Maurizio Margaglione

List of Publications by Citations

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	A polymorphism in the VKORC1 gene is associated with an interindividual variability in the dose-anticoagulant effect of warfarin. *Blood*, **2005**, 105, 645-9	2.2	577
135	F8 gene mutation type and inhibitor development in patients with severe hemophilia A: systematic review and meta-analysis. *Blood*, **2012**, 119, 2922-34	2.2	248
134	Prognostic factors in noncirrhotic patients with splanchnic vein thromboses. *American Journal of Gastroenterology*, **2007**, 102, 2464-70	0.7	218
133	Mutation of the angiopoietin-1 gene (ANGPT1) associates with a new type of hereditary angioedema. *Journal of Allergy and Clinical Immunology*, **2018**, 141, 1009-1017	11.5	150
132	Insight into the nature of the CRP-coronary event association using Mendelian randomization. *International Journal of Epidemiology*, **2006**, 35, 922-31	7.8	137
131	The Italian AICE-Genetics hemophilia A database: results and correlation with clinical phenotype. *Haematologica*, **2008**, 93, 722-8	6.6	73
130	Haplotype M2 in the annexin A5 (ANXA5) gene and the occurrence of obstetric complications. *Thrombosis and Haemostasis*, **2009**, 102, 309-13	7	59
129	Population genetics of venous thromboembolism. A narrative review. *Thrombosis and Haemostasis*, **2011**, 105, 221-31	7	56
128	Eosinophilia and thrombophilia in churg strauss syndrome: a clinical and pathogenetic overview. *Clinical and Applied Thrombosis/Hemostasis*, **2010**, 16, 628-36	3.3	53
127	Methylarginines and mortality in patients with end stage renal disease: a prospective cohort study. *Atherosclerosis*, **2009**, 207, 541-5	3.1	52
126	Oral anticoagulants: Pharmacogenetics Relationship between genetic and non-genetic factors. *Blood Reviews*, **2008**, 22, 127-40	11.1	48
125	The Relationship between Personality Traits, the 5HTT Polymorphisms, and the Occurrence of Anxiety and Depressive Symptoms in Elite Athletes. *PLoS ONE*, **2016**, 11, e0156601	3.7	45
124	Pharmacogenetics of dabigatran etexilate interindividual variability. *Thrombosis Research*, **2016**, 144, 1-5	8.2	45
123	Gene polymorphisms and sport attitude in Italian athletes. *Genetic Testing and Molecular Biomarkers*, **2011**, 15, 285-90	1.6	44
122	A new JAK2 gene mutation in patients with polycythemia vera and splanchnic vein thrombosis. *Blood*, **2007**, 110, 2768-9	2.2	42
121	Annexin V expression in human placenta is influenced by the carriership of the common haplotype M2. *Fertility and Sterility*, **2009**, 91, 940-2	4.8	40
120	Molecular diversity and thrombotic risk in protein S deficiency: the PROSIT study. *Human Mutation*, **2005**, 25, 259-69	4.7	40

119	Gain-of-function gene mutations and venous thromboembolism: distinct roles in different clinical settings. *Journal of Medical Genetics*, **2007**, 44, 412-6	5.8	37
118	The COX-2 G/C -765 polymorphism may modulate the occurrence of cerebrovascular ischemia. *Blood Coagulation and Fibrinolysis*, **2006**, 17, 93-6	1	37
117	Bleeding and re-thrombosis in primary antiphospholipid syndrome on oral anticoagulation: an 8-year longitudinal comparison with mitral valve replacement and inherited thrombophilia. *Thrombosis and Haemostasis*, **2005**, 93, 694-9	7	35
116	A myoferlin gain-of-function variant associates with a new type of hereditary angioedema. *Allergy: European Journal of Allergy and Clinical Immunology*, **2020**, 75, 2989-2992	9.3	34
115	Inherited platelet disorders: thrombocytopenias and thrombocytopathies. *Blood Transfusion*, **2009**, 7, 278-92	3.6	34
114	A G-to-A mutation in IVS-3 of the human gamma fibrinogen gene causing afibrinogenemia due to abnormal RNA splicing. *Blood*, **2000**, 96, 2501-2505	2.2	32
113	Human Fibrinogen: Molecular and Genetic Aspects of Congenital Disorders. *International Journal of Molecular Sciences*, **2018**, 19,	6.3	31
112	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
111	Identification of GLA gene deletions in Fabry patients by Multiplex Ligation-dependent Probe Amplification (MLPA). *Molecular Genetics and Metabolism*, **2008**, 94, 382-5	3.7	30
110	Homocysteine metabolism in families from southern Italy with neural tube defects: role of genetic and nutritional determinants. *Prenatal Diagnosis*, **2006**, 26, 1-5	3.2	28
109	International Consensus on the Use of Genetics in the Management of Hereditary Angioedema. *Journal of Allergy and Clinical Immunology: in Practice*, **2020**, 8, 901-911	5.4	28
108	Characterization of patients with angioedema without wheals: the importance of F12 gene screening. *Clinical Immunology*, **2015**, 157, 239-48	9	27
107	Venous Thromboembolism in Women Undergoing Assisted Reproductive Technologies: Data from the RIETE Registry. *Thrombosis and Haemostasis*, **2018**, 118, 1962-1968	7	27
106	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
105	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
104	ABCB1 SNP rs4148738 modulation of apixaban interindividual variability. *Thrombosis Research*, **2016**, 145, 24-6	8.2	23
103	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
102	Antithrombotic prophylaxis during pregnancy in women with deficiency of natural anticoagulants. *Blood Coagulation and Fibrinolysis*, **2008**, 19, 226-30	1	22

101	A rapid method for the quantification of the enantiomers of Warfarin, Phenprocoumon and Acenocoumarol by two-dimensional-enantioselective liquid chromatography/electrospray tandem mass spectrometry. *Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences*, **2007**, 850, 507-14	3.2	22
100	Role of the M2 haplotype within the annexin A5 gene in the occurrence of pregnancy-related venous thromboembolism. *American Journal of Obstetrics and Gynecology*, **2010**, 203, 461.e1-5	6.4	21
99	Angiopoietin-1 haploinsufficiency affects the endothelial barrier and causes hereditary angioedema. *Clinical and Experimental Allergy*, **2019**, 49, 626-635	4.1	21
98	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. *Thrombosis and Haemostasis*, **2012**, 107, 477-84	7	20
97	Occurrence of the JAK2 V617F mutation in the Budd-Chiari syndrome. *Blood Coagulation and Fibrinolysis*, **2008**, 19, 459-62	1	20
96	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. *Blood Coagulation and Fibrinolysis*, **2008**, 19, 197-202	1	20
95	The C677T methylenetetrahydrofolate reductase gene mutation does not influence cardiovascular risk in the dialysis population: results of a multicentre prospective study. *Nephrology Dialysis Transplantation*, **2005**, 20, 382-6	4.3	20
94	Inherited abnormalities of fibrinogen: 10-year clinical experience of an Italian group. *Blood Coagulation and Fibrinolysis*, **2006**, 17, 235-40	1	20
93	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. *Journal of Medical Virology*, **2021**, 93, 1739-1742	19.7	19
92	A novel congenital dysprothrombinemia leading to defective prothrombin maturation. *Thrombosis Research*, **2014**, 134, 1135-41	8.2	18
91	Summary of the Available Molecular Methods for Detection of SARS-CoV-2 during the Ongoing Pandemic. *International Journal of Molecular Sciences*, **2021**, 22,	6.3	18
90	Pregnancy-related venous thrombosis: comparison between spontaneous and ART conception in an Italian cohort. *BMJ Open*, **2015**, 5, e008213	3	17
89	The risk of occurrence of venous thrombosis: focus on protein Z. *Thrombosis Research*, **2011**, 128, 508-15	8.2	17
88	Increased plasma prothrombin concentration in cirrhotic patients with portal vein thrombosis and prothrombin G20210A mutation. *Thrombosis and Haemostasis*, **2006**, 95, 221-3	7	17
87	A reliable and rapid tool for plasma quantification of 18 psychotropic drugs by ESI tandem mass spectrometry. *Journal of Pharmaceutical and Biomedical Analysis*, **2012**, 67-68, 104-13	3.5	15
86	Sex modulation of the occurrence of jak2 v617f mutation in patients with splanchnic venous thrombosis. *Thrombosis Research*, **2011**, 128, 233-6	8.2	15
85	Role of thrombophilia in adverse obstetric outcomes and their prevention using antithrombotic therapy. *Seminars in Thrombosis and Hemostasis*, **2009**, 35, 630-43	5.3	14
84	A Frameshift Mutation in the Human Fibrinogen A□chain Gene (A□[499] Ala Frameshift Stop) Leading to Dysfibrinogen San Giovanni Rotondo. *Thrombosis and Haemostasis*, **2001**, 86, 1483-1488	7	14

83	Impaired control of the contact system in hereditary angioedema with normal C1-inhibitor. *Allergy: European Journal of Allergy and Clinical Immunology*, **2020**, 75, 1394-1403	9.3	14
82	Mutation analysis in hyperphenylalaninemia patients from South Italy. *Clinical Biochemistry*, **2013**, 46, 1896-8	3.5	12
81	Prevention and treatment of bleeding complications in patients receiving vitamin K antagonists, Part 1: Prevention. *American Journal of Hematology*, **2009**, 84, 579-83	7.1	12
80	Fetal sex identification in maternal plasma by means of short tandem repeats on chromosome x. *Annals of the New York Academy of Sciences*, **2008**, 1137, 148-56	6.5	12
79	Congenital nephrotic syndrome of Finnish type: detection of new nephrin mutations and prenatal diagnosis in an Italian family. *Prenatal Diagnosis*, **2005**, 25, 407-10	3.2	12
78	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. *Journal of Mass Spectrometry*, **2009**, 44, 346-52	2.2	11
77	Homocysteine levels in amniotic fluid. *Thrombosis and Haemostasis*, **2006**, 95, 625-628	7	11
76	A new method for determination of plasma homocystine by isotope dilution and electrospray tandem mass spectrometry. *Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences*, **2006**, 842, 64-9	3.2	11
75	Towards the genetic basis of cerebral venous thrombosis-the BEAST Consortium: a study protocol. *BMJ Open*, **2016**, 6, e012351	3	11
74	In vitro residual activity of phenylalanine hydroxylase variants and correlation with metabolic phenotypes in PKU. *Gene*, **2016**, 594, 138-143	3.8	11
73	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
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	Markers of haemostasis and angiogenesis in placentae from gestational vascular complications: impairment of mechanisms involved in maintaining intervillous blood flow. *Thrombosis Research*, **2010**, 125, 267-71	8.2	10
70	Stable-isotope dilution LC-ESI-MS/MS techniques for the quantification of total homocysteine in human plasma. *Journal of Chromatography B: Analytical Technologies in the Biomedical and Life Sciences*, **2009**, 877, 3292-9	3.2	10
69	Correlation between factors involved in the local haemostasis and angiogenesis in full term human placenta. *Thrombosis Research*, **2008**, 122, 376-82	8.2	10
68	Venous thrombosis in oral contraceptive users and the presence of the JAK2 V617F mutation. *Thrombosis and Haemostasis*, **2008**, 99, 640-2	7	10
67	Phenylalanine hydroxylase deficiency in south Italy: Genotype-phenotype correlations, identification of a novel mutant PAH allele and prediction of BH4 responsiveness. *Clinica Chimica Acta*, **2015**, 450, 51-5	6.2	9
66	Survival in primary antiphospholipid syndrome. A single-centre cohort study. *Thrombosis and Haemostasis*, **2016**, 115, 1200-8	7	9

65	Intra-familiar discordant PKU phenotype explained by mutation analysis in three pedigrees. *Clinical Biochemistry*, **2014**, 47, 233-5	3.5	9
64	Increased warfarin consumption and residual fibrin turnover in thrombotic patients with primary antiphospholipid syndrome. *Thrombosis Research*, **2011**, 127, 595-9	8.2	9
63	The role of genetics in the current diagnostic workup of idiopathic non-histaminergic angioedema. *Allergy: European Journal of Allergy and Clinical Immunology*, **2019**, 74, 810-812	9.3	9
62	Hereditary angioedema: Looking for bradykinin production and triggers of vascular permeability. *Clinical and Experimental Allergy*, **2019**, 49, 1395-1402	4.1	8
61	Role of CYP2D6 Polymorphisms in the Outcome of Postoperative Pain Treatment. *Pain Medicine*, **2015**, 16, 2012-23	2.8	8
60	The haplotype M2 within the ANXA5 gene is independently associated with the occurrence of deep venous thrombosis. *Thrombosis and Haemostasis*, **2010**, 103, 1102-3	7	8
59	Protein Z g-42a variant and the risk of pregnancy-related venous thromboembolism in a cohort of Italian patients. *Thrombosis Research*, **2009**, 123, 848-50	8.2	8
58	New TET2 gene mutations in patients with myeloproliferative neoplasms and splanchnic vein thrombosis. *Journal of Thrombosis and Haemostasis*, **2010**, 8, 1142-4	15.4	8
57	Janus kinase-2 mutation, cirrhosis and splanchnic vein thrombosis. *European Journal of Gastroenterology and Hepatology*, **2008**, 20, 245-6	2.2	8
56	The Genetics of Hereditary Angioedema: A Review. *Journal of Clinical Medicine*, **2021**, 10,	5.1	8
55	Variation in genes encoding for interferon □B and □4 in the prediction of HCV-1 treatment-induced viral clearance. *Liver International*, **2014**, 34, 1369-77	7.9	7
54	An unreported mutation within protein Z gene is associated with very low protein levels in women with fetal loss. *Fertility and Sterility*, **2008**, 90, 864-5	4.8	7
53	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. *Blood Coagulation and Fibrinolysis*, **2007**, 18, 203-4	1	7
52	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
51	Venous thromboembolism in assisted reproductive technologies: comparison between unsuccessful versus successful cycles in an Italian cohort. *Journal of Thrombosis and Thrombolysis*, **2018**, 45, 234-239	5.1	7
50	A novel allele variant of the SERPINF2 gene responsible for severe plasmin inhibitor (□antiplasmin) deficiency in an Italian patient. *Thrombosis Research*, **2018**, 166, 60-62	8.2	6
49	Clinical utility of antithrombotic prophylaxis in ART procedures: an Italian experience. *PLoS ONE*, **2014**, 9, e97604	3.7	6
48	Influence of the Gly1057Asp variant of the insulin receptor substrate 2 (IRS2) on insulin resistance and relationship with epicardial fat thickness in the elderly. *Experimental Gerontology*, **2012**, 47, 988-93	4.5	6

47	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
46	Fatal pulmonary thromboembolism. A retrospective autopsy study: searching for genetic thrombophilias (Factor V Leiden (G1691A) and FII (G20210A) gene variants) and dating the thrombus. *Forensic Science International*, **2012**, 214, 152-8	2.6	6
45	Detection of new deletions in a group of Italian patients with Hemophilia A by multiplex ligation-dependent probe amplification. *Genetic Testing and Molecular Biomarkers*, **2009**, 13, 573-6	1.6	5
44	Anticoagulation in Italian patients with venous thromboembolism and thrombophilic alterations: findings from START2 register study. *Blood Transfusion*, **2020**, 18, 486-495	3.6	5
43	Findings from a multicentre, observational study on reproductive outcomes in women with unexplained recurrent pregnancy loss: the OTTILIA registry. *Human Reproduction*, **2021**, 36, 2083-2090	5.7	5
42	Double de novo mutations in dilated cardiomyopathy with cardiac arrest. *Journal of Electrocardiology*, **2019**, 53, 40-43	1.4	5
41	Genetic Risk Factors and Inhibitor Development in Hemophilia: What Is Known and Searching for the Unknown. *Seminars in Thrombosis and Hemostasis*, **2018**, 44, 509-516	5.3	5
40	A beta3 Asp217-->Val substitution in a patient with variant Glanzmann Thrombasthenia severely affects integrin alphaIIBbeta3 functions. *Blood Coagulation and Fibrinolysis*, **2008**, 19, 657-62	1	4
39	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. *American Journal of Cardiology*, **2007**, 99, 1737-40	3	4
38	Genome-Wide Association Study Identifies First Locus Associated with Susceptibility to Cerebral Venous Thrombosis. *Annals of Neurology*, **2021**, 90, 777-788	9.4	4
37	Low Prevalence of Antibodies to SARS-CoV-2 and Undetectable Viral Load in Seropositive Blood Donors from South-Eastern Italy. *Acta Haematologica*, **2021**, 144, 580-584	2.7	4
36	Thromboelastography parameters in Italian pregnant women: do antithrombotic drugs change reference values?. *Journal of Investigative Medicine*, **2020**, 68, 902-905	2.9	3
35	Hereditary protein C deficiency and thrombosis risk: genotype and phenotype relation in a large Italian family. *European Journal of Haematology*, **2012**, 88, 336-9	3.8	3
34	Factor VII deficiency: a novel missense variant and genotype-phenotype correlation in patients from Southern Italy. *Human Genome Variation*, **2017**, 4, 17048	1.8	3
33	Structural analysis of protein Z gene variants in patients with foetal losses. *Thrombosis and Haemostasis*, **2013**, 110, 534-42	7	3
32	McKusick-Kaufman or Bardet-Biedl syndrome? A new borderline case in an Italian nonconsanguineous healthy family. *Indian Journal of Human Genetics*, **2011**, 17, 94-6		3
31	Coexistence of beta-thalassemia and hereditary hemochromatosis in homozygosity: a possible synergic effect?. *Hemoglobin*, **2009**, 33, 155-7	0.6	3
30	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. *Open AIDS Journal*, **2018**, 12, 6-10	0.6	3

29	Mortality and Transfusion Requirements in COVID-19 Hospitalized Italian Patients According to Severity of the Disease. *Journal of Clinical Medicine*, **2021**, 10,	5.1	3
28	Chronic thromboembolic pulmonary hypertension. *Netherlands Heart Journal*, **2015**, 23, 193	2.2	2
27	Severe bleeding and absent ADP-induced platelet aggregation associated with inherited combined CalDAG-GEFI and P2Y deficiencies. *Haematologica*, **2020**, 105, e361-e364	6.6	2
26	Adverse outcome in women with thrombophilia and bilateral uterine artery notches. *Fertility and Sterility*, **2006**, 86, 726-7	4.8	2
25	Use of low-molecular weight heparin, transfusion and mortality in COVID-19 patients not requiring ventilation. *Journal of Thrombosis and Thrombolysis*, **2021**, 52, 772-778	5.1	2
24	Modulation of factors involved in placental haemostasis and angiogenesis by low-molecular-weight-heparins. *Archives of Gynecology and Obstetrics*, **2016**, 294, 1323-1329	2.5	2
23	Obstetric outcomes in pregnant COVID-19 women: the imbalance of von Willebrand factor and ADAMTS13 axis.. *BMC Pregnancy and Childbirth*, **2022**, 22, 142	3.2	2
22	Identification of ten novel mutations in factor VIII gene: A study of a cohort of 52 haemophilia A patients. *Thrombosis Research*, **2015**, 135, 1031-4	8.2	1
21	The first case of a small supernumerary marker chromosome derived from chromosome 10 in an adult woman with an apparently normal phenotype. *Systems Biology in Reproductive Medicine*, **2015**, 61, 398-402	2.9	1
20	Novel AKAP9 mutation and long QT syndrome in a patient with torsades des pointes. *Journal of Interventional Cardiac Electrophysiology*, **2019**, 56, 171-172	2.4	1
19	How to handle low-molecular-weight heparins in patients with decreased renal function: an open issue. *Internal and Emergency Medicine*, **2008**, 3, 307-9	3.7	1
18	Italian experience with rVIII-single chain: a survey of patients with haemophilia A and their physicians. *Journal of Thrombosis and Thrombolysis*, **2021**, 1	5.1	1
17	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. *Infectious Diseases*, **2021**, 1-6	3.1	1
16	Homozygous methylentetrahydrofolate reductase C667T genotype anticipates age at venous thromboembolism by one decade. *Blood Coagulation and Fibrinolysis*, **2021**, 32, 382-386	1	1
15	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). *Genes*, **2021**, 12,	4.2	1
14	Severe systemic thrombosis in a young COVID-19 patient with a rare homozygous prothrombin G20210A mutation. *Infezioni in Medicina*, **2021**, 29, 259-262	3.6	1
13	Maladaptive Coping Strategies and Neuroticism Mediate the Relationship Between 5HTT-LPR Polymorphisms and Symptoms of Anxiety in Elite Athletes. **2019**, 16, 62-71		1
12	Psychosomatic syndromes are associated with IL-6 pro-inflammatory cytokine in heart failure patients.. *PLoS ONE*, **2022**, 17, e0265282	3.7	0

11	Anemone study: prevalence of risk factors for superficial vein thrombosis in a large Italian population of blood donors. *Journal of Thrombosis and Thrombolysis*, **2020**, 50, 689-696	5.1
10	Reply to "triple or dual therapy for HCV-1 naive patients? Optimizing selection tools". *Journal of Hepatology*, **2014**, 61, 179-80	13.4
9	Congenital Platelet Disorders **2015**, 761-772	
8	A platelet defect modulates bleeding in mild hemophilia: the tale of 2 brothers. *Clinical and Applied Thrombosis/Hemostasis*, **2009**, 15, 715-6	3.3
7	Mortality and clinical outcome of Italian patients undergoing orthopaedic surgery: effect of peri-operative blood transfusion. *Blood Transfusion*, **2021**, 19, 284-291	3.6
6	Type of Factor VIII Product as Inhibitor Risk Factor in Patients with Severe Hemophilia A and Null Mutations.. *Blood*, **2009**, 114, 3500-3500	2.2
5	Congenital Platelet Disorders915-927	
4	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. *BMC Pregnancy and Childbirth*, **2020**, 20, 772	3.2
3	Incidental splanchnic vein thrombosis: preliminary registry data. *Lancet Haematology,the*, **2016**, 3, e256-7	14.6
2	Antiphospholipid Antibodies in a General Obstetric Population: Clinical Impact on Pregnancy Outcome and Relationship with the M2 Haplotype in the Annexin A5 (ANXA5) Gene. *Hamostaseologie*, **2019**, 39, 203-207	1.9
1	An Italian Registry on Risk Factors for Venous Thromboembolism in Blood Donors Clinicaltrials. Gov: Nct03282747. *Proceedings (mdpi)*, **2018**, 2, 532	0.3