpura presented at the 29th congress of the International Society on Thrombosis and Haemostasis (ISTH 2021, Jul 17â€•21; virtual congress). Expert Review of Hematology, 0, , 1-10.	1.0	0
759	Evaluating Performance of Contemporary and Historical von Willebrand Factor (VWF) Assays in the Laboratory Identification of von Willebrand Disease (VWD): The Australasian Experience. Seminars in Thrombosis and Hemostasis, 2022, 48, 711-731.	1.5	11
760	Longitudinal bleeding assessment in von willebrand disease utilising an interim bleeding score. Journal of Thrombosis and Haemostasis, 0, , .	1.9	5
761	Diagnosis of Bleeding Disorders in Adolescents Hospitalized for Heavy Menstrual Bleeding. TH Open, 0, , .	0.7	0
762	Emergent Reversal of Antithrombotics and Treatment of Life-Threatening Bleeding from Coagulopathies: A Clinical Review. Journal of Emergency Medicine, 2022, 63, 17-48.	0.3	3
763	Impact of von Willebrand Disease on Women's Health Outcomes: A Matched Cohort Database Study. Journal of Women's Health, 2022, 31, 1262-1270.	1.5	6
764	Disorders of Hemostasis and Thrombosis. , 2023, , 173-211.		0
765	How Do Laboratories Perform von Willebrand Disease Diagnostics and Classification of von Willebrand Disease Patients? Results from External Quality Data and an International Survey. Seminars in Thrombosis and Hemostasis, 0, , .	1.5	0
766	Clinical, economic, and healthâ€•related quality of life burden associated with von Willebrand disease in adults and children: Systematic and targeted literature reviews. Haemophilia, 2023, 29, 411-422.	1.0	3



#	ARTICLE	IF	CITATIONS
767	Acquired von Willebrand Syndrome in a Patient with Multiple Comorbidities, Including MALT Lymphoma with IgA Monoclonal Gammopathy and Hyperviscosity Syndrome. <i>Internal Medicine</i> , 2023, 62, 605-611.	0.3	3
768	Diagnostic Testing for von Willebrand Disease: Trends and Insights from North American Laboratories over the Last Decade. <i>Seminars in Thrombosis and Hemostasis</i> , 2022, 48, 700-710.	1.5	6
769	How Do Laboratories Perform von Willebrand Disease Diagnostics and Classification of von Willebrand Disease Patients? Results from External Quality Data and an International Survey. <i>Seminars in Thrombosis and Hemostasis</i> , 0, , .	1.5	0
770	Laboratory Diagnosis of von Willebrand Disease (VWD): Geographical Perspectives. <i>Seminars in Thrombosis and Hemostasis</i> , 2022, 48, 750-766.	1.5	10
771	Evaluation for Bleeding Disorders in Suspected Child Abuse. <i>Pediatrics</i> , 2022, 150, .	1.0	5
772	Von Willebrand disease - detection, diagnostics and treatment. <i>Medicinski Pregled</i> , 2022, 75, 147-151.	0.1	0
773	How Do Laboratories Perform von Willebrand Disease Diagnostics and Classification of von Willebrand Disease Patients? Results from External Quality Data and an International Survey. <i>Seminars in Thrombosis and Hemostasis</i> , 2022, 48, 739-749.	1.5	3
774	Patients with von Willebrand disease in China: Results of an online survey. <i>Haemophilia</i> , 2023, 29, 230-239.	1.0	3
775	Effectiveness of individualized management using WILFACTIN <sup>®</sup> in patients with von Willebrand disease during surgical procedures: A single-center study. <i>Thrombosis Research</i> , 2022, 220, 88-90.	0.8	1
777	Memorial notice: William L. Nichols, MD. <i>International Journal of Laboratory Hematology</i> , 2023, 45, 139-140.	0.7	0
778	Diagnosis of von Willebrand disease during the management of deep neck abscess: A case report. <i>Acta Oto-Laryngologica Case Reports</i> , 2022, 7, 44-47.	0.1	0
779	Prevalence of Bleeding Symptoms in Denmark Using the Patient-administered Bleeding Assessment Tool (self-BAT). <i>Haemophilia</i> , 0, , .	1.0	1
780	The diagnosis, natural history, and management of von Willebrand disease in women in the age of guidelines. <i>Expert Review of Hematology</i> , 2023, 16, 435-450.	1.0	1
781	von Willebrand factor neutralizing and non-neutralizing alloantibodies in 213 subjects with type 3 von Willebrand disease enrolled in 3WINTERS-IPS. <i>Journal of Thrombosis and Haemostasis</i> , 2023, 21, 787-799.	1.9	0
782	Switch to pdVWF:pdFVIII concentrate for prophylaxis in a paediatric patient with Type 3 von Willebrand disease: a case report. <i>Hematology</i> , 2023, 28, .	0.7	0
783	Von Willebrand Disease, Hemophilia, and Other Inherited Bleeding Disorders in Pregnancy. <i>Obstetrics and Gynecology</i> , 2023, 141, 493-504.	1.2	1
784	Monoglyceride Lipase Deficiency Is Associated with Altered Thrombogenesis in Mice. <i>International Journal of Molecular Sciences</i> , 2023, 24, 3116.	1.8	0
786	The Role of the von Willebrand Factor Collagen-Binding Assay (VWF:CB) in the Diagnosis and Treatment of von Willebrand Disease (VWD) and Way Beyond: A Comprehensive 36-Year History. <i>Seminars in Thrombosis and Hemostasis</i> , 2024, 50, 043-080.	1.5	2

#	ARTICLE	IF	CITATIONS
787	The 2021 guidelines on the diagnosis of von Willebrand disease: A comparison with current clinical practice in Spanish centers. Haemophilia, 2023, 29, 925-927.	1.0	1
788	Perioperative Outcomes of Patients with Bleeding Disorders Undergoing Major Surgery at an Academic Hemophilia Treatment Center. Clinical and Applied Thrombosis/Hemostasis, 2023, 29, 107602962311650.	0.7	7
789	A comparative study in patients with type 2 von Willebrand disease using 4 different platelet-dependent von Willebrand factor assays. Research and Practice in Thrombosis and Haemostasis, 2023, 7, 100139.	1.0	3
790	Diagnóstico de la enfermedad de von Willebrand. Medicina Y Laboratorio, 2023, 27, 139-155.	0.0	1
792	Laboratory Testing for von Willebrand Factor: Factor VIII Binding for the Diagnosis or Exclusion of Type 2N von Willebrand Disease: An Update. Methods in Molecular Biology, 2023, , 679-691.	0.4	0
793	Laboratory Testing for von Willebrand Disease Using a Composite Rapid 3-Test Chemiluminescence-Based von Willebrand Factor Assay Panel. Methods in Molecular Biology, 2023, , 647-667.	0.4	3
802	Congenital bleeding and thrombotic disorders. , 2024, , 51-73.		0
807	Low von Willebrand Levels, von Willebrand Disease Does not Make. , 2023, , 351-353.		0
809	von Willebrand Disease: An Update on Diagnosis and Treatment. , 2023, , 77-105.		0