Christine Mannhalter

List of Publications by Citations

311 papers	10,026 citations	54 h-index	81 g-index
318 ext. papers	10,858 ext. citations	5.2 avg, IF	5.35 L-index

#	Paper	IF	Citations
311	Mean platelet volume is an independent risk factor for myocardial infarction but not for coronary artery disease. *British Journal of Haematology*, **2002**, 117, 399-404	4.5	312
310	Is elevated mean platelet volume associated with a worse outcome in patients with acute ischemic cerebrovascular events?. *Stroke*, **2004**, 35, 1688-91	6.7	220
309	Association of an ABCB1 gene haplotype with pharmacoresistance in temporal lobe epilepsy. *Neurology*, **2004**, 63, 1087-9	6.5	173
308	Most CD56+ intestinal lymphomas are CD8+CD5-T-cell lymphomas of monomorphic small to medium size histology. *American Journal of Pathology*, **1998**, 153, 1483-90	5.8	172
307	Risk of pregnancy-associated recurrent venous thromboembolism in women with a history of venous thrombosis. *Journal of Thrombosis and Haemostasis*, **2005**, 3, 949-54	15.4	170
306	A Single Genetic Origin for the Common Prothrombotic G20210A Polymorphism in the Prothrombin Gene. *Blood*, **1998**, 92, 1119-1124	2.2	162
305	Heme oxygenase-1 gene promoter microsatellite polymorphism is associated with restenosis after percutaneous transluminal angioplasty. *Journal of Endovascular Therapy*, **2001**, 8, 433-40	2.5	155
304	Morphine decreases clopidogrel concentrations and effects: a randomized, double-blind, placebo-controlled trial. *Journal of the American College of Cardiology*, **2014**, 63, 630-635	15.1	153
303	Swarm Intelligence-Enhanced Detection of Non-Small-Cell Lung Cancer Using Tumor-Educated Platelets. *Cancer Cell*, **2017**, 32, 238-252.e9	24.3	150
302	Temporary increase in the risk for recurrence during pregnancy in women with a history of venous thromboembolism. *Blood*, **2002**, 100, 1060-2	2.2	149
301	Primary follicular lymphoma of the duodenum is a distinct mucosal/submucosal variant of follicular lymphoma: a retrospective study of 63 cases. *Journal of Clinical Oncology*, **2011**, 29, 1445-51	2.2	148
300	The Risk of Recurrent Venous Thromboembolism in Patients with and without Factor V Leiden. *Thrombosis and Haemostasis*, **1997**, 77, 624-628	7	141
299	Dipeptidylpeptidase IV (CD26) defines leukemic stem cells (LSC) in chronic myeloid leukemia. *Blood*, **2014**, 123, 3951-62	2.2	140
298	Follicular lymphomas BCL-2/IgH junctions contain templated nucleotide insertions: novel insights into the mechanism of t(14;18) translocation. *Blood*, **2000**, 95, 3520-3529	2.2	135
297	High expression of lipoprotein lipase in poor risk B-cell chronic lymphocytic leukemia. *Leukemia*, **2005**, 19, 1216-23	10.7	118
296	The Risk of Early Recurrent Venous Thromboembolism after Oral Anticoagulant Therapy in Patients with the G20210A Transition in the Prothrombin Gene. *Thrombosis and Haemostasis*, **1999**, 81, 14-17	7	114
295	A meta-analysis of candidate gene polymorphisms and ischemic stroke in 6 study populations: association of lymphotoxin-alpha in nonhypertensive patients. *Stroke*, **2009**, 40, 683-95	6.7	102

294	Donor Leukocyte Infusion for Leukemic Relapse After Allogeneic Marrow Transplantation: Lack of Residual Donor Hematopoiesis Predicts Aplasia. *Blood*, **1997**, 89, 3113-3117	2.2	101
293	Origin of human mast cells: development from transplanted hematopoietic stem cells after allogeneic bone marrow transplantation. *Blood*, **1994**, 84, 2954-2959	2.2	94
292	Heme oxygenase-1 gene promoter polymorphism is associated with abdominal aortic aneurysm. *Thrombosis Research*, **2002**, 106, 131-6	8.2	91
291	High concentrations of soluble P-selectin are associated with risk of venous thromboembolism and the P-selectin Thr715 variant. *Clinical Chemistry*, **2007**, 53, 1235-43	5.5	90
290	Inclusion body myopathy and Paget disease is linked to a novel mutation in the VCP gene. *Neurology*, **2005**, 65, 1304-5	6.5	89
289	Mutation (677 C to T) in the methylenetetrahydrofolate reductase gene aggravates hyperhomocysteinemia in hemodialysis patients. *Kidney International*, **1997**, 52, 517-23	9.9	88
288	Thrombin as a multi-functional enzyme. Focus on in vitro and in vivo effects. *Thrombosis and Haemostasis*, **2011**, 106, 1020-33	7	86
287	Rituximab serum concentrations during immuno-chemotherapy of follicular lymphoma correlate with patient gender, bone marrow infiltration and clinical response. *Haematologica*, **2012**, 97, 1431-8	6.6	86
286	Evidence of a U-shaped association between factor XII activity and overall survival. *Journal of Thrombosis and Haemostasis*, **2007**, 5, 1143-8	15.4	86
285	Factor XII activity and antigen concentrations in patients suffering from recurrent thrombosis. *Fibrinolysis*, **1987**, 1, 259-263		85
284	Polymorphisms in coagulation factor genes and their impact on arterial and venous thrombosis. *Clinica Chimica Acta*, **2003**, 330, 31-55	6.2	83
283	The Prevalence of Factor XII Deficiency in 103 Orally Anticoagulated Outpatients Suffering from Recurrent Venous and/or Arterial Thromboembolism. *Thrombosis and Haemostasis*, **1992**, 68, 285-290	7	83
282	Matched case-control study on factor V Leiden and the prothrombin G20210A mutation in patients with ischemic stroke/transient ischemic attack up to the age of 60 years. *Stroke*, **2005**, 36, 1405-9	6.7	80
281	Association of a functional polymorphism in the clopidogrel target receptor gene, P2Y12, and the risk for ischemic cerebrovascular events in patients with peripheral artery disease. *Stroke*, **2005**, 36, 1394-9	6.7	77
280	Heme oxygenase-1 genotype and restenosis after balloon angioplasty: a novel vascular protective factor. *Journal of the American College of Cardiology*, **2004**, 43, 950-7	15.1	76
279	Protein S Deficiency: A Database of Mutations Summary of the First Update. *Thrombosis and Haemostasis*, **2000**, 84, 918-918	7	74
278	Templated nucleotide addition and immunoglobulin JH-gene utilization in t(11;14) junctions: implications for the mechanism of translocation and the origin of mantle cell lymphoma. *Cancer Research*, **2001**, 61, 1629-36	10.1	73
277	Human toll-like receptor 4 mutations are associated with susceptibility to invasive meningococcal disease in infancy. *Pediatric Infectious Disease Journal*, **2006**, 25, 80-1	3.4	71

276	Factor V Leiden mutation carriership and venous thromboembolism in polycythemia vera and essential thrombocythemia. *American Journal of Hematology*, **2002**, 71, 1-6	7.1	71
275	Molecular cytogenetic evidence of t(14;18)(IGH;BCL2) in a substantial proportion of primary cutaneous follicle center lymphomas. *American Journal of Surgical Pathology*, **2006**, 30, 529-36	6.7	70
274	Immunoreactivity of B-cell markers (CD79a, L26) in rare cases of extranodal cytotoxic peripheral T-(NK/T-) cell lymphomas. *Modern Pathology*, **2000**, 13, 766-72	9.8	70
273	High expression of activation-induced cytidine deaminase (AID) mRNA is associated with unmutated IGVH gene status and unfavourable cytogenetic aberrations in patients with chronic lymphocytic leukaemia. *Leukemia*, **2004**, 18, 756-62	10.7	69
272	A microsatellite polymorphism in the heme oxygenase-1 gene promoter is associated with increased bilirubin and HDL levels but not with coronary artery disease. *Thrombosis and Haemostasis*, **2004**, 91, 155-61	7	67
271	The Prevalence of Moderate and Severe FXII (Hageman Factor) Deficiency among the Normal Population: Evaluation of the Incidence of FXII Deficiency among 300 Healthy Blood Donors. *Thrombosis and Haemostasis*, **1994**, 71, 068-072	7	67
270	The KIT D816V allele burden predicts survival in patients with mastocytosis and correlates with the WHO type of the disease. *Allergy: European Journal of Allergy and Clinical Immunology*, **2014**, 69, 810-3	9.3	64
269	The (-174) G/C polymorphism in the interleukin-6 gene is associated with the severity of acute cerebrovascular events. *Thrombosis Research*, **2003**, 110, 181-6	8.2	63
268	Major determinants of hyperhomocysteinemia in peritoneal dialysis patients. *Kidney International*, **1998**, 53, 1775-82	9.9	62
267	ABO mismatch increases transplant-related morbidity and mortality in patients given nonmyeloablative allogeneic HPC transplantation. *Transfusion*, **2003**, 43, 1153-61	2.9	62
266	Oral contraceptives enhance the risk of clinical manifestation of venous thrombosis at a young age in females homozygous for factor V Leiden. *British Journal of Haematology*, **1996**, 93, 487-90	4.5	60
265	Preeclampsia and fetal loss in women with a history of venous thromboembolism. *Arteriosclerosis, Thrombosis, and Vascular Biology*, **2001**, 21, 874-9	9.4	59
264	Interleukin-1 receptor antagonist genotype is associated with coronary atherosclerosis in patients with type 2 diabetes. *Diabetes*, **2002**, 51, 3582-5	0.9	59
263	Variable prognostic value of FLT3 internal tandem duplications in patients with de novo AML and a normal karyotype, t(15;17), t(8;21) or inv(16). *The Hematology Journal*, **2002**, 3, 283-9		59
262	Donor heme oxygenase-1 genotype is associated with renal allograft function. *Transplantation*, **2004**, 77, 538-42	1.8	58
261	Evaluation of RNA isolation methods and reference genes for RT-PCR analyses of rare target RNA. *Clinical Chemistry and Laboratory Medicine*, **2000**, 38, 171-7	5.9	56
260	4G4G genotype of the plasminogen activator inhibitor-1 promoter polymorphism associates with disseminated intravascular coagulation in children with systemic meningococcemia. *Journal of Thrombosis and Haemostasis*, **2007**, 5, 2049-54	15.4	55
259	Influence of functional haplotypes in the drug transporter gene ABCB1 on central nervous system drug distribution in humans. *Clinical Pharmacology and Therapeutics*, **2005**, 78, 182-90	6.1	55

258	Regression of colonic low grade B cell lymphoma of the mucosa associated lymphoid tissue type after eradication of Helicobacter pylori. *Gut*, **2000**, 46, 133-5	19.2	55
257	FLAG (fludarabine, cytosine arabinoside, G-CSF) for refractory and relapsed acute myeloid leukemia. *Annals of Hematology*, **1996**, 73, 265-71	3	54
256	4G/5G promoter polymorphism in the plasminogen-activator-inhibitor-1 gene in children with systemic meningococcaemia. *European Journal of Pediatrics*, **2005**, 164, 486-90	4.1	52
255	Polymorphisms of the inflammatory system and risk of ischemic cerebrovascular events. *Clinical Chemistry and Laboratory Medicine*, **2006**, 44, 918-23	5.9	51
254	The risk of recurrent venous thromboembolism in heterozygous carriers of factor V Leiden and a first spontaneous venous thromboembolism. *Archives of Internal Medicine*, **2002**, 162, 2357-60		51
253	Clonality analysis using X-chromosome inactivation at the human androgen receptor gene (Humara). Evaluation of large cohorts of patients with chronic myeloproliferative diseases, secondary neutrophilia, and reactive thrombocytosis. *American Journal of Clinical Pathology*, **1999**,	1.9	51
252	Low-density lipoprotein receptor-related protein 1 polymorphism 663 C > T affects clotting factor VIII activity and increases the risk of venous thromboembolism. *Journal of Thrombosis and Haemostasis*, **2007**, 5, 497-502	15.4	50
251	A microsatellite polymorphism in the heme oxygenase-1 gene promoter is associated with risk for melanoma. *International Journal of Cancer*, **2006**, 119, 1312-5	7.5	49
250	Coagulation factor XII (FXII) activity, activated FXII, distribution of FXII C46T gene polymorphism and coronary risk. *Journal of Thrombosis and Haemostasis*, **2008**, 6, 291-296	15.4	48
249	CD5 expression in a lymphoma of the mucosa-associated lymphoid tissue (MALT)-type as a marker for early dissemination and aggressive clinical behaviour. *Leukemia and Lymphoma*, **2001**, 42, 823-9	1.9	48
248	The interleukin-6 G(-174)C promoter polymorphism does not determine plasma interleukin-6 concentrations in experimental endotoxemia in humans. *Clinical Chemistry*, **2004**, 50, 195-200	5.5	47
247	Pregnancy-associated risk for venous thromboembolism and pregnancy outcome in women homozygous for factor V Leiden. *The Hematology Journal*, **2000**, 1, 37-41		46
246	Biomarkers predictive of venous thromboembolism in patients with newly diagnosed high-grade gliomas. *Neuro-Oncology*, **2014**, 16, 1645-51	1	45
245	Quantification of minimal residual disease in patients with BCR-ABL-positive acute lymphoblastic leukaemia using quantitative competitive polymerase chain reaction. *British Journal of Haematology*, **1999**, 106, 634-43	4.5	45
244	Genetic fingerprinting in mouthwashes of patients after allogeneic bone marrow transplantation. *Bone Marrow Transplantation*, **1999**, 24, 95-8	4.4	45
243	Heme Oxygenase-1 Gene Promoter Microsatellite Polymorphism Is Associated With Restenosis After Percutaneous Transluminal Angioplasty. *Journal of Endovascular Therapy*, **2001**, 8, 433-440	2.5	45
242	Human high molecular weight kininogen. Effects of cleavage by kallikrein on protein structure and procoagulant activity.. *Journal of Biological Chemistry*, **1980**, 255, 6433-6438	5.4	45
241	Routinely available biomarkers improve prediction of long-term mortality in stable coronary artery disease: the Vienna and Ludwigshafen Coronary Artery Disease (VILCAD) risk score. *European Heart Journal*, **2012**, 33, 2282-9	9.5	44

240	Next-generation sequencing identifies major DNA methylation changes during progression of Ph+ chronic myeloid leukemia. *Leukemia*, **2016**, 30, 1861-8	10.7	44
239	C677T MTHFR mutation and factor V Leiden mutation in patients with TIA/minor stroke: a case-control study. *Thrombosis Research*, **1999**, 93, 61-9	8.2	43
238	Circulating Free Methylated Tumor DNA Markers for Sensitive Assessment of Tumor Burden and Early Response Monitoring in Patients Receiving Systemic Chemotherapy for Colorectal Cancer Liver Metastasis. *Annals of Surgery*, **2018**, 268, 894-902	7.8	43
237	High frequency of concomitant mastocytosis in patients with acute myeloid leukemia exhibiting the transforming KIT mutation D816V. *Molecular Oncology*, **2010**, 4, 335-46	7.9	42
236	Haem oxygenase-1 genotype and cardiovascular adverse events in patients with peripheral artery disease. *European Journal of Clinical Investigation*, **2005**, 35, 731-7	4.6	42
235	Human high molecular weight kininogen. Effects of cleavage by kallikrein on protein structure and procoagulant activity. *Journal of Biological Chemistry*, **1980**, 255, 6433-8	5.4	42
234	Factor V Leiden mutation increases the risk for venous thromboembolism in cancer patients - results from the Vienna Cancer And Thrombosis Study (CATS). *Journal of Thrombosis and Haemostasis*, **2015**, 13, 17-22	15.4	41
233	Defective diacylglycerol-induced Ca2+ entry but normal agonist-induced activation responses in TRPC6-deficient mouse platelets. *Journal of Thrombosis and Haemostasis*, **2012**, 10, 419-29	15.4	41
232	Rapid establishment of long-term culture-initiating cells of donor origin after nonmyeloablative allogeneic hematopoietic stem-cell transplantation, and significant prognostic impact of donor T-cell chimerism on stable engraftment and progression-free survival. *Transplantation*, **2003**, 76, 230-6	1.8	41
231	Diffuse large B-cell lymphomas with plasmablastic/plasmacytoid features are associated with TP53 deletions and poor clinical outcome. *Leukemia*, **2004**, 18, 146-55	10.7	41
230	The 4G/4G genotype at nucleotide position -675 in the promotor region of the plasminogen activator inhibitor 1 (PAI-1) gene is less frequent in young patients with minor stroke than in controls. *British Journal of Haematology*, **2000**, 110, 469-71	4.5	41
229	Protein S deficiency type I: identification of point mutations in 9 of 10 families. *Blood*, **1995**, 86, 3444-3451	11.2	40
228	Diagnosis of Wilson's disease in an asymptomatic sibling by DNA linkage analysis. *Gastroenterology*, **1995**, 109, 2015-8	13.3	40
227	Parameters influencing FVIII pharmacokinetics in patients with severe and moderate haemophilia A. *Haemophilia*, **2015**, 21, 343-50	3.3	39
226	Identification of oncostatin M as a JAK2 V617F-dependent amplifier of cytokine production and bone marrow remodeling in myeloproliferative neoplasms. *FASEB Journal*, **2012**, 26, 894-906	0.9	39
225	GAB2 is a novel target of 11q amplification in AML/MDS. *Genes Chromosomes and Cancer*, **2006**, 45, 798-807	3.9	39
224	A common C-->T polymorphism at nt 46 in the promoter region of coagulation factor XII is associated with decreased factor XII activity. *Thrombosis Research*, **2001**, 101, 255-60	8.2	39
223	Identification of CD25 as STAT5-Dependent Growth Regulator of Leukemic Stem Cells in Ph+ CML. *Clinical Cancer Research*, **2016**, 22, 2051-61	12.9	38

222	Multiplex PCR for rapid detection of T-cell receptor-gamma chain gene rearrangements in patients with lymphoproliferative diseases. *British Journal of Haematology*, **1996**, 94, 136-9	4.5	38
221	Opposite effects of CX3CR1 receptor polymorphisms V249I and T280M on the development of acute coronary syndrome. A possible implication of fractalkine in inflammatory activation. *Thrombosis and Haemostasis*, **2005**, 93, 949-54	7	37
220	Monitoring of minimal residual leukemia in patients with MLL-AF9 positive acute myeloid leukemia by RT-PCR. *Leukemia*, **1999**, 13, 1519-24	10.7	37
219	Common IgE-epitopes of recombinant Phl p I, the major timothy grass pollen allergen and natural group I grass pollen isoallergens. *Molecular Immunology*, **1996**, 33, 417-26	4.3	37
218	Mechanism of the chromosomal translocation t(14;18) in lymphoma: detection of a 45-Kd breakpoint binding protein. *Blood*, **1993**, 81, 1833-1840	2.2	37
217	Homocysteine levels in polycythaemia vera and essential thrombocythaemia. *British Journal of Haematology*, **1999**, 105, 551-555	4.5	36
216	Hepatosplenic gammadelta T-cell lymphoma successfully treated with a combination of alemtuzumab and cladribine. *Annals of Oncology*, **2008**, 19, 1025-6	10.3	35
215	Heme oxygenase-1 genotype is a vascular anti-inflammatory factor following balloon angioplasty. *Journal of Endovascular Therapy*, **2002**, 9, 385-94	2.5	35
214	Association of low-grade inflammation with nephropathy in type 2 diabetic patients: role of elevated CRP-levels and 2 different gene-polymorphisms of proinflammatory cytokines. *Experimental and Clinical Endocrinology and Diabetes*, **2007**, 115, 38-41	2.3	34
213	The effect of a promoter polymorphism in the heme oxygenase-1 gene on the risk of ischaemic cerebrovascular events: the influence of other vascular risk factors. *Thrombosis Research*, **2004**, 113, 217-23	2.2	34
212	Transcription of AML1/ETO in bone marrow and cord blood of individuals without acute myelogenous leukemia. *Blood*, **2002**, 100, 2267-8	2.2	34
211	Long-term treatment with imatinib results in profound mast cell deficiency in Ph+ chronic myeloid leukemia. *Oncotarget*, **2015**, 6, 3071-84	3.3	34
210	Prognostic significance of molecular staging by PCR-amplification of immunoglobulin gene rearrangements in diffuse large B-cell lymphoma (DLBCL). *Leukemia*, **2004**, 18, 1102-7	10.7	33
209	Detection of engraftment and mixed chimerism following bone marrow transplantation using PCR amplification of a highly variable region-variable number of tandem repeats (VNTR) in the von Willebrand factor gene. *Annals of Hematology*, **1991**, 63, 227-8	3	33
208	Overexpression of uridine diphospho glucuronosyltransferase 2B17 in high-risk chronic lymphocytic leukemia. *Blood*, **2013**, 121, 1175-83	2.2	32
207	Resistance to activated protein C (APC): mutation at Arg506 of coagulation factor V and vascular access thrombosis in haemodialysis patients. *Nephrology Dialysis Transplantation*, **1996**, 11, 668-72	4.3	31
206	Risk of recurrence after a first venous thromboembolic event in young women. *Haematologica*, **2007**, 92, 1201-7	6.6	31
205	Quantitative comparison of the expression of EVI1 and its presumptive antagonist, MDS1/EVI1, in patients with myeloid leukemia. *Genes Chromosomes and Cancer*, **2003**, 36, 80-9	5	31

204	Homozygosity for the C-->T polymorphism at nucleotide 46 in the 5Nuntranslated region of the factor XII gene protects from development of acute coronary syndrome. *British Journal of Haematology*, **2001**, 115, 1007-9	4.5	31
203	Impact of variables of the P-selectin - P-selectin glycoprotein ligand-1 axis on leukocyte-platelet interactions in cardiovascular disease. *Thrombosis and Haemostasis*, **2015**, 113, 806-12	7	30
202	PCR-monitoring of minimal residual leukaemia after conventional chemotherapy and bone marrow transplantation in BCR-ABL-positive acute lymphoblastic leukaemia. *British Journal of Haematology*, **1995**, 89, 937-41	4.5	30
201	Polymorphisms in the interleukin-1 gene cluster in children and young adults with systemic meningococcemia. *Clinical Chemistry*, **2006**, 52, 511-4	5.5	30
200	CYP2C9*2 and CYP2C9*3 alleles confer a lower risk for myocardial infarction. *Clinical Chemistry*, **2004**, 50, 2395-8	5.5	30
199	The K121Q polymorphism in the plasma cell membrane glycoprotein 1 gene predisposes to early myocardial infarction. *Journal of Molecular Medicine*, **2002**, 80, 791-5	5.5	30
198	High proportion of patients with bleeding of unknown cause in persons with a mild-to-moderate bleeding tendency: Results from the Vienna Bleeding Biobank (VIBB). *Haemophilia*, **2018**, 24, 405-413	3.3	29
197	Determinants of factor VIII plasma levels in carriers of haemophilia A and in control women. *Haemophilia*, **2010**, 16, 111-7	3.3	29
196	The C-reactive protein (+)1444C/T alteration modulates the inflammation and coagulation response in human endotoxemia. *Clinical Chemistry*, **2006**, 52, 1952-7	5.5	29
195	Is the factor XIII 34Val/Leu polymorphism a protective factor for cerebrovascular disease?. *British Journal of Haematology*, **2003**, 120, 310-4	4.5	29
194	Combined polymerase chain reaction approach for clonality detection in lymphoid neoplasms. *Diagnostic Molecular Pathology*, **1999**, 8, 80-91		29
193	The effect of p22-PHOX (CYBA) polymorphisms on premature coronary artery disease (≤40 years of age). *Thrombosis and Haemostasis*, **2011**, 105, 529-34	7	28
192	Interleukin-6 and interleukin-6 promoter polymorphism (−174) G>C in patients with spontaneous venous thromboembolism. *Thrombosis and Haemostasis*, **2006**, 95, 802-806	7	28
191	T- and B-cell clonality and frequency of human herpes viruses-6, -8 and Epstein Barr virus in angioimmunoblastic T-cell lymphoma. *Hematological Oncology*, **2004**, 22, 169-77	1.3	28
190	P-selectin gene haplotypes modulate soluble P-selectin concentrations and contribute to the risk of venous thromboembolism. *Thrombosis and Haemostasis*, **2008**, 99, 899-904	7	27
189	Improved characteristics of aPC-resistance assay: Coatest aPC resistance by predilution of samples with factor V deficient plasma. *American Journal of Clinical Pathology*, **1996**, 106, 588-93	1.9	27
188	Relative seroprevalence of human herpes viruses in patients with chronic lymphocytic leukaemia. *European Journal of Clinical Investigation*, **2009**, 39, 497-506	4.6	26
187	Leukemia-free survival and mortality in patients with refractory or relapsed acute leukemia given marrow transplants from sibling and unrelated donors. *Bone Marrow Transplantation*, **1998**, 21, 673-8	4.4	26

186	The biological and clinical significance of MLL abnormalities in haematological malignancies. *European Journal of Clinical Investigation*, **2004**, 34 Suppl 2, 12-24	4.6	26
185	Multiplexed Mutagenically Separated PCR: Simultaneous Single-Tube Detection of the Factor V R506Q (G1691A), the Prothrombin G20210A, and the Methylenetetrahydrofolate Reductase A223V (C677T) Variants. *Clinical Chemistry*, **2001**, 47, 333-335	5.5	26
184	Genetic and nongenetic factors influencing plasma homocysteine levels in patients with ischemic cerebrovascular disease and in healthy control subjects. *Translational Research*, **1999**, 133, 575-82		26
183	The role of fibrinogen plasma levels, the -455G>A fibrinogen and the factor XIII A subunit (FXIII-A) Val34Leu polymorphism in cancer-associated venous thrombosis. *Thrombosis and Haemostasis*, **2011**, 106, 908-13	7	25
182	Polymorphisms associated with both noncardioembolic stroke and coronary heart disease: vienna stroke registry. *Cerebrovascular Diseases*, **2009**, 28, 499-504	3.2	25
181	Clinical restenosis after coronary stent implantation is associated with the heme oxygenase-1 gene promoter polymorphism and the heme oxygenase-1 +99G/C variant. *Clinical Chemistry*, **2005**, 51, 1661-5	5.5	25
180	Protein C promoter polymorphisms associate with sepsis in children with systemic meningococcemia. *Human Genetics*, **2007**, 122, 183-90	6.3	24
179	Candidate genetic risk factors of stroke: results of a multilocus genotyping assay. *Clinical Chemistry*, **2007**, 53, 600-5	5.5	24
178	Is low serum bilirubin an independent risk factor for coronary artery disease in men but not in women?. *Clinical Chemistry*, **2003**, 49, 1201-4	5.5	24
177	Glycoprotein Ib polymorphisms influence platelet plug formation under high shear rates. *British Journal of Haematology*, **2003**, 120, 652-5	4.5	24
176	Twelve novel and two recurrent mutations in 14 Austrian families with hereditary protein C deficiency. *Blood Coagulation and Fibrinolysis*, **1993**, 4, 273-80	1	24
175	Coagulation factor XII (FXII) activity, activated FXII, distribution of FXII C46T gene polymorphism and coronary risk. *Journal of Thrombosis and Haemostasis*, **2008**, 6, 291-6	15.4	24
174	Polymorphisms in inflammatory genes and the risk of ischemic stroke and transient ischemic attack: results of a multilocus genotyping assay. *Clinical Chemistry*, **2009**, 55, 134-8	5.5	23
173	Relapse of Philadelphia chromosome positive acute lymphoblastic leukaemia after marrow transplantation: sustained molecular remission after early and dose-escalating infusion of donor leucocytes. *British Journal of Haematology*, **1997**, 97, 161-4	4.5	23
172	Impact of environmental and hereditary risk factors on the clinical manifestation of thrombophilia in homozygous carriers of factor V:G1691A. *Journal of Thrombosis and Haemostasis*, **2004**, 2, 430-6	15.4	23
171	No evidence for an increased risk of venous thrombosis in patients with factor V Leiden by the homozygous 677 C to T mutation in the methylenetetrahydrofolate-reductase gene. *Blood Coagulation and Fibrinolysis*, **1999**, 10, 101-5	1	23
170	Differential impact of cytochrome 2C9 allelic variants on clopidogrel-mediated platelet inhibition determined by five different platelet function tests. *International Journal of Cardiology*, **2013**, 166, 126-31	3.2	22
169	DNA repair polymorphisms associated with cytogenetic subgroups in B-cell chronic lymphocytic leukemia. *Genes Chromosomes and Cancer*, **2009**, 48, 760-7	5	22

168	The endocardial binary appearance (‘binary sign’) is an unreliable marker for echocardiographic detection of Fabry disease in patients with left ventricular hypertrophy. *European Journal of Echocardiography*, **2011**, 12, 744-9		22
167	The EGF A61G polymorphism is associated with disease-free period and survival in malignant melanoma. *Journal of Investigative Dermatology*, **2006**, 126, 2242-6	4.3	22
166	Chronic myeloid leukemia: pathophysiology, diagnostic parameters, and current treatment concepts. *Wiener Klinische Wochenschrift*, **2003**, 115, 485-504	2.3	22
165	FVIII-binding IgG modulates FVIII half-life in patients with severe and moderate hemophilia A without inhibitors. *Blood*, **2016**, 128, 293-6	2.2	22
164	Butyrylcholinesterase activity predicts long-term survival in patients with coronary artery disease. *Clinical Chemistry*, **2012**, 58, 1055-8	5.5	21
163	Evaluation of a new screening assay ProC Global for identification of defects in the protein C/protein S anticoagulant pathway. *Thrombosis Research*, **1997**, 87, 501-10	8.2	21
162	Thrombophilic polymorphisms--factor V Leiden, prothrombin G20210A, and methylenetetrahydrofolate reductase C677T mutations--and preterm birth. *Wiener Klinische Wochenschrift*, **2004**, 116, 622-6	2.3	21
161	Monitoring of minimal residual disease in patients with MLL-AF6-positive acute myeloid leukaemia by reverse transcriptase polymerase chain reaction. *British Journal of Haematology*, **2000**, 109, 622-8	4.5	21
160	Clinical features of thrombophilia in families with gene defects in protein C or protein S combined with factor V Leiden. *Blood Coagulation and Fibrinolysis*, **1998**, 9, 85-9	1	21
159	Prothrombin Fragment F1+2 Is not Predictive for Recurrent Venous Thromboembolism. *Thrombosis and Haemostasis*, **1997**, 77, 0829-0833	7	21
158	Mean platelet volume predicts outcome in patients with asymptomatic carotid artery disease. *European Journal of Clinical Investigation*, **2014**, 44, 22-8	4.6	20
157	Prognostic value of neutrophils in patients with asymptomatic carotid artery disease. *Atherosclerosis*, **2013**, 231, 274-80	3.1	20
156	No impact of endogenous prothrombotic conditions on the risk of central venous line-related thrombotic events in children: results of the KIDCAT study (KIDs with Catheter Associated Thrombosis). *Journal of Thrombosis and Haemostasis*, **2014**, 12, 1610-5	15.4	20
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