

Maurizio Margaglione

List of Publications by Year in Descending Order

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The third column is the impact factor (IF) of the journal, and the fourth column is the number of citations of the article.

136
papers

3,215
citations

28
h-index

52
g-index

143
ext. papers

3,700
ext. citations

4.5
avg, IF

4.9
L-index

#	Paper	IF	Citations
136	Obstetric outcomes in pregnant COVID-19 women: the imbalance of von Willebrand factor and ADAMTS13 axis.. <i>BMC Pregnancy and Childbirth</i> , 2022 , 22, 142	3.2	2
135	Psychosomatic syndromes are associated with IL-6 pro-inflammatory cytokine in heart failure patients.. <i>PLoS ONE</i> , 2022 , 17, e0265282	3.7	0
134	Mortality and clinical outcome of Italian patients undergoing orthopaedic surgery: effect of peri-operative blood transfusion. <i>Blood Transfusion</i> , 2021 , 19, 284-291	3.6	
133	Italian experience with rVIII-single chain: a survey of patients with haemophilia A and their physicians. <i>Journal of Thrombosis and Thrombolysis</i> , 2021 , 1	5.1	1
132	Increased SARS-CoV-2 seroprevalence in healthy blood donors after the second pandemic wave in South-Eastern Italy: evidence for asymptomatic young donors as potential virus spreaders. <i>Infectious Diseases</i> , 2021 , 1-6	3.1	1
131	Use of low-molecular weight heparin, transfusion and mortality in COVID-19 patients not requiring ventilation. <i>Journal of Thrombosis and Thrombolysis</i> , 2021 , 52, 772-778	5.1	2
130	Homozygous methylentetrahydrofolate reductase C667T genotype anticipates age at venous thromboembolism by one decade. <i>Blood Coagulation and Fibrinolysis</i> , 2021 , 32, 382-386	1	1
129	The Genetics of Hereditary Angioedema: A Review. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	8
128	Findings from a multicentre, observational study on reproductive outcomes in women with unexplained recurrent pregnancy loss: the OTTILIA registry. <i>Human Reproduction</i> , 2021 , 36, 2083-2090	5.7	5
127	Results from a survey in healthy blood donors in South Eastern Italy indicate that we are far away from herd immunity to SARS-CoV-2. <i>Journal of Medical Virology</i> , 2021 , 93, 1739-1742	19.7	19
126	Mortality and Transfusion Requirements in COVID-19 Hospitalized Italian Patients According to Severity of the Disease. <i>Journal of Clinical Medicine</i> , 2021 , 10,	5.1	3
125	Summary of the Available Molecular Methods for Detection of SARS-CoV-2 during the Ongoing Pandemic. <i>International Journal of Molecular Sciences</i> , 2021 , 22,	6.3	18
124	Pathogenic Variant (1085G>A) Linked to Infantile Progressive Neurological Disorder: Evidence of Maternal Transmission by Germline Mosaicism and Influence of a Contemporary in cis Variant (1535T>C). <i>Genes</i> , 2021 , 12,	4.2	1
123	Genome-Wide Association Study Identifies First Locus Associated with Susceptibility to Cerebral Venous Thrombosis. <i>Annals of Neurology</i> , 2021 , 90, 777-788	9.4	4
122	Low Prevalence of Antibodies to SARS-CoV-2 and Undetectable Viral Load in Seropositive Blood Donors from South-Eastern Italy. <i>Acta Haematologica</i> , 2021 , 144, 580-584	2.7	4
121	Severe systemic thrombosis in a young COVID-19 patient with a rare homozygous prothrombin G20210A mutation. <i>Infezioni in Medicina</i> , 2021 , 29, 259-262	3.6	1
120	Anemone study: prevalence of risk factors for superficial vein thrombosis in a large Italian population of blood donors. <i>Journal of Thrombosis and Thrombolysis</i> , 2020 , 50, 689-696	5.1	

119	Severe bleeding and absent ADP-induced platelet aggregation associated with inherited combined CalDAG-GEFI and P2Y deficiencies. <i>Haematologica</i> , 2020 , 105, e361-e364	6.6	2
118	A myoferlin gain-of-function variant associates with a new type of hereditary angioedema. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020 , 75, 2989-2992	9.3	34
117	Thromboelastography parameters in Italian pregnant women: do antithrombotic drugs change reference values?. <i>Journal of Investigative Medicine</i> , 2020 , 68, 902-905	2.9	3
116	Anticoagulation in Italian patients with venous thromboembolism and thrombophilic alterations: findings from START2 register study. <i>Blood Transfusion</i> , 2020 , 18, 486-495	3.6	5
115	International Consensus on the Use of Genetics in the Management of Hereditary Angioedema. <i>Journal of Allergy and Clinical Immunology: in Practice</i> , 2020 , 8, 901-911	5.4	28
114	Impaired control of the contact system in hereditary angioedema with normal C1-inhibitor. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2020 , 75, 1394-1403	9.3	14
113	The curious incident of a cavum velum interpositum cyst in twins of a mother carrying May-Hegglin anomaly: a case report and short literature review. <i>BMC Pregnancy and Childbirth</i> , 2020 , 20, 772	3.2	
112	Novel AKAP9 mutation and long QT syndrome in a patient with torsades des pointes. <i>Journal of Interventional Cardiac Electrophysiology</i> , 2019 , 56, 171-172	2.4	1
111	Hereditary angioedema: Looking for bradykinin production and triggers of vascular permeability. <i>Clinical and Experimental Allergy</i> , 2019 , 49, 1395-1402	4.1	8
110	Angiopoietin-1 haploinsufficiency affects the endothelial barrier and causes hereditary angioedema. <i>Clinical and Experimental Allergy</i> , 2019 , 49, 626-635	4.1	21
109	Antiphospholipid Antibodies in a General Obstetric Population: Clinical Impact on Pregnancy Outcome and Relationship with the M2 Haplotype in the Annexin A5 (ANXA5) Gene. <i>Hamostaseologie</i> , 2019 , 39, 203-207	1.9	
108	The role of genetics in the current diagnostic workup of idiopathic non-histaminergic angioedema. <i>Allergy: European Journal of Allergy and Clinical Immunology</i> , 2019 , 74, 810-812	9.3	9
107	Double de novo mutations in dilated cardiomyopathy with cardiac arrest. <i>Journal of Electrocardiology</i> , 2019 , 53, 40-43	1.4	5
106	Maladaptive Coping Strategies and Neuroticism Mediate the Relationship Between 5HTT-LPR Polymorphisms and Symptoms of Anxiety in Elite Athletes. 2019 , 16, 62-71		1
105	A novel allele variant of the SERPINF2 gene responsible for severe plasmin inhibitor (Eantiplasmin) deficiency in an Italian patient. <i>Thrombosis Research</i> , 2018 , 166, 60-62	8.2	6
104	Mutation of the angiopoietin-1 gene (ANGPT1) associates with a new type of hereditary angioedema. <i>Journal of Allergy and Clinical Immunology</i> , 2018 , 141, 1009-1017	11.5	150
103	Human Fibrinogen: Molecular and Genetic Aspects of Congenital Disorders. <i>International Journal of Molecular Sciences</i> , 2018 , 19,	6.3	31
102	Preliminary Data From the Study of Coagulative Profile of HIV Infected Individuals Suggest a Role For Point Mutations in the Gene in Protein S Deficiency in Individuals Undergoing Highly Antiretroviral Therapy. <i>Open AIDS Journal</i> , 2018 , 12, 6-10	0.6	3

101	Venous thromboembolism in assisted reproductive technologies: comparison between unsuccessful versus successful cycles in an Italian cohort. <i>Journal of Thrombosis and Thrombolysis</i> , 2018 , 45, 234-239	5.1	7
100	Venous Thromboembolism in Women Undergoing Assisted Reproductive Technologies: Data from the RIETE Registry. <i>Thrombosis and Haemostasis</i> , 2018 , 118, 1962-1968	7	27
99	An Italian Registry on Risk Factors for Venous Thromboembolism in Blood Donors Clinicaltrials. Gov: Nct03282747. <i>Proceedings (mdpi)</i> , 2018 , 2, 532	0.3	
98	Genetic Risk Factors and Inhibitor Development in Hemophilia: What Is Known and Searching for the Unknown. <i>Seminars in Thrombosis and Hemostasis</i> , 2018 , 44, 509-516	5.3	5
97	Factor VII deficiency: a novel missense variant and genotype-phenotype correlation in patients from Southern Italy. <i>Human Genome Variation</i> , 2017 , 4, 17048	1.8	3
96	Survival in primary antiphospholipid syndrome. A single-centre cohort study. <i>Thrombosis and Haemostasis</i> , 2016 , 115, 1200-8	7	9
95	The Relationship between Personality Traits, the 5HTT Polymorphisms, and the Occurrence of Anxiety and Depressive Symptoms in Elite Athletes. <i>PLoS ONE</i> , 2016 , 11, e0156601	3.7	45
94	Towards the genetic basis of cerebral venous thrombosis-the BEAST Consortium: a study protocol. <i>BMJ Open</i> , 2016 , 6, e012351	3	11
93	Incidental splanchnic vein thrombosis: preliminary registry data. <i>Lancet Haematology,the</i> , 2016 , 3, e256-74.6		
92	In vitro residual activity of phenylalanine hydroxylase variants and correlation with metabolic phenotypes in PKU. <i>Gene</i> , 2016 , 594, 138-143	3.8	11
91	ABCB1 SNP rs4148738 modulation of apixaban interindividual variability. <i>Thrombosis Research</i> , 2016 , 145, 24-6	8.2	23
90	Modulation of factors involved in placental haemostasis and angiogenesis by low-molecular-weight-heparins. <i>Archives of Gynecology and Obstetrics</i> , 2016 , 294, 1323-1329	2.5	2
89	Pharmacogenetics of dabigatran etexilate interindividual variability. <i>Thrombosis Research</i> , 2016 , 144, 1-5	8.2	45
88	Characterization of patients with angioedema without wheals: the importance of F12 gene screening. <i>Clinical Immunology</i> , 2015 , 157, 239-48	9	27
87	Chronic thromboembolic pulmonary hypertension. <i>Netherlands Heart Journal</i> , 2015 , 23, 193	2.2	2
86	Phenylalanine hydroxylase deficiency in south Italy: Genotype-phenotype correlations, identification of a novel mutant PAH allele and prediction of BH4 responsiveness. <i>Clinica Chimica Acta</i> , 2015 , 450, 51-5	6.2	9
85	Identification of ten novel mutations in factor VIII gene: A study of a cohort of 52 haemophilia A patients. <i>Thrombosis Research</i> , 2015 , 135, 1031-4	8.2	1
84	The first case of a small supernumerary marker chromosome derived from chromosome 10 in an adult woman with an apparently normal phenotype. <i>Systems Biology in Reproductive Medicine</i> , 2015 , 61, 398-402	2.9	1

83 Congenital Platelet Disorders **2015**, 761-772

82 Clinical utility of screening for CALR gene exon 9 mutations in patients with splanchnic venous thrombosis. *Thrombosis and Haemostasis*, **2015**, 113, 1381-2 7 10

81 Role of CYP2D6 Polymorphisms in the Outcome of Postoperative Pain Treatment. *Pain Medicine*, **2015**, 16, 2012-23 2.8 8

80 Pregnancy-related venous thrombosis: comparison between spontaneous and ART conception in an Italian cohort. *BMJ Open*, **2015**, 5, e008213 3 17

79 Assessment of the olfactory function in Italian patients with type 3 von Willebrand disease caused by a homozygous 253 Kb deletion involving VWF and TMEM16B/ANO2. *PLoS ONE*, **2015**, 10, e0116483 3.7 7

78 A novel congenital dysprothrombinemia leading to defective prothrombin maturation. *Thrombosis Research*, **2014**, 134, 1135-41 8.2 18

77 Mutational spectrum of the c1 inhibitor gene in a cohort of Italian patients with hereditary angioedema: description of nine novel mutations. *Annals of Human Genetics*, **2014**, 78, 73-82 2.2 31

76 Intra-familial discordant PKU phenotype explained by mutation analysis in three pedigrees. *Clinical Biochemistry*, **2014**, 47, 233-5 3.5 9

75 Reply to "triple or dual therapy for HCV-1 naive patients? Optimizing selection tools". *Journal of Hepatology*, **2014**, 61, 179-80 13.4

74 Clinical utility of antithrombotic prophylaxis in ART procedures: an Italian experience. *PLoS ONE*, **2014**, 9, e97604 3.7 6

73 Variation in genes encoding for interferon β and η in the prediction of HCV-1 treatment-induced viral clearance. *Liver International*, **2014**, 34, 1369-77 7.9 7

72 Perceived challenges and attitudes to regimen and product selection from Italian haemophilia treaters: the 2013 AICE survey. *Haemophilia*, **2014**, 20, e128-35 3.3 10

71 Identification of naïve HCV-1 patients with chronic hepatitis who may benefit from dual therapy with peg-interferon and ribavirin. *Journal of Hepatology*, **2014**, 60, 16-21 13.4 23

70 Outcome of patients with splanchnic venous thrombosis presenting without overt MPN: a role for the JAK2 V617F mutation re-evaluation. *Thrombosis Research*, **2013**, 132, e99-e104 8.2 25

69 Mutation analysis in hyperphenylalaninemia patients from South Italy. *Clinical Biochemistry*, **2013**, 46, 1896-8 3.5 12

68 De novo homozygous mutation of the C1 inhibitor gene in a patient with hereditary angioedema. *Journal of Allergy and Clinical Immunology*, **2013**, 132, 748-750.e3 11.5 22

67 Homozygosity by descent of a 3Mb chromosome 17 haplotype causes coinheritance of Glanzmann thrombasthenia and primary ciliary dyskinesia. *Blood*, **2013**, 122, 4289-91 2.2 6

66 Structural analysis of protein Z gene variants in patients with foetal losses. *Thrombosis and Haemostasis*, **2013**, 110, 534-42 7 3

65	Hereditary protein C deficiency and thrombosis risk: genotype and phenotype relation in a large Italian family. <i>European Journal of Haematology</i> , 2012 , 88, 336-9	3.8	3
64	Influence of the Gly1057Asp variant of the insulin receptor substrate 2 (IRS2) on insulin resistance and relationship with epicardial fat thickness in the elderly. <i>Experimental Gerontology</i> , 2012 , 47, 988-93	4.5	6
63	F8 gene mutation type and inhibitor development in patients with severe hemophilia A: systematic review and meta-analysis. <i>Blood</i> , 2012 , 119, 2922-34	2.2	248
62	A reliable and rapid tool for plasma quantification of 18 psychotropic drugs by ESI tandem mass spectrometry. <i>Journal of Pharmaceutical and Biomedical Analysis</i> , 2012 , 67-68, 104-13	3.5	15
61	Obstetric complications and pregnancy-related venous thromboembolism: the effect of low-molecular-weight heparin on their prevention in carriers of factor V Leiden or prothrombin G20210A mutation. <i>Thrombosis and Haemostasis</i> , 2012 , 107, 477-84	7	20
60	Fatal pulmonary thromboembolism. A retrospective autopsy study: searching for genetic thrombophilias (Factor V Leiden (G1691A) and FII (G20210A) gene variants) and dating the thrombus. <i>Forensic Science International</i> , 2012 , 214, 152-8	2.6	6
59	Increased warfarin consumption and residual fibrin turnover in thrombotic patients with primary antiphospholipid syndrome. <i>Thrombosis Research</i> , 2011 , 127, 595-9	8.2	9
58	Sex modulation of the occurrence of jak2 v617f mutation in patients with splanchnic venous thrombosis. <i>Thrombosis Research</i> , 2011 , 128, 233-6	8.2	15
57	The risk of occurrence of venous thrombosis: focus on protein Z. <i>Thrombosis Research</i> , 2011 , 128, 508-158.2		17
56	Gene polymorphisms and sport attitude in Italian athletes. <i>Genetic Testing and Molecular Biomarkers</i> , 2011 , 15, 285-90	1.6	44
55	Population genetics of venous thromboembolism. A narrative review. <i>Thrombosis and Haemostasis</i> , 2011 , 105, 221-31	7	56
54	McKusick-Kaufman or Bardet-Biedl syndrome? A new borderline case in an Italian nonconsanguineous healthy family. <i>Indian Journal of Human Genetics</i> , 2011 , 17, 94-6		3
53	Eosinophilia and thrombophilia in churg strauss syndrome: a clinical and pathogenetic overview. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2010 , 16, 628-36	3.3	53
52	The haplotype M2 within the ANXA5 gene is independently associated with the occurrence of deep venous thrombosis. <i>Thrombosis and Haemostasis</i> , 2010 , 103, 1102-3	7	8
51	Markers of haemostasis and angiogenesis in placentae from gestational vascular complications: impairment of mechanisms involved in maintaining intervillous blood flow. <i>Thrombosis Research</i> , 2010 , 125, 267-71	8.2	10
50	Role of the M2 haplotype within the annexin A5 gene in the occurrence of pregnancy-related venous thromboembolism. <i>American Journal of Obstetrics and Gynecology</i> , 2010 , 203, 461.e1-5	6.4	21
49	New TET2 gene mutations in patients with myeloproliferative neoplasms and splanchnic vein thrombosis. <i>Journal of Thrombosis and Haemostasis</i> , 2010 , 8, 1142-4	15.4	8
48	Detection of new deletions in a group of Italian patients with Hemophilia A by multiplex ligation-dependent probe amplification. <i>Genetic Testing and Molecular Biomarkers</i> , 2009 , 13, 573-6	1.6	5

47	Role of thrombophilia in adverse obstetric outcomes and their prevention using antithrombotic therapy. <i>Seminars in Thrombosis and Hemostasis</i> , 2009 , 35, 630-43	5.3	14
46	A platelet defect modulates bleeding in mild hemophilia: the tale of 2 brothers. <i>Clinical and Applied Thrombosis/Hemostasis</i> , 2009 , 15, 715-6	3.3	
45	Prevention and treatment of bleeding complications in patients receiving vitamin K antagonists, Part 1: Prevention. <i>American Journal of Hematology</i> , 2009 , 84, 579-83	7.1	12
44	Setting up a 2D-LC/MS/MS method for the rapid quantitation of the prostanoid metabolites 6-oxo-PGF(1alpha) and TXB2 as markers for hemostasis assessment. <i>Journal of Mass Spectrometry</i> , 2009 , 44, 346-52	2.2	11
43	Stable-isotope dilution LC-ESI-MS/MS techniques for the quantification of total homocysteine in human plasma. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2009 , 877, 3292-9	3.2	10
42	Annexin V expression in human placenta is influenced by the carriership of the common haplotype M2. <i>Fertility and Sterility</i> , 2009 , 91, 940-2	4.8	40
41	Protein Z g-42a variant and the risk of pregnancy-related venous thromboembolism in a cohort of Italian patients. <i>Thrombosis Research</i> , 2009 , 123, 848-50	8.2	8
40	Coexistence of beta-thalassemia and hereditary hemochromatosis in homozygosity: a possible synergic effect?. <i>Hemoglobin</i> , 2009 , 33, 155-7	0.6	3
39	Methylarginines and mortality in patients with end stage renal disease: a prospective cohort study. <i>Atherosclerosis</i> , 2009 , 207, 541-5	3.1	52
38	Haplotype M2 in the annexin A5 (ANXA5) gene and the occurrence of obstetric complications. <i>Thrombosis and Haemostasis</i> , 2009 , 102, 309-13	7	59
37	Inherited platelet disorders: thrombocytopenias and thrombocytopathies. <i>Blood Transfusion</i> , 2009 , 7, 278-92	3.6	34
36	Type of Factor VIII Product as Inhibitor Risk Factor in Patients with Severe Hemophilia A and Null Mutations.. <i>Blood</i> , 2009 , 114, 3500-3500	2.2	
35	Identification of GLA gene deletions in Fabry patients by Multiplex Ligation-dependent Probe Amplification (MLPA). <i>Molecular Genetics and Metabolism</i> , 2008 , 94, 382-5	3.7	30
34	Correlation between factors involved in the local haemostasis and angiogenesis in full term human placenta. <i>Thrombosis Research</i> , 2008 , 122, 376-82	8.2	10
33	An unreported mutation within protein Z gene is associated with very low protein levels in women with fetal loss. <i>Fertility and Sterility</i> , 2008 , 90, 864-5	4.8	7
32	Occurrence of the JAK2 V617F mutation in the Budd-Chiari syndrome. <i>Blood Coagulation and Fibrinolysis</i> , 2008 , 19, 459-62	1	20
31	The Italian AICE-Genetics hemophilia A database: results and correlation with clinical phenotype. <i>Haematologica</i> , 2008 , 93, 722-8	6.6	73
30	A beta3 Asp217-->Val substitution in a patient with variant Glanzmann Thrombasthenia severely affects integrin alphaIIb beta3 functions. <i>Blood Coagulation and Fibrinolysis</i> , 2008 , 19, 657-62	1	4

29	Janus kinase-2 mutation, cirrhosis and splanchnic vein thrombosis. <i>European Journal of Gastroenterology and Hepatology</i> , 2008 , 20, 245-6	2.2	8
28	Screening of mutations of hemophilia A in 40 Italian patients: a novel G-to-A mutation in intron 10 of the F8 gene as a putative cause of mild hemophilia A in southern Italy. <i>Blood Coagulation and Fibrinolysis</i> , 2008 , 19, 197-202	1	20
27	Antithrombotic prophylaxis during pregnancy in women with deficiency of natural anticoagulants. <i>Blood Coagulation and Fibrinolysis</i> , 2008 , 19, 226-30	1	22
26	Venous thrombosis in oral contraceptive users and the presence of the JAK2 V617F mutation. <i>Thrombosis and Haemostasis</i> , 2008 , 99, 640-2	7	10
25	How to handle low-molecular-weight heparins in patients with decreased renal function: an open issue. <i>Internal and Emergency Medicine</i> , 2008 , 3, 307-9	3.7	1
24	Oral anticoagulants: Pharmacogenetics Relationship between genetic and non-genetic factors. <i>Blood Reviews</i> , 2008 , 22, 127-40	11.1	48
23	Fetal sex identification in maternal plasma by means of short tandem repeats on chromosome x. <i>Annals of the New York Academy of Sciences</i> , 2008 , 1137, 148-56	6.5	12
22	A rapid method for the quantification of the enantiomers of Warfarin, Phenprocoumon and Acenocoumarol by two-dimensional-enantioselective liquid chromatography/electrospray tandem mass spectrometry. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2007 , 852, 107-14	3.2	22
21	A novel mutation in human ether-a-go-go-related gene, alanine to proline at position 490, found in a large family with autosomal dominant long QT syndrome. <i>American Journal of Cardiology</i> , 2007 , 99, 1737-40	3	4
20	Prognostic factors in noncirrhotic patients with splanchnic vein thromboses. <i>American Journal of Gastroenterology</i> , 2007 , 102, 2464-70	0.7	218
19	Gain-of-function gene mutations and venous thromboembolism: distinct roles in different clinical settings. <i>Journal of Medical Genetics</i> , 2007 , 44, 412-6	5.8	37
18	A new JAK2 gene mutation in patients with polycythemia vera and splanchnic vein thrombosis. <i>Blood</i> , 2007 , 110, 2768-9	2.2	42
17	A new case of combined factor V and factor VIII deficiency further suggests that the LMAN1 M1T mutation is a frequent cause in Italian patients. <i>Blood Coagulation and Fibrinolysis</i> , 2007 , 18, 203-4	1	7
16	Insight into the nature of the CRP-coronary event association using Mendelian randomization. <i>International Journal of Epidemiology</i> , 2006 , 35, 922-31	7.8	137
15	Adverse outcome in women with thrombophilia and bilateral uterine artery notches. <i>Fertility and Sterility</i> , 2006 , 86, 726-7	4.8	2
14	Homocysteine levels in amniotic fluid. <i>Thrombosis and Haemostasis</i> , 2006 , 95, 625-628	7	11
13	Increased plasma prothrombin concentration in cirrhotic patients with portal vein thrombosis and prothrombin G20210A mutation. <i>Thrombosis and Haemostasis</i> , 2006 , 95, 221-3	7	17
12	The COX-2 G/C -765 polymorphism may modulate the occurrence of cerebrovascular ischemia. <i>Blood Coagulation and Fibrinolysis</i> , 2006 , 17, 93-6	1	37

11	Inherited abnormalities of fibrinogen: 10-year clinical experience of an Italian group. <i>Blood Coagulation and Fibrinolysis</i> , 2006 , 17, 235-40	1	20
10	Homocysteine metabolism in families from southern Italy with neural tube defects: role of genetic and nutritional determinants. <i>Prenatal Diagnosis</i> , 2006 , 26, 1-5	3.2	28
9	A new method for determination of plasma homocysteine by isotope dilution and electrospray tandem mass spectrometry. <i>Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences</i> , 2006 , 842, 64-9	3.2	11
8	The C677T methylenetetrahydrofolate reductase gene mutation does not influence cardiovascular risk in the dialysis population: results of a multicentre prospective study. <i>Nephrology Dialysis Transplantation</i> , 2005 , 20, 382-6	4.3	20
7	A polymorphism in the VKORC1 gene is associated with an interindividual variability in the dose-anticoagulant effect of warfarin. <i>Blood</i> , 2005 , 105, 645-9	2.2	577
6	Molecular diversity and thrombotic risk in protein S deficiency: the PROSIT study. <i>Human Mutation</i> , 2005 , 25, 259-69	4.7	40
5	Congenital nephrotic syndrome of Finnish type: detection of new nephrin mutations and prenatal diagnosis in an Italian family. <i>Prenatal Diagnosis</i> , 2005 , 25, 407-10	3.2	12
4	Bleeding and re-thrombosis in primary antiphospholipid syndrome on oral anticoagulation: an 8-year longitudinal comparison with mitral valve replacement and inherited thrombophilia. <i>Thrombosis and Haemostasis</i> , 2005 , 93, 694-9	7	35
3	A Frameshift Mutation in the Human Fibrinogen A α chain Gene (A[499] Ala Frameshift Stop) Leading to Dysfibrinogen San Giovanni Rotondo. <i>Thrombosis and Haemostasis</i> , 2001 , 86, 1483-1488	7	14
2	A G-to-A mutation in IVS-3 of the human gamma fibrinogen gene causing afibrinogenemia due to abnormal RNA splicing. <i>Blood</i> , 2000 , 96, 2501-2505	2.2	32
1	Congenital Platelet Disorders915-927		