Bernhard Lmmle

List of Publications by Citations

274 papers	14,662 citations	54 h-index	116 g-index
282 ext. papers	16,378 ext. citations	6 avg, IF	6.3 L-index

#	Paper	IF	Citations
274	von Willebrand factor-cleaving protease in thrombotic thrombocytopenic purpura and the hemolytic-uremic syndrome. *New England Journal of Medicine*, **1998**, 339, 1578-84	59.2	1449
273	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	2.2	755
272	Elevated nucleosome levels in systemic inflammation and sepsis. *Critical Care Medicine*, **2003**, 31, 1947-51	1.4	677
271	C1-inhibitor in patients with severe sepsis and septic shock: beneficial effect on renal dysfunction. *Critical Care Medicine*, **2002**, 30, 1722-8	1.4	615
270	ADAMTS13 activity in thrombotic thrombocytopenic purpura-hemolytic uremic syndrome: relation to presenting features and clinical outcomes in a prospective cohort of 142 patients. *Blood*, **2003**, 102, 60-8	2.2	552
269	Deficient Activity of von Willebrand Factor□leaving Protease in Chronic Relapsing Thrombotic Thrombocytopenic Purpura. *Blood*, **1997**, 89, 3097-3103	2.2	470
268	Survival and relapse in patients with thrombotic thrombocytopenic purpura. *Blood*, **2010**, 115, 1500-11; quiz 1662	2.2	377
267	Partial amino acid sequence of purified von Willebrand factor-cleaving protease. *Blood*, **2001**, 98, 1654-61	2.2	333
266	Von Willebrand factor-cleaving protease (ADAMTS13) in thrombocytopenic disorders: a severely deficient activity is specific for thrombotic thrombocytopenic purpura. *Blood*, **2002**, 100, 710-3	2.2	302
265	von Willebrand factor-mediated platelet adhesion is critical for deep vein thrombosis in mouse models. *Blood*, **2011**, 117, 1400-7	2.2	294
264	Acquired Deficiency of von Willebrand Factor-Cleaving Protease in a Patient With Thrombotic Thrombocytopenic Purpura. *Blood*, **1998**, 91, 2839-2846	2.2	292
263	Assay of von Willebrand Factor (vWF)-cleaving Protease Based on Decreased Collagen Binding Affinity of Degraded vWF. *Thrombosis and Haemostasis*, **1999**, 82, 1386-1389	7	277
262	Severe COVID-19 infection associated with endothelial activation. *Thrombosis Research*, **2020**, 190, 62	8.2	272
261	Diagnostic criteria for hematopoietic stem cell transplant-associated microangiopathy: results of a consensus process by an International Working Group. *Haematologica*, **2007**, 92, 95-100	6.6	259
260	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, 1432-6	15.4	247
259	Consensus on the standardization of terminology in thrombotic thrombocytopenic purpura and related thrombotic microangiopathies. *Journal of Thrombosis and Haemostasis*, **2017**, 15, 312-322	15.4	236
258	ADAMTS13 autoantibodies in patients with thrombotic microangiopathies and other immunomediated diseases. *Blood*, **2005**, 106, 1262-7	2.2	229

257	Circulating DNA and myeloperoxidase indicate disease activity in patients with thrombotic microangiopathies. *Blood*, **2012**, 120, 1157-64	2.2	198
256	Cloning, expression, and functional characterization of the von Willebrand factor-cleaving protease (ADAMTS13). *Blood*, **2002**, 100, 3626-32	2.2	197
255	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, 437-54	4.2	187
254	Association of two silent polymorphisms of platelet glycoprotein Ia/IIa receptor with risk of myocardial infarction: a case-control study. *Lancet, The*, **1999**, 353, 351-4	40	184
253	Epitope mapping of ADAMTS13 autoantibodies in acquired thrombotic thrombocytopenic purpura. *Blood*, **2004**, 103, 4514-9	2.2	178
252	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	7	161
251	Thrombotic thrombocytopenic purpura. *Nature Reviews Disease Primers*, **2017**, 3, 17020	51.1	139
250	Thrombotic thrombocytopenic purpura. *Journal of Thrombosis and Haemostasis*, **2005**, 3, 1663-75	15.4	138
249	ADAMTS-13, von Willebrand factor and related parameters in severe sepsis and septic shock. *Journal of Thrombosis and Haemostasis*, **2007**, 5, 2284-90	15.4	135
248	Fatal congenital thrombotic thrombocytopenic purpura with apparent ADAMTS13 inhibitor: in vitro inhibition of ADAMTS13 activity by hemoglobin. *Blood*, **2005**, 105, 542-4	2.2	135
247	Fibrin glue in surgery: frequent development of inhibitors of bovine thrombin and human factor V. *British Journal of Haematology*, **1993**, 85, 528-32	4.5	130
246	Multiple major morbidities and increased mortality during long-term follow-up after recovery from thrombotic thrombocytopenic purpura. *Blood*, **2013**, 122, 2023-9; quiz 2142	2.2	112
245	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	7	108
244	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, 1882-7	15.4	84
243	Coagulation factors II, V, VII, and X, prothrombin gene 20210G-->A transition, and factor V Leiden in coronary artery disease: high factor V clotting activity is an independent risk factor for myocardial infarction. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **1999**, 19, 1020-5	9.4	84
242	Clinical outcomes after platelet transfusions in patients with thrombotic thrombocytopenic purpura. *Transfusion*, **2009**, 49, 873-87	2.9	83
241	Impaired DNase1-mediated degradation of neutrophil extracellular traps is associated with acute thrombotic microangiopathies. *Journal of Thrombosis and Haemostasis*, **2015**, 13, 732-42	15.4	81
240	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, 649-51	4.5	80

239	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, 821-4	4.5	79
238	Measurement of ADAMTS-13 activity in plasma by the FRETS-VWF73 assay: comparison with other assay methods. *Journal of Thrombosis and Haemostasis*, **2006**, 4, 1146-8	15.4	77
237	Plasma DNA is Elevated in Patients with Deep Vein Thrombosis. *Journal of Vascular Surgery: Venous and Lymphatic Disorders*, **2013**, 1,	3.2	75
236	Gut microbiota regulate hepatic von Willebrand factor synthesis and arterial thrombus formation via Toll-like receptor-2. *Blood*, **2017**, 130, 542-553	2.2	70
235	Stability of coagulation assays performed in plasma from citrated whole blood transported at ambient temperature. *Thrombosis and Haemostasis*, **2008**, 99, 416-26	7	70
234	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	4	69
233	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, 7503-7	11.5	68
232	Accuracy of D-dimer/fibrinogen ratio to predict pulmonary embolism: a prospective diagnostic study. *Journal of Thrombosis and Haemostasis*, **2003**, 1, 708-13	15.4	65
231	A common origin of the 4143insA ADAMTS13 mutation. *Thrombosis and Haemostasis*, **2006**, 96, 3-6	7	64
230	Protein C replacement in severe meningococcemia: rationale and clinical experience. *Clinical Infectious Diseases*, **2001**, 32, 1338-46	11.6	62
229	Different disparities of gender and race among the thrombotic thrombocytopenic purpura and hemolytic-uremic syndromes. *American Journal of Hematology*, **2010**, 85, 844-7	7.1	61
228	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, 936-43	6.6	60
227	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, 100-2	4.5	60
226	Initial experience from a double-blind, placebo-controlled, clinical outcome study of ARC1779 in patients with thrombotic thrombocytopenic purpura. *American Journal of Hematology*, **2012**, 87, 430-2	7.1	59
225	Von Willebrand Factor-cleaving Protease in Childhood Diarrhoea-associated Haemolytic Uraemic Syndrome. *Thrombosis and Haemostasis*, **2001**, 85, 975-978	7	59
224	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, 866-7	15.4	58
223	Dural puncture and activated protein C resistance: risk factors for cerebral venous sinus thrombosis. *Journal of Neurology, Neurosurgery and Psychiatry*, **1997**, 63, 351-6	5.5	56
222	Rapid determination of anti-heparin/platelet factor 4 antibody titers in the diagnosis of heparin-induced thrombocytopenia. *American Journal of Medicine*, **2003**, 114, 528-36	2.4	56

221	Severe osteoporosis due to systemic mast cell disease: successful treatment with interferon alpha-2B. *Rheumatology*, **1996**, 35, 898-900	3.9	55
220	Prekallikrein deficiency: the characteristic normalization of the severely prolonged aPTT following increased preincubation time is due to autoactivation of factor XII. *Thrombosis Research*, **2002**, 105, 463-70	8.2	54
219	Cognitive deficits after recovery from thrombotic thrombocytopenic purpura. *Transfusion*, **2009**, 49, 1092-101	2.9	53
218	Evidence for a role of anti-ADAMTS13 autoantibodies despite normal ADAMTS13 activity in recurrent thrombotic thrombocytopenic purpura. *Haematologica*, **2012**, 97, 297-303	6.6	53
217	The International Hereditary Thrombotic Thrombocytopenic Purpura Registry: key findings at enrollment until 2017. *Haematologica*, **2019**, 104, 2107-2115	6.6	52
216	D-dimers predict stroke subtype when assessed early. *Cerebrovascular Diseases*, **2010**, 29, 82-6	3.2	52
215	Familial acquired thrombotic thrombocytopenic purpura: ADAMTS13 inhibitory autoantibodies in identical twins. *Blood*, **2004**, 103, 4195-7	2.2	52
214	Plasma prekallikrein, factor XII, antithrombin III, C1(-)-inhibitor and alpha 2-macroglobulin in critically ill patients with suspected disseminated intravascular coagulation (DIC). *American Journal of Clinical Pathology*, **1984**, 82, 396-404	1.9	52
213	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	2.4	51
212	Treatment of thrombotic thrombocytopenic purpura. *Vox Sanguinis*, **2006**, 90, 245-54	3.1	51
211	The Swiss cohort of elderly patients with venous thromboembolism (SWITCO65+): rationale and methodology. *Journal of Thrombosis and Thrombolysis*, **2013**, 36, 475-83	5.1	50
210	Elevated levels of plasma prekallikrein, high molecular weight kininogen and factor XI in coronary heart disease. *Atherosclerosis*, **2002**, 161, 261-7	3.1	50
209	Role of ADAMTS13 in the pathogenesis, diagnosis, and treatment of thrombotic thrombocytopenic purpura. *Hematology American Society of Hematology Education Program*, **2012**, 2012, 610-616	3.1	50
208	Polypharmacy is associated with an increased risk of bleeding in elderly patients with venous thromboembolism. *Journal of General Internal Medicine*, **2015**, 30, 17-24	4	49
207	Decreasing frequency of plasma exchange complications in patients treated for thrombotic thrombocytopenic purpura-hemolytic uremic syndrome, 1996 to 2011. *Transfusion*, **2012**, 52, 2525-32; quiz 2524	2.9	49
206	Immunoblotting studies of the molecular forms of protein C in plasma. *Thrombosis Research*, **1988**, 52, 33-43	8.2	49
205	Effect of low-molecular weight dextran sulfate on coagulation and platelet function tests. *Thrombosis Research*, **2002**, 105, 441-6	8.2	48
204	Platelets: thrombotic thrombocytopenic purpura. *Hematology American Society of Hematology Education Program*, **2002**, 315-34	3.1	48

203	Sporadic bloody diarrhoea-associated thrombotic thrombocytopenic purpura-haemolytic uraemic syndrome: an adult and paediatric comparison. *British Journal of Haematology*, **2008**, 141, 696-707	4.5	47
202	Protein Z in ischaemic stroke. *British Journal of Haematology*, **2001**, 114, 169-73	4.5	46
201	ADAMTS13 activity in sickle cell disease. *American Journal of Hematology*, **2006**, 81, 492-8	7.1	44
200	Titre of anti-heparin/PF4-antibodies and extent of in vivo activation of the coagulation and fibrinolytic systems. *Thrombosis and Haemostasis*, **2004**, 91, 276-82	7	44
199	Rapid exclusion or confirmation of heparin-induced thrombocytopenia: a single-center experience with 1,291 patients. *Haematologica*, **2012**, 97, 89-97	6.6	43
198	ADAMTS13 activity, von Willebrand factor, factor VIII and D-dimers in COVID-19 inpatients. *Thrombosis Research*, **2020**, 192, 174-175	8.2	42
197	Pregnancy outcomes following recovery from acquired thrombotic thrombocytopenic purpura. *Blood*, **2014**, 123, 1674-80	2.2	40
196	Splenectomy in relapsing and plasma-refractory acquired thrombotic thrombocytopenic purpura. *Haematologica*, **2004**, 89, 320-4	6.6	40
195	Thrombo-Inflammation in Cardiovascular Disease: An Expert Consensus Document from the Third Maastricht Consensus Conference on Thrombosis. *Thrombosis and Haemostasis*, **2020**, 120, 538-564	7	39
194	Frequency and significance of HIV infection among patients diagnosed with thrombotic thrombocytopenic purpura. *Clinical Infectious Diseases*, **2009**, 48, 1129-37	11.6	38
193	The Oklahoma Thrombotic Thrombocytopenic Purpura-Hemolytic Uremic Syndrome Registry: the Swiss connection. *European Journal of Haematology*, **2008**, 80, 277-86	3.8	38
192	Deficiency of von Willebrand factor-cleaving protease in familial and acquired thrombotic thrombocytopenic purpura. *Best Practice and Research: Clinical Haematology*, **1998**, 11, 509-14		36
191	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	4	35
190	Dosing lepirudin in patients with heparin-induced thrombocytopenia and normal or impaired renal function: a single-center experience with 68 patients. *Blood*, **2009**, 113, 2402-9	2.2	34
189	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	7	34
188	Monitoring of heparin treatment. Comparison of 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	1.9	34
187	Hereditary thrombotic thrombocytopenic purpura and the hereditary TTP registry. *Hamostaseologie*, **2013**, 33, 138-43	1.9	33
186	Protein Z in healthy human individuals and in patients with a bleeding tendency. *British Journal of Haematology*, **1998**, 102, 1219-23	4.5	32

185	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, 165-70	1	32
184	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	15.4	32
183	High prevalence of hereditary thrombotic thrombocytopenic purpura in central Norway: from clinical observation to evidence. *Journal of Thrombosis and Haemostasis*, **2016**, 14, 73-82	15.4	32
182	Mycobacterium genavense infection in a patient with long-standing chronic lymphocytic leukaemia. *Journal of Internal Medicine*, **2000**, 248, 343-8	10.8	31
181	No association of APC resistance with myocardial infarction. *Blood Coagulation and Fibrinolysis*, **1995**, 6, 456-9	1	31
180	The splenic autoimmune response to ADAMTS13 in thrombotic thrombocytopenic purpura contains recurrent antigen-binding CDR3 motifs. *Blood*, **2014**, 124, 3469-79	2.2	30
179	Thrombotic microangiopathic syndromes associated with drugs, HIV infection, hematopoietic stem cell transplantation and cancer. *Presse Medicale*, **2012**, 41, e177-88	2.2	30
178	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, 721-4	2.4	29
177	In vitro evaluation of the efficacy and biocompatibility of new, synthetic ABO immunoabsorbents. *Transplantation*, **1995**, 60, 425-30	1.8	28
176	Factor XII clotting activity and antigen levels in patients with thromboembolic disease. *Blood Coagulation and Fibrinolysis*, **1992**, 3, 555-61	1	28
175	Redefining outcomes in immune TTP: an international working group consensus report. *Blood*, **2021** , 137, 1855-1861	2.2	28
174	Use of the pentasaccharide fondaparinux as an anticoagulant during haemodialysis. *Thrombosis and Haemostasis*, **2007**, 98, 1200-7	7	27
173	Evaluation of a Platelet Function Analyser (PFA-100) in patients with a bleeding tendency. *Swiss Medical Weekly*, **2002**, 132, 443-8	3.1	27
172	Depression and cognitive deficits as long-term consequences of thrombotic thrombocytopenic purpura. *Transfusion*, **2017**, 57, 1152-1162	2.9	26
171	Binding of alpha-thrombin to fibrin depends on the quality of the fibrin network. *Biochemical Journal*, **1994**, 298 (Pt 1), 157-63	3.8	26
170	Plasma protein C inhibitor is elevated in survivors of myocardial infarction. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **1997**, 17, 114-8	9.4	26
169	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	7	26
168	Thromboembolism in patients with congenital afibrinogenaemia. Long-term observational data and systematic review. *Thrombosis and Haemostasis*, **2016**, 116, 722-32	7	25

167	Circulating extracellular DNA is an independent predictor of mortality in elderly patients with venous thromboembolism. *PLoS ONE*, **2018**, 13, e0191150	3.7	25
166	Variability of anti-PF4/heparin antibody results obtained by the rapid testing system ID-H/PF4-PaGIA. *Journal of Thrombosis and Haemostasis*, **2009**, 7, 1649-55	15.4	25
165	Von Willebrand Factor in Thrombotic Thrombocytopenic Purpura. *Thrombosis and Haemostasis*, **1999**, 82, 592-600	7	25
164	Open ADAMTS13, induced by antibodies, is a biomarker for subclinical immune-mediated thrombotic thrombocytopenic purpura. *Blood*, **2020**, 136, 353-361	2.2	24
163	Current insights into thrombotic microangiopathies: Thrombotic thrombocytopenic purpura and pregnancy. *Thrombosis Research*, **2015**, 135 Suppl 1, S30-3	8.2	24
162	IFNalpha treatment in systemic mastocytosis. *Annals of Hematology*, **1999**, 78, 483-4	3	23
161	New strategies in diagnosis and treatment of thrombotic thrombocytopenic purpura: case report and review. *European Journal of Pediatrics*, **1999**, 158, 883-7	4.1	23
160	Thrombotic Thrombocytopenic Purpura: Pathophysiology, Diagnosis, and Management. *Journal of Clinical Medicine*, **2021**, 10,	5.1	23
159	Prospective comparison of clinical prognostic scores in elder patients with a pulmonary embolism. *Journal of Thrombosis and Haemostasis*, **2012**, 10, 2270-6	15.4	22
158	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, 663-6	3	22
157	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, 167-72	5.3	22
156	Blood group O and black race are independent risk factors for thrombotic thrombocytopenic purpura associated with severe ADAMTS13 deficiency. *Transfusion*, **2011**, 51, 2237-43	2.9	21
155	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, 1235-7	5.6	21
154	Factor XIII in severe sepsis and septic shock. *Thrombosis Research*, **2007**, 119, 311-8	8.2	20
153	A new substitution, gamma 358 Ser-->Cys, in fibrinogen Milano VII causes defective fibrin polymerization. *Blood*, **1994**, 84, 1874-1880	2.2	20
152	Enhanced specificity of immunoblotting using radiolabeled antigen overlay: studies of blood coagulation factor XII and prekallikrein in plasma. *Analytical Biochemistry*, **1986**, 156, 118-25	3.1	20
151	Purified Human Plasma Kallikrein Does Not Stimulate but Primes Neutrophils for Superoxide Production. *Thrombosis and Haemostasis*, **1989**, 62, 1121-1125	7	20
150	The impact of congenital thrombotic thrombocytopenic purpura on pregnancy complications. *Thrombosis and Haemostasis*, **2014**, 111, 1180-3	7	19

149	The prothrombin time/international normalized ratio (PT/INR) Line: derivation of local INR with commercial thromboplastins and coagulometers--two independent studies. *Journal of Thrombosis and Haemostasis*, **2011**, 9, 140-8	15.4	19
148	Acquired deficiency of von Willebrand factor-cleaving protease in a patient suffering from acute systemic lupus erythematosus. *Rheumatology*, **2001**, 40, 940-2	3.9	19
147	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	7	19
146	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	7	19
145	Quantitative immunoblotting assay of blood coagulation factor XII. *Thrombosis Research*, **1986**, 41, 747-59	3.2	19
144	Progressive multifocal leukoencephalopathy in common variable immunodeficiency: mitigated course under mirtazapine and mefloquine. *Journal of NeuroVirology*, **2015**, 21, 694-701	3.9	18
143	Low molecular weight heparin-induced thrombocytopenia and skin necrosis distant from injection sites. *European Journal of Haematology*, **1994**, 53, 61-3	3.8	18
142	Rituximab for acute plasma-refractory thrombotic thrombocytopenic purpura. A case report and concise review of the literature. *Swiss Medical Weekly*, **2007**, 137, 518-24	3.1	18
141	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, 1132-5	2.7	17
140	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, 417-21		16
139	Usefulness of the D-dimer/fibrinogen ratio to predict deep venous thrombosis. *Journal of Thrombosis and Haemostasis*, **2005**, 3, 385-7	15.4	16
138	How high is the true fibrinogen content of fibrinogen standards?. *Thrombosis Research*, **1989**, 56, 583-92	8.2	16
137	Relapse Rate in Survivors of Acute Autoimmune Thrombotic Thrombocytopenic Purpura Treated with or without Rituximab. *Thrombosis and Haemostasis*, **2018**, 118, 1743-1751	7	16
136	Genetic variations in complement factors in patients with congenital thrombotic thrombocytopenic purpura with renal insufficiency. *International Journal of Hematology*, **2016**, 103, 283-91	2.3	15
135	Bilateral periorbital ecchymoses. An often missed sign of amyloid purpura. *Hamostaseologie*, **2014**, 34, 249-52	1.9	15
134	Performance of a New Fibrin Monomer Assay to Exclude Deep Vein Thrombosis in Symptomatic Outpatients. *Thrombosis and Haemostasis*, **1999**, 81, 50-53	7	15
133	Contact phase of blood coagulation is not activated in edema of high altitude. *Journal of Applied Physiology*, **1989**, 67, 1336-40	3.7	15
132	Role of ADAMTS13 in the pathogenesis, diagnosis, and treatment of thrombotic thrombocytopenic purpura. *Hematology American Society of Hematology Education Program*, **2012**, 2012, 610-6	3.1	15

131	Subcutaneous low-molecular-weight heparin for treatment of Trousseau's syndrome. *Annals of Hematology*, **1997**, 75, 165-7	3	13
130	High Molecular Weight Kininogen Is Cleaved by FXIa at Three Sites: Arg409-Arg410, Lys502-Thr503 and Lys325-Lys326. *Thrombosis and Haemostasis*, **2000**, 83, 709-714	7	13
129	Is factor V Leiden a risk factor for thrombotic microangiopathies without severe ADAMTS13 deficiency?. *Thrombosis and Haemostasis*, **2005**, 94, 1186-1189	7	12
128	Agreement of D-dimer Results Measured by a Rapid ELISA (VIDAS) before and after Storage during 24 h or Transportation of the Original Whole Blood Samples. *Thrombosis and Haemostasis*, **2000**, 83, 177-178	7	12
127	Hemostatic and fibrinolytic parameters in survivors of myocardial infarction: a low plasma level of plasmin-alpha2-antiplasmin complex is an independent predictor of coronary re-events. *Blood Coagulation and Fibrinolysis*, **2001**, 12, 17-24	1	12
126	Thrombotic thrombocytopenic purpura: advances in pathophysiology, diagnosis, and treatment--introduction. *Seminars in Hematology*, **2004**, 41, 1-3	4	11
125	Frequency of protein Z deficiency in patients with ischaemic stroke. *Lancet, The*, **2001**, 358, 840-1	40	11
124	Patent ductus arteriosus generates neonatal hemolytic jaundice with thrombocytopenia in Upshaw-Schulman syndrome. *Blood Advances*, **2019**, 3, 3191-3195	7.8	11
123	Severe plasma prekallikrein deficiency: Clinical characteristics, novel KLKB1 mutations, and estimated prevalence. *Journal of Thrombosis and Haemostasis*, **2020**, 18, 1598-1617	15.4	10
122	Predictors and Outcomes of Recurrent Venous Thromboembolism in Elderly Patients. *American Journal of Medicine*, **2018**, 131, 703.e7-703.e16	2.4	10
121	Rumpel-Leede sign in thrombocytopenia due to Epstein-Barr virus-induced mononucleosis. *British Journal of Haematology*, **2010**, 148, 2	4.5	10
120	Images in clinical medicine. Capnocytophaga canimorsus sepsis. *New England Journal of Medicine*, **1998**, 339, 1827	59.2	10
119	Fibrinogens Bern IV, Bern V and Milano XI. *Blood Coagulation and Fibrinolysis*, **1999**, 10, 93-100	1	10
118	Annual incidence and severity of acute episodes in hereditary thrombotic thrombocytopenic purpura. *Blood*, **2021**, 137, 3563-3575	2.2	10
117	Association between thyroid dysfunction and venous thromboembolism in the elderly: a prospective cohort study. *Journal of Thrombosis and Haemostasis*, **2016**, 14, 685-94	15.4	10
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