Bernhard LÃ¤mmle

List of Publications by Year
in descending order

277 papers	17,992 citations	18482 62 h-index	14208 128 g-index
282 all docs	282 docs citations	282 times ranked	9516 citing authors

#	Article	IF	Citations
1	von Willebrand Factorâ€“Cleaving Protease in Thrombotic Thrombocytopenic Purpura and the Hemolyticâ€“Uremic Syndrome. New England Journal of Medicine, 1998, 339, 1578-1584.	27.0	1,717
2	Partial purification and characterization of a protease from human plasma cleaving von Willebrand factor to fragments produced by in vivo proteolysis. Blood, 1996, 87, 4223-4234.	1.4	858
3	Elevated nucleosome levels in systemic inflammation and sepsis*. Critical Care Medicine, 2003, 31, 1947-1951.	0.9	715
4	ADAMTS13 activity in thrombotic thrombocytopenic purpuraâ€“hemolytic uremic syndrome: relation to presenting features and clinical outcomes in a prospective cohort of 142 patients. Blood, 2003, 102, 60-68.	1.4	649
5	C1-inhibitor in patients with severe sepsis and septic shock: Beneficial effect on renal dysfunction. Critical Care Medicine, 2002, 30, 1722-1728.	0.9	634
6	Deficient Activity of von Willebrand Factorâ€“Cleaving Protease in Chronic Relapsing Thrombotic Thrombocytopenic Purpura. Blood, 1997, 89, 3097-3103.	1.4	536
7	Survival and relapse in patients with thrombotic thrombocytopenic purpura. Blood, 2010, 115, 1500-1511.	1.4	477
8	Partial amino acid sequence of purified von Willebrand factorâ€“cleaving protease. Blood, 2001, 98, 1654-1661.	1.4	375
9	von Willebrand factorâ€“mediated platelet adhesion is critical for deep vein thrombosis in mouse models. Blood, 2011, 117, 1400-1407.	1.4	369
10	Consensus on the standardization of terminology in thrombotic thrombocytopenic purpura and related thrombotic microangiopathies. Journal of Thrombosis and Haemostasis, 2017, 15, 312-322.	3.8	362
11	Severe COVID-19 infection associated with endothelial activation. Thrombosis Research, 2020, 190, 62.	1.7	358
12	Von Willebrand factorâ€“cleaving protease (ADAMTS13) in thrombocytopenic disorders: a severely deficient activity is specific for thrombotic thrombocytopenic purpura. Blood, 2002, 100, 710-713.	1.4	343
13	Diagnostic criteria for hematopoietic stem cell transplant-associated microangiopathy: results of a consensus process by an International Working Group. Haematologica, 2007, 92, 95-100.	3.5	341
14	Acquired Deficiency of von Willebrand Factor-Cleaving Protease in a Patient With Thrombotic Thrombocytopenic Purpura. Blood, 1998, 91, 2839-2846.	1.4	324
15	Assay of von Willebrand Factor (vWF)-cleaving Protease Based on Decreased Collagen Binding Affinity of Degraded vWF. Thrombosis and Haemostasis, 1999, 82, 1386-1389.	3.4	310
16	The incidence of thrombotic thrombocytopenic purpura-hemolytic uremic syndrome: all patients, idiopathic patients, and patients with severe ADAMTS-13 deficiency. Journal of Thrombosis and Haemostasis, 2005, 3, 1432-1436.	3.8	305
17	ADAMTS13 autoantibodies in patients with thrombotic microangiopathies and other immunomediated diseases. Blood, 2005, 106, 1262-1267.	1.4	275
18	Circulating DNA and myeloperoxidase indicate disease activity in patients with thrombotic microangiopathies. Blood, 2012, 120, 1157-1164.	1.4	249

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19	Thrombotic thrombocytopenic purpura. Nature Reviews Disease Primers, 2017, 3, 17020.	30.5	242
20	Aetiology and pathogenesis of thrombotic thrombocytopenic purpura and haemolytic uraemic syndrome: the role of von Willebrand factor-cleaving protease. Best Practice and Research in Clinical Haematology, 2001, 14, 437-454.	1.7	217
21	Cloning, expression, and functional characterization of the von Willebrand factorâ€“cleaving protease (ADAMTS13). Blood, 2002, 100, 3626-3632.	1.4	216
22	Epitope mapping of ADAMTS13 autoantibodies in acquired thrombotic thrombocytopenic purpura. Blood, 2004, 103, 4514-4519.	1.4	204
23	Recovery and Half-Life of von Willebrand Factor-Cleaving Protease after Plasma Therapy in Patients with Thrombotic Thrombocytopenic Purpura. Thrombosis and Haemostasis, 1999, 81, 8-13.	3.4	203
24	Association of two silent polymorphisms of platelet glycoprotein Ia/IIa receptor with risk of myocardial infarction: a case-control study. Lancet, The, 1999, 353, 351-354.	13.7	200
25	Multiple major morbidities and increased mortality during long-term follow-up after recovery from thrombotic thrombocytopenic purpura. Blood, 2013, 122, 2023-2029.	1.4	161
26	Thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2005, 3, 1663-1675.	3.8	159
27	Fibrin glue in surgery: frequent development of inhibitors of bovine thrombin and human factor V. British Journal of Haematology, 1993, 85, 528-532.	2.5	157
28	ADAMTSâ€13, von Willebrand factor and related parameters in severe sepsis and septic shock. Journal of Thrombosis and Haemostasis, 2007, 5, 2284-2290.	3.8	153
29	Fatal congenital thrombotic thrombocytopenic purpura with apparent ADAMTS13 inhibitor: in vitro inhibition of ADAMTS13 activity by hemoglobin. Blood, 2005, 105, 542-544.	1.4	152
30	Thromboembolism and Bleeding Tendency in Congenital Factor XII DeficienCy - A Study on 74 Subjects from 14 Swiss Families. Thrombosis and Haemostasis, 1991, 65, 117-121.	3.4	146
31	Gut microbiota regulate hepatic von Willebrand factor synthesis and arterial thrombus formation via Toll-like receptor-2. Blood, 2017, 130, 542-553.	1.4	119
32	Impaired DNase1â€mediated degradation of neutrophil extracellular traps is associated with acute thrombotic microangiopathies. Journal of Thrombosis and Haemostasis, 2015, 13, 732-742.	3.8	109
33	Redefining outcomes in immune TTP: an international working group consensus report. Blood, 2021, 137, 1855-1861.	1.4	103
34	Clinical outcomes after platelet transfusions in patients with thrombotic thrombocytopenic purpura. Transfusion, 2009, 49, 873-887.	1.6	99
35	Plasma DNA is elevated in patients with deep vein thrombosis. Journal of Vascular Surgery: Venous and Lymphatic Disorders, 2013, 1, 341-348.e1.	1.6	99
36	The International Hereditary Thrombotic Thrombocytopenic Purpura Registry: key findings at enrollment until 2017. Haematologica, 2019, 104, 2107-2115.	3.5	99

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37	Coagulation Factors II, V, VII, and X, Prothrombin Gene 20210Gât'A Transition, and Factor V Leiden in Coronary Artery Disease. Arteriosclerosis, Thrombosis, and Vascular Biology, 1999, 19, 1020-1025.	2.4	97
38	Ten years of prophylactic treatment with fresh-frozen plasma in a child with chronic relapsing thrombotic thrombocytopenic purpura as a result of a congenital deficiency of von Willebrand factor-cleaving protease. British Journal of Haematology, 2001, 113, 649-651.	2.5	95
39	Measurement of von Willebrand factor-cleaving protease (ADAMTS-13) activity in plasma: a multicenter comparison of different assay methods. Journal of Thrombosis and Haemostasis, 2003, 1, 1882-1887.	3.8	95
40	Thrombotic Thrombocytopenic Purpura: Pathophysiology, Diagnosis, and Management. Journal of Clinical Medicine, 2021, 10, 536.	2.4	94
41	ADAMTS13 gene defects in two brothers with constitutional thrombotic thrombocytopenic purpura and normalization of von Willebrand factor-cleaving protease activity by recombinant human ADAMTS13. British Journal of Haematology, 2003, 120, 821-824.	2.5	89
42	Measurement of ADAMTS-13 activity in plasma by the FRETS-VWF73 assay: comparison with other assay methods. Journal of Thrombosis and Haemostasis, 2006, 4, 1146-1148.	3.8	89
43	Accuracy of d-dimer/fibrinogen ratio to predict pulmonary embolism: a prospective diagnostic study. Journal of Thrombosis and Haemostasis, 2003, 1, 708-713.	3.8	82
44	Stability of coagulation assays performed in plasma from citrated whole blood transported at ambient temperature. Thrombosis and Haemostasis, 2008, 99, 416-426.	3.4	81
45	Von Willebrand factor-cleaving protease (ADAMTS-13) activity determination in the diagnosis of thrombotic microangiopathies: the Swiss experience. Seminars in Hematology, 2004, 41, 75-82.	3.4	79
46	Triplet structure of von Willebrand factor reflects proteolytic degradation of high molecular weight multimers.. Proceedings of the National Academy of Sciences of the United States of America, 1993, 90, 7503-7507.	7.1	78
47	Predictors and Causes of Long-Term Mortality in Elderly Patients with Acute Venous Thromboembolism: A Prospective Cohort Study. American Journal of Medicine, 2017, 130, 198-206.	1.5	78
48	Dural puncture and activated protein C resistance: risk factors for cerebral venous sinus thrombosis. Journal of Neurology, Neurosurgery and Psychiatry, 1997, 63, 351-356.	1.9	77
49	Pancreatitis preceding acute episodes of thrombotic thrombocytopenic purpura-hemolytic uremic syndrome: report of five patients with a systematic review of published reports. Haematologica, 2007, 92, 936-943.	3.5	75
50	Different disparities of gender and race among the thrombotic thrombocytopenic purpura and hemolyticâ€uremic syndromes. American Journal of Hematology, 2010, 85, 844-847.	4.1	75
51	Protein C Replacement in Severe Meningococcemia: Rationale and Clinical Experience. Clinical Infectious Diseases, 2001, 32, 1338-1346.	5.8	74
52	A common origin of the 4143insA ADAMTS13 mutation. Thrombosis and Haemostasis, 2006, 96, 3-6.	3.4	74
53	Cognitive deficits after recovery from thrombotic thrombocytopenic purpura. Transfusion, 2009, 49, 1092-1101.	1.6	73
54	Rapid determination of anti-heparin/platelet factor 4 antibody titers in the diagnosis of heparin-induced thrombocytopenia. American Journal of Medicine, 2003, 114, 528-536.	1.5	72

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55	Hyperbilirubinemia interferes with ADAMTS-13 activity measurement by FRETS-VWF73 assay: diagnostic relevance in patients suffering from acute thrombotic microangiopathies. Journal of Thrombosis and Haemostasis, 2007, 5, 866-867.	3.8	72
56	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.	4.1	71
57	Microangiopathic haemolytic anaemia in metastasizing malignant tumours is not associated with a severe deficiency of the von Willebrand factor-cleaving protease. British Journal of Haematology, 2001, 113, 100-102.	2.5	70
58	Evidence for a role of anti-ADAMTS13 autoantibodies despite normal ADAMTS13 activity in recurrent thrombotic thrombocytopenic purpura. Haematologica, 2012, 97, 297-303.	3.5	69
59	SEVERE OSTEOPOROSIS DUE TO SYSTEMIC MAST CELL DISEASE: SUCCESSFUL TREATMENT WITH INTERFERON ALPHA-2B. Rheumatology, 1996, 35, 898-900.	1.9	66
60	ADAMTS13 activity, von Willebrand factor, factor VIII and D-dimers in COVID-19 inpatients. Thrombosis Research, 2020, 192, 174-175.	1.7	66
61	Plasma Prekallikrein, Factor XII, Antithrombin III, C1â”-Inhibitor and Î±2-Macroglobulin in Critically Ill Patients with Suspected Disseminated Intravascular Coagulation (DIC). American Journal of Clinical Pathology, 1984, 82, 396-404.	0.7	64
62	Von Willebrand Factor-cleaving Protease in Childhood Diarrhoea-associated Haemolytic Uraemic Syndrome. Thrombosis and Haemostasis, 2001, 85, 975-978.	3.4	64
63	Polypharmacy is Associated with an Increased Risk of Bleeding in Elderly Patients with Venous Thromboembolism. Journal of General Internal Medicine, 2015, 30, 17-24.	2.6	64
64	Validation of the ISTH/SSC bleeding assessment tool for inherited platelet disorders: A communication from the Platelet Physiology SSC. Journal of Thrombosis and Haemostasis, 2020, 18, 732-739.	3.8	64
65	Thrombo-Inflammation in Cardiovascular Disease: An Expert Consensus Document from the Third Maastricht Consensus Conference on Thrombosis. Thrombosis and Haemostasis, 2020, 120, 538-564.	3.4	64
66	Rapid exclusion or confirmation of heparin-induced thrombocytopenia: a single-center experience with 1,291 patients. Haematologica, 2012, 97, 89-97.	3.5	63
67	Decreasing frequency of plasma exchange complications in patients treated for thrombotic thrombocytopenic purpuraâ€hemolytic uremic syndrome, 1996 to 2011 (CME). Transfusion, 2012, 52, 2525-2532.	1.6	63
68	The Swiss cohort of elderly patients with venous thromboembolism (SWITCO65+): rationale and methodology. Journal of Thrombosis and Thrombolysis, 2013, 36, 475-483.	2.1	63
69	Immunoblotting studies of the molecular forms of protein C in plasma. Thrombosis Research, 1988, 52, 33-43.	1.7	61
70	Elevated levels of plasma prekallikrein, high molecular weight kininogen and factor XI in coronary heart disease. Atherosclerosis, 2002, 161, 261-267.	0.8	61
71	Familial acquired thrombotic thrombocytopenic purpura: ADAMTS13 inhibitory autoantibodies in identical twins. Blood, 2004, 103, 4195-4197.	1.4	61
72	Pregnancy outcomes following recovery from acquired thrombotic thrombocytopenic purpura. Blood, 2014, 123, 1674-1680.	1.4	61

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73	Prekallikrein deficiency. Thrombosis Research, 2002, 105, 463-470.	1.7	60
74	Protein Z in ischaemic stroke. British Journal of Haematology, 2001, 114, 169-173.	2.5	59
75	D-Dimers Predict Stroke Subtype when Assessed Early. Cerebrovascular Diseases, 2010, 29, 82-86.	1.7	58
76	Platelets: Thrombotic Thrombocytopenic Purpura. Hematology American Society of Hematology Education Program, 2002, 2002, 315-334.	2.5	58
77	Role of ADAMTS13 in the pathogenesis, diagnosis, and treatment of thrombotic thrombocytopenic purpura. Hematology American Society of Hematology Education Program, 2012, 2012, 610-616.	2.5	58
78	Treatment of thrombotic thrombocytopenic purpura. Vox Sanguinis, 2006, 90, 245-254.	1.5	57
79	ADAMTS13 activity in sickle cell disease. American Journal of Hematology, 2006, 81, 492-498.	4.1	54
80	Titre of anti-heparin/PF4-antibodies and extent of in vivo activation of the coagulation and fibrinolytic systems. Thrombosis and Haemostasis, 2004, 91, 276-282.	3.4	50
81	Sporadic bloody diarrhoeaâ€associated thrombotic thrombocytopenic purpuraâ€haemolytic uraemic syndrome: an adult and paediatric comparison. British Journal of Haematology, 2008, 141, 696-707.	2.5	50
82	Effect of low-molecular weight dextran sulfate on coagulation and platelet function tests. Thrombosis Research, 2002, 105, 441-446.	1.7	49
83	High prevalence of hereditary thrombotic thrombocytopenic purpura in central Norway: from clinical observation to evidence. Journal of Thrombosis and Haemostasis, 2016, 14, 73-82.	3.8	49
84	Frequency and Significance of HIV Infection among Patients Diagnosed with Thrombotic Thrombocytopenic Purpura. Clinical Infectious Diseases, 2009, 48, 1129-1137.	5.8	48
85	Splenectomy in relapsing and plasma-refractory acquired thrombotic thrombocytopenic purpura. Haematologica, 2004, 89, 320-4.	3.5	46
86	SimpliRED D-dimer Assay: Comparability of Capillary and Citrated Venous Whole Blood, Between-assay Variability, and Performance of the Test for Exclusion of Deep Vein Thrombosis in Symptomatic Outpatients. Thrombosis and Haemostasis, 1998, 79, 1217-1219.	3.4	44
87	Acquired thrombotic thrombocytopenic purpura: ADAMTS13 activity, anti-ADAMTS13 autoantibodies and risk of recurrent disease. Haematologica, 2008, 93, 172-177.	3.5	44
88	Rapid D-dimer testing and pre-test clinical probability in the exclusion of deep venous thrombosis in symptomatic outpatients. Blood Coagulation and Fibrinolysis, 2001, 12, 165-170.	1.0	43
89	Dosing lepirudin in patients with heparin-induced thrombocytopenia and normal or impaired renal function: a single-center experience with 68 patients. Blood, 2009, 113, 2402-2409.	1.4	43
90	Hereditary thrombotic thrombocytopenic purpura and the hereditary TTP registry. Hamostaseologie, 2013, 33, 138-143.	1.9	43

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91	16 Deficiency of von Willebrand factor-cleaving protease in familial and acquired thrombotic thrombocytopenic purpura. Best Practice and Research: Clinical Haematology, 1998, 11, 509-514.	1.1	41
92	Protein Z in healthy human individuals and in patients with a bleeding tendency. British Journal of Haematology, 1998, 102, 1219-1223.	2.5	41
93	The Oklahoma Thrombotic Thrombocytopenic Purpuraâ€“Hemolytic Uremic Syndrome Registry: the Swiss connection. European Journal of Haematology, 2008, 80, 277-286.	2.2	40
94	The splenic autoimmune response to ADAMTS13 in thrombotic thrombocytopenic purpura contains recurrent antigen-binding CDR3 motifs. Blood, 2014, 124, 3469-3479.	1.4	40
95	Depression and cognitive deficits as longâ€term consequences of thrombotic thrombocytopenic purpura. Transfusion, 2017, 57, 1152-1162.	1.6	40
96	Plasma therapy in thrombotic thrombocytopenic purpura: review of the literature and the Bern experience in a subgroup of patients with severe acquired ADAMTS-13 deficiency. Seminars in Hematology, 2004, 41, 48-59.	3.4	39
97	Monitoring of Heparin Treatment. Comparison between Thrombin Time, Activated Partial Thromboplastin Time, and Plasma Heparin Concentration, and Analysis of the Behavior of Antithrombin III. American Journal of Clinical Pathology, 1980, 74, 68-73.	0.7	38
98	Mycobacterium genavense infection in a patient with long-standing chronic lymphocytic leukaemia. Journal of Internal Medicine, 2000, 248, 343-348.	6.0	38
99	Factor XII clotting activity and antigen levels in patients with thromboembolic disease. Blood Coagulation and Fibrinolysis, 1992, 3, 555-562.	1.0	35
100	No association of APC resistance with myocardial infarction. Blood Coagulation and Fibrinolysis, 1995, 6, 456-459.	1.0	35
101	Haemolyticâ€uraemic syndrome and thrombotic thrombocytopenic purpuraâ€”new insights into underlying biochemical mechanisms. Nephrology Dialysis Transplantation, 2000, 15, 1112-1114.	0.7	35
102	Thrombotic microangiopathic syndromes associated with drugs, HIV infection, hematopoietic stem cell transplantation and cancer. Presse Medicale, 2012, 41, e177-e188.	1.9	35
103	Open ADAMTS13, induced by antibodies, is a biomarker for subclinical immune-mediated thrombotic thrombocytopenic purpura. Blood, 2020, 136, 353-361.	1.4	35
104	Detection and Quantitation of Cleaved and Uncleaved High Molecular Weight Kininogen in Plasma by Ligand Blotting with Radiolabeled Plasma Prekallikrein or Factor XI. Thrombosis and Haemostasis, 1988, 59, 151-161.	3.4	33
105	Heparin-dependent in vitro aggregation of normal platelets by plasma of a patient with heparin-induced skin necrosis: specific diagnostic test for a rare side effect. American Journal of Medicine, 1988, 85, 721-724.	1.5	32
106	IN VITRO EVALUATION OF THE EFFICACY AND BIOCOMPATIBILITY OF NEW, SYNTHETIC ABO IMMUNOABSORBENTS. Transplantation, 1995, 60, 425-429.	1.0	32
107	Current insights into thrombotic microangiopathies: Thrombotic thrombocytopenic purpura and pregnancy. Thrombosis Research, 2015, 135, S30-S33.	1.7	32
108	Thromboembolism in patients with congenital afibrinogenaemia. Thrombosis and Haemostasis, 2016, 116, 722-732.	3.4	32

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109	Binding of Î±-thrombin to fibrin depends on the quality of the fibrin network. Biochemical Journal, 1994, 298, 157-163.	3.7	31
110	Relapse Rate in Survivors of Acute Autoimmune Thrombotic Thrombocytopenic Purpura Treated with or without Rituximab. Thrombosis and Haemostasis, 2018, 118, 1743-1751.	3.4	31
111	Annual incidence and severity of acute episodes in hereditary thrombotic thrombocytopenic purpura. Blood, 2021, 137, 3563-3575.	1.4	31
112	Enhanced specificity of immunoblotting using radiolabeled antigen overlay: Studies of blood coagulation factor XII and prekallikrein in plasma. Analytical Biochemistry, 1986, 156, 118-125.	2.4	30
113	Circulating extracellular DNA is an independent predictor of mortality in elderly patients with venous thromboembolism. PLoS ONE, 2018, 13, e0191150.	2.5	30
114	Use of the pentasaccharide fondaparinux as an anticoagulant during haemodialysis. Thrombosis and Haemostasis, 2007, 98, 1200-1207.	3.4	29
115	Variability of antiâ€PF4/heparin antibody results obtained by the rapid testing system IDâ€H/PF4â€PaGIA. Journal of Thrombosis and Haemostasis, 2009, 7, 1649-1655.	3.8	29
116	Prospective comparison of clinical prognostic scores in elder patients with a pulmonary embolism. Journal of Thrombosis and Haemostasis, 2012, 10, 2270-2276.	3.8	29
117	Evaluation of a Platelet Function Analyser (PFA-100) in patients with a bleeding tendency. Swiss Medical Weekly, 2002, 132, 443-8.	1.6	29
118	Von Willebrand Factor in Thrombotic Thrombocytopenic Purpura. Thrombosis and Haemostasis, 1999, 82, 592-600.	3.4	28
119	New strategies in diagnosis and treatment of thrombotic thrombocytopenic purpura: case report and review. European Journal of Pediatrics, 1999, 158, 883-887.	2.7	28
120	Plasma Protein C Inhibitor Is Elevated in Survivors of Myocardial Infarction. Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 114-118.	2.4	28
121	The novel acceptor splice site mutation 11396(G->A) in the factor XII gene causes a truncated transcript in cross-reacting material negative patients. Human Molecular Genetics, 1995, 4, 1235-1237.	2.9	27
122	Blood group O and black race are independent risk factors for thrombotic thrombocytopenic purpura associated with severe ADAMTS13 deficiency. Transfusion, 2011, 51, 2237-2243.	1.6	27
123	IFNÎ± treatment in systemic mastocytosis. Annals of Hematology, 1999, 78, 483-484.	1.8	26
124	Genetic Predisposition to Bleeding during Oral Anticoagulant Therapy: Evidence for Common Founder Mutations (FIXVal-10 and FIXThr-10) and an Independent CpG Hotspot Mutation (FIXThr-10). Thrombosis and Haemostasis, 2001, 85, 454-457.	3.4	25
125	Assays of von Willebrand Factor- Cleaving Protease: A Test for Diagnosis of Familial and Acquired Thrombotic Thrombocytopenic Purpura. Seminars in Thrombosis and Hemostasis, 2002, 28, 167-172.	2.7	25
126	Factor XIII in severe sepsis and septic shock. Thrombosis Research, 2007, 119, 311-318.	1.7	25

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127	Acquired deficiency of von Willebrand factorâ€cleaving protease in a patient suffering from acute systemic lupus erythematosus. Rheumatology, 2001, 40, 940-942.	1.9	24
128	Usefulness of the D-dimer/fibrinogen ratio to predict deep venous thrombosis. Journal of Thrombosis and Haemostasis, 2005, 3, 385-387.	3.8	24
129	Low molecular weight heparinâ€induced thrombocytopenia and skin necrosis distant from injection sites. European Journal of Haematology, 1994, 53, 61-63.	2.2	24
130	Quantitative immunoblotting assay of blood coagulation factor XII. Thrombosis Research, 1986, 41, 747-759.	1.7	23
131	A first case of congenital TTP on the African continent due to a new homozygous mutation in the catalytic domain of ADAMTS13. Annals of Hematology, 2008, 87, 663-666.	1.8	23
132	The impact of congenital thrombotic thrombocytopenic purpura on pregnancy complications. Thrombosis and Haemostasis, 2014, 111, 1180-1183.	3.4	23
133	Severe plasma prekallikrein deficiency: Clinical characteristics, novel KLKB1 mutations, and estimated prevalence. Journal of Thrombosis and Haemostasis, 2020, 18, 1598-1617.	3.8	23
134	Purified Human Plasma Kallikrein Does Not Stimulate but Primes Neutrophils for Superoxide Production. Thrombosis and Haemostasis, 1989, 62, 1121-1125.	3.4	23
135	Rituximab for acute plasma-refractory thrombotic thrombocytopenic purpura. A case report and concise review of the literature. Swiss Medical Weekly, 2007, 137, 518-24.	1.6	23
136	Progressive multifocal leukoencephalopathy in common variable immunodeficiency: mitigated course under mirtazapine and mefloquine. Journal of NeuroVirology, 2015, 21, 694-701.	2.1	22
137	Influence of Low Molecular Weight Heparin and Low Molecular Weight Dextran Sulfate on the Inhibition of Coagulation Factor XIa by Serpins. Thrombosis and Haemostasis, 1998, 80, 82-86.	3.4	21
138	Hemophilia A Pseudoaneurysm in a Patient with High Responding Inhibitors Complicating Total Knee Arthroplasty: Embolization: A Cost-Reducing Alternative to Medical Therapy. CardioVascular and Interventional Radiology, 2006, 29, 1132-1135.	2.0	21
139	The prothrombin time/international normalized ratio (PT/INR) Line: derivation of local INR with commercial thromboplastins and coagulometers â€“ two independent studies. Journal of Thrombosis and Haemostasis, 2011, 9, 140-148.	3.8	21
140	Role of ADAMTS13 in the pathogenesis, diagnosis, and treatment of thrombotic thrombocytopenic purpura. Hematology American Society of Hematology Education Program, 2012, 2012, 610-6.	2.5	21
141	A new substitution, gamma 358 Ser-->Cys, in fibrinogen Milano VII causes defective fibrin polymerization. Blood, 1994, 84, 1874-1880.	1.4	20
142	Bilateral periorbital ecchymoses. Hamostaseologie, 2014, 34, 249-252.	1.9	20
143	Genetic variations in complement factors in patients with congenital thrombotic thrombocytopenic purpura with renal insufficiency. International Journal of Hematology, 2016, 103, 283-291.	1.6	20
144	CapnocytophagaÂ canimorsusSepsis. New England Journal of Medicine, 1998, 339, 1827-1827.	27.0	19

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145	Beta2-Glycoprotein I: Implications for a Regulatory Role in Thrombotic Thrombocytopenic Purpura.. Blood, 2007, 110, 278-278.	1.4	19
146	Contact phase of blood coagulation is not activated in edema of high altitude. Journal of Applied Physiology, 1989, 67, 1336-1340.	2.5	18
147	Performance of a New Fibrin Monomer Assay to Exclude Deep Vein Thrombosis in Symptomatic Outpatients. Thrombosis and Haemostasis, 1999, 81, 50-53.	3.4	18
148	The von Willebrand Factor-Cleaving Protease (ADAMTS-13) and the Diagnosis of Thrombotic Thrombocytopenic Purpura (TTP). Pathophysiology of Haemostasis and Thrombosis: International Journal on Haemostasis and Thrombosis Research, 2003, 33, 417-421.	0.3	18
149	Patent ductus arteriosus generates neonatal hemolytic jaundice with thrombocytopenia in Upshaw-Schulman syndrome. Blood Advances, 2019, 3, 3191-3195.	5.2	18
150	How high is the true fibrinogen content of fibrinogen standards?. Thrombosis Research, 1989, 56, 583-592.	1.7	17
151	Predictors and Outcomes of Recurrent Venous Thromboembolism in Elderly Patients. American Journal of Medicine, 2018, 131, 703.e7-703.e16.	1.5	17
152	Quantitative Immunoblotting of Plasma and Platelet Protein S. Thrombosis and Haemostasis, 1986, 56, 382-386.	3.4	17
153	Association between thyroid dysfunction and venous thromboembolism in the elderly: a prospective cohort study. Journal of Thrombosis and Haemostasis, 2016, 14, 685-694.	3.8	16
154	Derivation and validation of a novel bleeding risk score for elderly patients with venous thromboembolism on extended anticoagulation. Thrombosis and Haemostasis, 2017, 117, 1930-1936.	3.4	16
155	Immunogenic hotspots in the spacer domain of ADAMTS13 in immuneâ€mediated thrombotic thrombocytopenic purpura. Journal of Thrombosis and Haemostasis, 2021, 19, 478-488.	3.8	16
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