Flora Peyvandi

List of Publications by Year
in descending order

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663 papers	33,075 citations	7672 79 h-index	7427 157 g-index
685 all docs	685 docs citations	685 times ranked	39477 citing authors

#	Article	IF	Citations
1	Genetic studies of body mass index yield new insights for obesity biology. Nature, 2015, 518, 197-206.	13.7	3,823
2	Large-scale association analysis identifies 13 new susceptibility loci for coronary artery disease. Nature Genetics, 2011, 43, 333-338.	9.4	1,685
3	Genomewide Association Study of Severe Covid-19 with Respiratory Failure. New England Journal of Medicine, 2020, 383, 1522-1534.	13.9	1,548
4	Hypercoagulability of COVIDâ€19 patients in intensive care unit: A report of thromboelastography findings and other parameters of hemostasis. Journal of Thrombosis and Haemostasis, 2020, 18, 1738-1742.	1.9	1,070
5	Genome-wide association of early-onset myocardial infarction with single nucleotide polymorphisms and copy number variants. Nature Genetics, 2009, 41, 334-341.	9.4	990
6	Guidelines on the diagnosis and management of thrombotic thrombocytopenic purpura and other thrombotic microangiopathies. British Journal of Haematology, 2012, 158, 323-335.	1.2	700
7	Caplacizumab Treatment for Acquired Thrombotic Thrombocytopenic Purpura. New England Journal of Medicine, 2019, 380, 335-346.	13.9	625
8	Caplacizumab for Acquired Thrombotic Thrombocytopenic Purpura. New England Journal of Medicine, 2016, 374, 511-522.	13.9	480
9	Recessively inherited coagulation disorders. Blood, 2004, 104, 1243-1252.	0.6	479
10	New susceptibility locus for coronary artery disease on chromosome 3q22.3. Nature Genetics, 2009, 41, 280-282.	9.4	440
11	A Randomized Trial of Factor VIII and Neutralizing Antibodies in Hemophilia A. New England Journal of Medicine, 2016, 374, 2054-2064.	13.9	414
12	Coagulation factor activity and clinical bleeding severity in rare bleeding disorders: results from the European Network of Rare Bleeding Disorders. Journal of Thrombosis and Haemostasis, 2012, 10, 615-621.	1.9	362
13	Consensus on the standardization of terminology in thrombotic thrombocytopenic purpura and related thrombotic microangiopathies. Journal of Thrombosis and Haemostasis, 2017, 15, 312-322.	1.9	362
14	Genome-Wide Association Identifies Nine Common Variants Associated With Fasting Proinsulin Levels and Provides New Insights Into the Pathophysiology of Type 2 Diabetes. Diabetes, 2011, 60, 2624-2634.	0.3	335
15	The past and future of haemophilia: diagnosis, treatments, and its complications. Lancet, The, 2016, 388, 187-197.	6.3	331
16	ADAMTS13 autoantibodies in patients with thrombotic microangiopathies and other immunomediated diseases. Blood, 2005, 106, 1262-1267.	0.6	275
17	The Thrombogram in Rare Inherited Coagulation Disorders: Its Relation to Clinical Bleeding. Thrombosis and Haemostasis, 2002, 88, 576-582.	1.8	261
18	ADAMTS13 and anti-ADAMTS13 antibodies as markers for recurrence of acquired thrombotic thrombocytopenic purpura during remission. Haematologica, 2008, 93, 232-239.	1.7	250

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19	Rare bleeding disorders: diagnosis and treatment. Blood, 2015, 125, 2052-2061.	0.6	244
20	Bleeding and thrombosis in 55 patients with inherited afibrinogenaemia. British Journal of Haematology, 1999, 107, 204-206.	1.2	233
21	Clinical phenotypes and factor VII genotype in congenital factor VII deficiency. Thrombosis and Haemostasis, 2005, 93, 481-487.	1.8	218
22	Rare coagulation deficiencies. Haemophilia, 2002, 8, 308-321.	1.0	215
23	Complement activation in patients with COVID-19: AÂnovel therapeutic target. Journal of Allergy and Clinical Immunology, 2020, 146, 215-217.	1.5	210
24	Large-Scale Gene-Centric Analysis Identifies Novel Variants for Coronary Artery Disease. PLoS Genetics, 2011, 7, e1002260.	1.5	203
25	No Evidence of Association Between Prothrombotic Gene Polymorphisms and the Development of Acute Myocardial Infarction at a Young Age. Circulation, 2003, 107, 1117-1122.	1.6	191
26	ISTH guidelines for treatment of thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2020, 18, 2496-2502.	1.9	188
27	Abdominal Aortic Aneurysm Is Associated with a Variant in Low-Density Lipoprotein Receptor-Related Protein 1. American Journal of Human Genetics, 2011, 89, 619-627.	2.6	185
28	Rare Coagulation Disorders. Thrombosis and Haemostasis, 1999, 82, 1207-1214.	1.8	180
29	Current and novel biomarkers of thrombotic risk in COVID-19: a Consensus Statement from the International COVID-19 Thrombosis Biomarkers Colloquium. Nature Reviews Cardiology, 2022, 19, 475-495.	6.1	180
30	The ADAMTS13â€von Willebrand factor axis in COVIDâ€19 patients. Journal of Thrombosis and Haemostasis, 2021, 19, 513-521.	1.9	176
31	Pregnancy complications and obstetric care in women with inherited bleeding disorders. Haemophilia, 2013, 19, 1-10.	1.0	173
32	Evaluation and management of postpartum hemorrhage: consensus from an international expert panel. Transfusion, 2014, 54, 1756-1768.	0.8	167
33	ADAMTS13mutations and polymorphisms in congenital thrombotic thrombocytopenic purpura. Human Mutation, 2010, 31, 11-19.	1.1	165
34	Clinical manifestations and complications of childbirth and replacement therapy in 385 Iranian patients with type 3 von Willebrand disease. British Journal of Haematology, 2000, 111, 1236-1239.	1.2	152
35	Procoagulant imbalance in patients with non-alcoholic fatty liver disease. Journal of Hepatology, 2014, 61, 148-154.	1.8	149
36	von Willebrand factor cleaving protease (ADAMTS-13) and ADAMTS-13 neutralizing autoantibodies in 100 patients with thrombotic thrombocytopenic purpura. British Journal of Haematology, 2004, 127, 433-439.	1.2	142

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37	ISTH guidelines for the diagnosis of thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2020, 18, 2486-2495.	1.9	142
38	Smoking and the Risk of Mortality and Vascular and Respiratory Events in Patients Undergoing Major Surgery. JAMA Surgery, 2013, 148, 755.	2.2	140
39	ADAMTS13 activity to antigen ratio in physiological and pathological conditions associated with an increased risk of thrombosis. British Journal of Haematology, 2007, 138, 534-540.	1.2	135
40	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
41	Classification of rare bleeding disorders (RBDs) based on the association between coagulant factor activity and clinical bleeding severity. Journal of Thrombosis and Haemostasis, 2012, 10, 1938-1943.	1.9	129
42	A critical appraisal of oneâ€stage and chromogenic assays of factor VIII activity. Journal of Thrombosis and Haemostasis, 2016, 14, 248-261.	1.9	127
43	Complement activation and endothelial perturbation parallel COVID-19 severity and activity. Journal of Autoimmunity, 2021, 116, 102560.	3.0	127
44	Factor X Deficiency. Seminars in Thrombosis and Hemostasis, 2009, 35, 407-415.	1.5	125
45	Fibrinogen replacement therapy for congenital fibrinogen deficiency. Journal of Thrombosis and Haemostasis, 2011, 9, 1687-1704.	1.9	124
46	Genetic diagnosis of haemophilia and other inherited bleeding disorders. Haemophilia, 2006, 12, 82-89.	1.0	123
47	Introduction: Rare Bleeding Disorders: General Aspects of Clinical Features, Diagnosis, and Management. Seminars in Thrombosis and Hemostasis, 2009, 35, 349-355.	1.5	123
48	The bleeding score predicts clinical outcomes and replacement therapy in adults with von Willebrand disease. Blood, 2014, 123, 4037-4044.	0.6	123
49	Incidence of bleeding symptoms in 100 patients with inherited afibrinogenemia or hypofibrinogenemia. Journal of Thrombosis and Haemostasis, 2006, 4, 1634-1637.	1.9	122
50	Congenital factor X deficiency: spectrum of bleeding symptoms in 32 Iranian patients. British Journal of Haematology, 1998, 102, 626-628.	1.2	121
51	Multimorbidity and polypharmacy in the elderly: lessons from REPOSI. Internal and Emergency Medicine, 2014, 9, 723-734.	1.0	121
52	Effect of anakinra on mortality in patients with COVID-19: a systematic review and patient-level meta-analysis. Lancet Rheumatology, The, 2021, 3, e690-e697.	2.2	121
53	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
54	Factor V Deficiency. Seminars in Thrombosis and Hemostasis, 2009, 35, 382-389.	1.5	114

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55	Combined deficiency of factor V and factor VIII is due to mutations in either LMAN1 or MCFD2. Blood, 2006, 107, 1903-1907.	0.6	111
56	Symptoms of inherited factor V deficiency in 35 Iranian patients. British Journal of Haematology, 1998, 103, 1067-1069.	1.2	109
57	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
58	Residual plasmatic activity of ADAMTS13 is correlated with phenotype severity in congenital thrombotic thrombocytopenic purpura. Blood, 2012, 120, 440-448.	0.6	107
59	Gene Polymorphisms Predicting High Plasma Levels of Coagulation and Fibrinolysis Proteins. Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 755-759.	1.1	103
60	ADAMTS-13 assays in thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2010, 8, 631-640.	1.9	103
61	Inhibitor development in haemophilia according to concentrate. Thrombosis and Haemostasis, 2015, 113, 968-975.	1.8	103
62	Redefining outcomes in immune TTP: an international working group consensus report. Blood, 2021, 137, 1855-1861.	0.6	103
63	Pattern of symptoms in 93 Iranian patients with severe factor XIII deficiency. Journal of Thrombosis and Haemostasis, 2003, 1, 1852-1853.	1.9	101
64	Pharmacokinetics and safety of fibrinogen concentrate. Journal of Thrombosis and Haemostasis, 2009, 7, 2064-2069.	1.9	100
65	Molecular defects in type 3 von Willebrand disease: updated results from 40 multiethnic patients. Blood Cells, Molecules, and Diseases, 2003, 30, 264-270.	0.6	95
66	The Italian AICE-Genetics hemophilia A database: results and correlation with clinical phenotype. Haematologica, 2008, 93, 722-728.	1.7	95
67	Presentation and pattern of symptoms in 382 patients with Glanzmann thrombasthenia in Iran. American Journal of Hematology, 2004, 77, 198-199.	2.0	94
68	Force fluctuations during the Maximum Isometric Voluntary Contraction of the quadriceps femoris in haemophilic patients. Haemophilia, 2007, 13, 65-70.	1.0	94
69	Caplacizumab reduces the frequency of major thromboembolic events, exacerbations and death in patients with acquired thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2017, 15, 1448-1452.	1.9	94
70	Epidemiology and treatment of congenital fibrinogen deficiency. Thrombosis Research, 2012, 130, S7-S11.	0.8	93
71	Clinical manifestations in 28 Italian and Iranian patients with severe factor VII deficiency. Haemophilia, 1997, 3, 242-246.	1.0	92
72	Treatment of rare factor deficiencies other than hemophilia. Blood, 2019, 133, 415-424.	0.6	92

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73	Clinical manifestations and complications of childbirth and replacement therapy in 385 Iranian patients with type 3 von Willebrand disease. British Journal of Haematology, 2000, 111, 1236-1239.	1.2	91
74	CYP2C9 genotypes and dose requirements during the induction phase of oral anticoagulant therapy. Clinical Pharmacology and Therapeutics, 2004, 75, 198-203.	2.3	90
75	Factor VIII products and inhibitor development: the SIPPET study (survey of inhibitors in) Tj ETQq1 1 0.784314 rgBT /Overlock 10 Tf 50	1.0	90
76	Anakinra combined with methylprednisolone in patients with severe COVID-19 pneumonia and hyperinflammation: An observational cohort study. Journal of Allergy and Clinical Immunology, 2021, 147, 561-566.e4.	1.5	90
77	Role of von Willebrand factor in the haemostasis. Blood Transfusion, 2011, 9 Suppl 2, s3-8.	0.3	89
78	Thrombosis in Inflammatory Bowel Diseases: Role of Inherited Thrombophilia. American Journal of Gastroenterology, 2005, 100, 2036-2041.	0.2	85
79	Bleeding symptoms in 27 Iranian patients with the combined deficiency of factor V and factor VIII. British Journal of Haematology, 1998, 100, 773-776.	1.2	84
80	Hemophilic arthropathy: Current knowledge and future perspectives. Journal of Thrombosis and Haemostasis, 2021, 19, 2112-2121.	1.9	84
81	Molecular Analysis of the ERGIC-53 Gene in 35 Families With Combined Factor V-Factor VIII Deficiency. Blood, 1999, 93, 2253-2260.	0.6	83
82	Rare bleeding disorders. Haemophilia, 2008, 14, 202-210.	1.0	82
83	Hemostasis and menstruation: appropriate investigation for underlying disorders of hemostasis in women with excessive menstrual bleeding. Fertility and Sterility, 2005, 84, 1345-1351.	0.5	81
84	Rare bleeding disorders. Haemophilia, 2006, 12, 137-142.	1.0	76
85	Gender-differences in disease distribution and outcome in hospitalized elderly: Data from the REPOSI study. European Journal of Internal Medicine, 2014, 25, 617-623.	1.0	75
86	ADAMTSâ€13 activity and autoantibodies classes and subclasses as prognostic predictors in acquired thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2012, 10, 1556-1565.	1.9	74
87	Rotational thromboelastography for monitoring of fibrinogen concentrate therapy in fibrinogen deficiency. Blood Coagulation and Fibrinolysis, 2008, 19, 777-783.	0.5	71
88	Initial experience from a doubleâ€blind, placeboâ€controlled, clinical outcome study of ARC1779 in patients with thrombotic thrombocytopenic purpura. American Journal of Hematology, 2012, 87, 430-432.	2.0	71
89	Congenital afibrinogenemia: mutations leading to premature termination codons in fibrinogen Aî±-chain gene are not associated with the decay of the mutant mRNAs. Blood, 2001, 98, 3685-3692.	0.6	68
90	Anti-beta 2 glycoprotein I antibodies and the risk of myocardial infarction in young premenopausal women. Journal of Thrombosis and Haemostasis, 2007, 5, 2421-2428.	1.9	67

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91	Novel aspects of factor XIII deficiency. Current Opinion in Hematology, 2011, 18, 366-372.	1.2	67
92	Rare bleeding disorders â€“ bleeding assessment tools, laboratory aspects and phenotype and therapy of FXI deficiency. Haemophilia, 2014, 20, 71-75.	1.0	67
93	The European Hematology Association Roadmap for European Hematology Research: a consensus document. Haematologica, 2016, 101, 115-208.	1.7	67
94	Risk factors for inhibitor development in severe hemophilia A. Thrombosis Research, 2018, 168, 20-27.	0.8	67
95	Molecular Characterisation and Three-Dimensional Structural Analysis of Mutations in 21 Unrelated Families with Inherited Factor VII Deficiency. Thrombosis and Haemostasis, 2000, 84, 250-257.	1.8	67
96	Influence of 9p21.3 Genetic Variants on Clinical and Angiographic Outcomes in Early-Onset Myocardial Infarction. Journal of the American College of Cardiology, 2011, 58, 426-434.	1.2	66
97	Advances in the treatment of bleeding disorders. Journal of Thrombosis and Haemostasis, 2016, 14, 2095-2106.	1.9	66
98	Factor XIII â€“ an under diagnosed deficiency â€“ are we using the right assays?. Journal of Thrombosis and Haemostasis, 2010, 8, 2478-2482.	1.9	65
99	Gynecological and obstetrical manifestations of inherited bleeding disorders in women. Journal of Thrombosis and Haemostasis, 2011, 9, 236-245.	1.9	64
100	Genotype-phenotype correlation in combined deficiency of factor V and factor VIII. Blood, 2008, 111, 5592-5600.	0.6	63
101	Adherence to antithrombotic therapy guidelines improves mortality among elderly patients with atrial fibrillation: insights from the REPOSI study. Clinical Research in Cardiology, 2016, 105, 912-920.	1.5	63
102	The effect of emicizumab prophylaxis on healthâ€related outcomes in persons with haemophilia A with inhibitors: HAVEN 1 Study. Haemophilia, 2019, 25, 33-44.	1.0	63
103	Molecular Characterization of a Multiethnic Group of 21 Patients with Type 3 von Willebrand Disease. Thrombosis and Haemostasis, 2000, 84, 536-540.	1.8	61
104	Gene mutations and three-dimensional structural analysis in 13 families with severe factor X deficiency. British Journal of Haematology, 2002, 117, 685-692.	1.2	61
105	Future of coagulation factor replacement therapy. Journal of Thrombosis and Haemostasis, 2013, 11, 84-98.	1.9	61
106	A comparative evaluation of a new automated assay for von Willebrand factor activity. Haemophilia, 2013, 19, 338-342.	1.0	61
107	A fatal case of COVID-19 pneumonia occurring in a patient with severe acute ulcerative colitis. Gut, 2020, 69, 1148-1149.	6.1	60
108	The thrombogram in rare inherited coagulation disorders: its relation to clinical bleeding. Thrombosis and Haemostasis, 2002, 88, 576-82.	1.8	60

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109	A new hemophilia carrier nomenclature to define hemophilia in women and girls: Communication from the SSC of the ISTH. Journal of Thrombosis and Haemostasis, 2021, 19, 1883-1887.	1.9	59
110	Rare bleeding disorders. Haemophilia, 2012, 18, 148-153.	1.0	58
111	Rare Bleeding Disorders: Worldwide Efforts for Classification, Diagnosis, and Management. Seminars in Thrombosis and Hemostasis, 2013, 39, 579-584.	1.5	58
112	Variants of OCTN1-2 cation transporter genes are associated with both Crohn's disease and ulcerative colitis. Alimentary Pharmacology and Therapeutics, 2006, 23, 497-506.	1.9	57
113	Second international collaborative study evaluating performance characteristics of methods measuring the von Willebrand factor cleaving protease (ADAMTS-13). Journal of Thrombosis and Haemostasis, 2008, 6, 1534-1541.	1.9	57
114	Clinical advances in gene therapy updates on clinical trials of gene therapy in haemophilia. Haemophilia, 2019, 25, 738-746.	1.0	57
115	Formation of methionine sulfoxide by peroxynitrite at position 1606 of von Willebrand factor inhibits its cleavage by ADAMTS-13: A new prothrombotic mechanism in diseases associated with oxidative stress. Free Radical Biology and Medicine, 2010, 48, 446-456.	1.3	56
116	TTP and ADAMTS13: When Is Testing Appropriate?. Hematology American Society of Hematology Education Program, 2007, 2007, 121-126.	0.9	54
117	Combined FV and FVIII deficiency. Haemophilia, 2008, 14, 1201-1208.	1.0	54
118	Disorders of hemostasis and excessive menstrual bleeding: prevalence and clinical impact. Fertility and Sterility, 2005, 84, 1338-1344.	0.5	53
119	Treatment of rare factor deficiencies in 2016. Hematology American Society of Hematology Education Program, 2016, 2016, 663-669.	0.9	53
120	The thrombospondin-1 N700S polymorphism is associated with early myocardial infarction without altering von Willebrand factor multimer size. Blood, 2006, 108, 1280-1283.	0.6	52
121	Nextâ€generation sequencing study finds an excess of rare, coding singleâ€nucleotide variants of ADAMTS13 in patients with deep vein thrombosis. Journal of Thrombosis and Haemostasis, 2013, 11, 1228-1239.	1.9	52
122	Plasma ADAMTSâ€13 levels and the risk of myocardial infarction: an individual patient data metaâ€analysis. Journal of Thrombosis and Haemostasis, 2015, 13, 1396-1404.	1.9	52
123	Mortality rate and risk factors for gastrointestinal bleeding in elderly patients. European Journal of Internal Medicine, 2019, 61, 54-61.	1.0	52
124	Pulmonary embolism in a young pregnant woman with COVID-19. Thrombosis Research, 2020, 191, 36-37.	0.8	52
125	Principles of treatment and update of recommendations for the management of haemophilia and congenital bleeding disorders in Italy. Blood Transfusion, 2014, 12, 575-98.	0.3	52
126	Prospective study on the behaviour of the metalloprotease ADAMTS13 and of von Willebrand factor after bone marrow transplantation. British Journal of Haematology, 2006, 134, 187-195.	1.2	51

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127	The association of factor V Leiden with myocardial infarction is replicated in 1880 patients with premature disease. Journal of Thrombosis and Haemostasis, 2010, 8, 2116-2121.	1.9	50
128	Combined Factor V and Factor VIII Deficiency. Seminars in Thrombosis and Hemostasis, 2009, 35, 390-399.	1.5	49
129	Hypercoagulability Is a Stronger Risk Factor for Ischaemic Stroke than for Myocardial Infarction: A Systematic Review. PLoS ONE, 2015, 10, e0133523.	1.1	49
130	Prothrombin 20210G>A is an ancestral prothrombotic mutation that occurred in whites approximately 24 000 years ago. Blood, 2006, 107, 4666-4668.	0.6	48
131	Genetic sequence analysis of inherited bleeding diseases. Blood, 2013, 122, 3423-3431.	0.6	48
132	Acquired inhibitors of clotting factors: AICE recommendations for diagnosis and management. Blood Transfusion, 2015, 13, 498-513.	0.3	48
133	Phenotype and genotype report on homozygous and heterozygous patients with congenital factor X deficiency. Haematologica, 2008, 93, 934-938.	1.7	47
134	Strong association of the APOA5-1131T>C gene variant and early-onset acute myocardial infarction. Atherosclerosis, 2011, 214, 397-403.	0.4	47
135	Factor <scp>XIII</scp> deficiency diagnosis: Challenges and tools. International Journal of Laboratory Hematology, 2018, 40, 3-11.	0.7	47
136	Short-term Exposure to High Altitude Causes Coagulation Activation and Inhibits Fibrinolysis. Thrombosis and Haemostasis, 2002, 87, 342-343.	1.8	46
137	Different clinical severity of first episodes and recurrences of thrombotic thrombocytopenic purpura. British Journal of Haematology, 2010, 151, 488-494.	1.2	46
138	Phase 3 study of recombinant von Willebrand factor in patients with severe von Willebrand disease who are undergoing elective surgery. Journal of Thrombosis and Haemostasis, 2019, 17, 52-62.	1.9	46
139	Detailed stratified GWAS analysis for severe COVID-19 in four European populations. Human Molecular Genetics, 2022, 31, 3945-3966.	1.4	46
140	Phenotype-genotype characterization of 10 families with severe a subunit factor XIII deficiency. Human Mutation, 2004, 23, 98-98.	1.1	45
141	Polypharmacy in older people: lessons from 10Âyears of experience with the REPOSIÂregister. Internal and Emergency Medicine, 2018, 13, 1191-1200.	1.0	45
142	Efficacy and safety of openâ€label caplacizumab in patients with exacerbations of acquired thrombotic thrombocytopenic purpura in the HERCULES study. Journal of Thrombosis and Haemostasis, 2020, 18, 479-484.	1.9	45
143	Genetic risk stratification to reduce inhibitor development in the early treatment of hemophilia A: a SIPPET analysis. Blood, 2017, 130, 1757-1759.	0.6	44
144	Chromosome 3 cluster rs11385942 variant links complement activation with severe COVID-19. Journal of Autoimmunity, 2021, 117, 102595.	3.0	44

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145	Role of Chloride Ions in Modulation of the Interaction between von Willebrand Factor and ADAMTS-13. Journal of Biological Chemistry, 2005, 280, 23295-23302.	1.6	43
146	Gynaecological and obstetrical problems in women with different bleeding disorders. Haemophilia, 2009, 15, 1291-1299.	1.0	43
147	Ndufc2 Gene Inhibition Is Associated With Mitochondrial Dysfunction and Increased Stroke Susceptibility in an Animal Model of Complex Human Disease. Journal of the American Heart Association, 2016, 5, .	1.6	43
148	Platelet to Lymphocyte Ratio and Neutrophil to Lymphocyte Ratio as Risk Factors for Venous Thrombosis. Clinical and Applied Thrombosis/Hemostasis, 2018, 24, 808-814.	0.7	43
149	Molecular analysis of the ERGIC-53 gene in 35 families with combined factor V-factor VIII deficiency. Blood, 1999, 93, 2253-60.	0.6	43
150	Relatively Poor Performance of Clinical Laboratories for DNA Analyses in the Detection of Two Thrombophilic Mutations â€“ A Cause for Concern. Thrombosis and Haemostasis, 2002, 88, 690-691.	1.8	41
151	Arg2074Cys missense mutation in the C2 domain of factor V causing moderately severe factor V deficiency: molecular characterization by expression of the recombinant protein. Blood, 2003, 101, 173-177.	0.6	41
152	Effects of PCSK9 genetic variants on plasma LDL cholesterol levels and risk of premature myocardial infarction in the Italian population. Journal of Lipid Research, 2010, 51, 3342-3349.	2.0	41
153	Defining Aging Phenotypes and Related Outcomes: Clues to Recognize Frailty in Hospitalized Older Patients. Journals of Gerontology - Series A Biological Sciences and Medical Sciences, 2016, 72, glw188.	1.7	41
154	Exploring the global landscape of genetic variation in coagulation factor XI deficiency. Blood, 2017, 130, e1-e6.	0.6	41
155	Nonneutralizing antibodies against factor VIII and risk of inhibitor development in severe hemophilia A. Blood, 2017, 129, 1245-1250.	0.6	41
156	Clinical and Laboratory Features of Patients with Acquired Thrombotic Thrombocytopenic Purpura: Fourteen Years of the Milan TTP Registry. Thrombosis and Haemostasis, 2019, 119, 695-704.	1.8	41
157	Management of pregnancy and delivery in women with inherited bleeding disorders. Seminars in Fetal and Neonatal Medicine, 2011, 16, 311-317.	1.1	40
158	Factor XI deficiency in Iranians: its clinical manifestations in comparison with those of classic hemophilia. Haematologica, 2002, 87, 512-4.	1.7	40
159	Mechanisms of the interaction between twoADAMTS13 gene mutations leading to severe deficiency of enzymatic activity. Human Mutation, 2006, 27, 330-336.	1.1	39
160	Discrepancies between ADAMTS13 activity assays in patients with thrombotic microangiopathies. Thrombosis and Haemostasis, 2013, 109, 488-496.	1.8	39
161	A twoâ€centre comparative evaluation of new automated assays for von Willebrand factor ristocetin cofactor activity and antigen. Haemophilia, 2014, 20, 147-153.	1.0	39
162	Timing and severity of inhibitor development in recombinant versus plasmaâ€derived factor VIII concentrates: a SIPPET analysis. Journal of Thrombosis and Haemostasis, 2018, 16, 39-43.	1.9	39

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163	Caplacizumab prevents refractoriness and mortality in acquired thrombotic thrombocytopenic purpura: integrated analysis. Blood Advances, 2021, 5, 2137-2141.	2.5	39
164	Clinical and molecular characterization of 6 patients affected by severe deficiency of coagulation factor V: broadening of the mutational spectrum of factor V gene and in vitro analysis of the newly identified missense mutations. Blood, 2003, 102, 3210-3216.	0.6	38
165	Effects ofCYP2C9andVKORC1on INR variations and dose requirements during initial phase of anticoagulant therapy. Pharmacogenomics, 2008, 9, 1237-1250.	0.6	38
166	How the Direct Oral Anticoagulant Apixaban Affects Thrombin Generation Parameters. Thrombosis Research, 2015, 135, 1186-1190.	0.8	38
167	Pulmonary immuno-thrombosis in COVID-19 ARDS pathogenesis. Intensive Care Medicine, 2021, 47, 899-902.	3.9	38
168	Congenital afibrinogenemia: first identification of splicing mutations in the fibrinogen BÎ²-chain gene causing activation of cryptic splice sites. Blood, 2002, 100, 4478-4484.	0.6	37
169	The genetics of the alternative pathway of complement in the pathogenesis of HELLP syndrome. Journal of Maternal-Fetal and Neonatal Medicine, 2012, 25, 2322-2325.	0.7	37
170	Evaluation of a New, Rapid, Fully Automated Assay for the Measurement of ADAMTS13 Activity. Thrombosis and Haemostasis, 2019, 119, 1767-1772.	1.8	37
171	Comparison of adverse drug reactions among four COVIDâ€19 vaccines in Europe using the EudraVigilance database: Thrombosis at unusual sites. Journal of Thrombosis and Haemostasis, 2021, 19, 2554-2558.	1.9	37
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