Elvira Grandone

List of Publications by Year
in descending order

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281 papers	11,791 citations	31976 53 h-index	31849 101 g-index
290 all docs	290 docs citations	290 times ranked	13510 citing authors

#	Article	IF	Citations
1	Multiple common variants for celiac disease influencing immune gene expression. Nature Genetics, 2010, 42, 295-302.	21.4	871
2	A polymorphism in the VKORC1 gene is associated with an interindividual variability in the dose-anticoagulant effect of warfarin. Blood, 2005, 105, 645-649.	1.4	647
3	Aspirin for Preventing the Recurrence of Venous Thromboembolism. New England Journal of Medicine, 2012, 366, 1959-1967.	27.0	545
4	Risk factors and clinical presentation of portal vein thrombosis in patients with liver cirrhosis. Journal of Hepatology, 2004, 40, 736-741.	3.7	512
5	Demographic and clinical data in acquired hemophilia A: results from the European Acquired Haemophilia Registry (EACH2). Journal of Thrombosis and Haemostasis, 2012, 10, 622-631.	3.8	395
6	Management of bleeding in acquired hemophilia A: results from the European Acquired Haemophilia (EACH2) Registry. Blood, 2012, 120, 39-46.	1.4	326
7	Meta-Analysis of Genome-Wide Association Studies in Celiac Disease and Rheumatoid Arthritis Identifies Fourteen Non-HLA Shared Loci. PLoS Genetics, 2011, 7, e1002004.	3.5	307
8	Immunosuppression for acquired hemophilia A: results from the European Acquired Haemophilia Registry (EACH2). Blood, 2012, 120, 47-55.	1.4	284
9	Prognostic Factors in Noncirrhotic Patients With Splanchnic Vein Thromboses. American Journal of Gastroenterology, 2007, 102, 2464-2470.	0.4	254
10	Genetic Modulation of Oral Anticoagulation with Warfarin. Thrombosis and Haemostasis, 2000, 84, 775-778.	3.4	225
11	Increased Risk for Venous Thrombosis in Carriers of the Prothrombin Gâ†'A^20210 Gene Variant. Annals of Internal Medicine, 1998, 129, 89.	3.9	209
12	Factor V Leiden, C>T MTHFR Polymorphism and Genetic Susceptibility to Preeclampsia. Thrombosis and Haemostasis, 1997, 77, 1052-1054.	3.4	205
13	Factor V Leiden Is Associated with Repeated and Recurrent Unexplained Fetal Losses. Thrombosis and Haemostasis, 1997, 77, 0822-0824.	3.4	204
14	Long-term Clinical Outcomes of Splanchnic Vein Thrombosis. JAMA Internal Medicine, 2015, 175, 1474.	5.1	180
15	Genetic susceptibility to pregnancy-related venous thromboembolism: Roles of factor V Leiden, prothrombin G20210A, and methylenetetrahydrofolate reductase C677T mutations. American Journal of Obstetrics and Gynecology, 1998, 179, 1324-1328.	1.3	168
16	Genetic Susceptibility to Nonsteroidal Anti-Inflammatory Drugâ€"Related Gastroduodenal Bleeding: Role of Cytochrome P450 2C9 Polymorphisms. Gastroenterology, 2007, 133, 465-471.	1.3	161
17	The JAK2 V617F mutation frequently occurs in patients with portal and mesenteric venous thrombosis. Journal of Thrombosis and Haemostasis, 2007, 5, 55-61.	3.8	150
18	Abnormally high thromboxane biosynthesis in homozygous homocystinuria. Evidence for platelet involvement and probucol-sensitive mechanism.. Journal of Clinical Investigation, 1993, 92, 1400-1406.	8.2	141

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19	PAI-1 Plasma Levels in a General Population Without Clinical Evidence of Atherosclerosis. Arteriosclerosis, Thrombosis, and Vascular Biology, 1998, 18, 562-567.	2.4	140
20	The Methylenetetrahydrofolate Reductase TT677 Genotype Is Associated with Venous Thrombosis Independently of the Coexistence of the FV Leiden and the Prothrombin. Thrombosis and Haemostasis, 1998, 79, 907-911.	3.4	138
21	Longâ€term outcomes of patients with cerebral vein thrombosis: a multicenter study. Journal of Thrombosis and Haemostasis, 2012, 10, 1297-1302.	3.8	129
22	The PAI-1 Gene Locus 4G/5G Polymorphism Is Associated With a Family History of Coronary Artery Disease. Arteriosclerosis, Thrombosis, and Vascular Biology, 1998, 18, 152-156.	2.4	128
23	Coexistence of Factor V Leiden and Factor II A20210 Mutations and Recurrent Venous Thromboembolism. Thrombosis and Haemostasis, 1999, 82, 1583-1587.	3.4	123
24	Risk of Pregnancy-related Venous Thrombosis in Carriers of Severe Inherited Thrombophilia. Thrombosis and Haemostasis, 2001, 86, 800-803.	3.4	119
25	Inherited Prothrombotic Conditions and Premature Ischemic Stroke. Arteriosclerosis, Thrombosis, and Vascular Biology, 1999, 19, 1751-1756.	2.4	113
26	FactorÂVIII gene (F8) mutations as predictors of outcome in immune tolerance induction of hemophiliaÂA patients with highâ€responding inhibitors. Journal of Thrombosis and Haemostasis, 2009, 7, 1809-1815.	3.8	103
27	Inherited thrombophilia and in vitro fertilization implantation failure. Fertility and Sterility, 2001, 76, 201-202.	1.0	101
28	Abnormally high circulation levels of tissue plasminogen activator and plasminogen activator inhibitor-1 in patients with a history of ischemic stroke.. Arteriosclerosis and Thrombosis: A Journal of Vascular Biology, 1994, 14, 1741-1745.	3.9	96
29	The Italian AICE-Genetics hemophilia A database: results and correlation with clinical phenotype. Haematologica, 2008, 93, 722-728.	3.5	95
30	Preventing adverse obstetric outcomes in women with genetic thrombophilia. Fertility and Sterility, 2002, 78, 371-375.	1.0	90
31	Use of hydroxychloroquine in hospitalised COVID-19 patients is associated with reduced mortality: Findings from the observational multicentre Italian CORIST study. European Journal of Internal Medicine, 2020, 82, 38-47.	2.2	88
32	Heparin in COVID-19 Patients Is Associated with Reduced In-Hospital Mortality: The Multicenter Italian CORIST Study. Thrombosis and Haemostasis, 2021, 121, 1054-1065.	3.4	87
33	The Clinical Course of Venous Thromboembolism May Differ According to Cancer Site. American Journal of Medicine, 2017, 130, 337-347.	1.5	83
34	Prevalence of Apolipoprotein E Alleles in Healthy Subjects and Survivors of Ischemic Stroke. Stroke, 1998, 29, 399-403.	2.0	80
35	Inherited Thrombophilic Risk Factors and Venous Thromboembolism. Chest, 2000, 118, 1405-1411.	0.8	79
36	Anticardiolipin Antibodies in Patients With Liver Disease. American Journal of Gastroenterology, 1999, 94, 2983-2987.	0.4	78

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37	Pulmonary thrombosis in 2019â€nCoV pneumonia?. Journal of Thrombosis and Haemostasis, 2020, 18, 1511-1513.	3.8	75
38	Prothrombotic Genetic Risk Factors and the Occurrence of Gestational Hypertension with or without Proteinuria. Thrombosis and Haemostasis, 1999, 81, 349-352.	3.4	74
39	Plasminogen Activator Inhibitor-1 (PAI-1) Antigen Plasma Levels in Subjects Attending a Metabolic Ward: Relation to Polymorphisms of PAI-1 and Angiontensin Converting Enzyme (ACE) Genes. Arteriosclerosis, Thrombosis, and Vascular Biology, 1997, 17, 2082-2087.	2.4	69
40	Pharmacogenetics of dabigatran etexilate interindividual variability. Thrombosis Research, 2016, 144, 1-5.	1.7	69
41	An Alternative Method for PAI-1 Promoter Polymorphism (4G/5G) Typing. Thrombosis and Haemostasis, 1997, 77, 605-606.	3.4	69
42	Deletion Polymorphism in the Angiotensin-Converting Enzyme Gene in Patients With a History of Ischemic Stroke. Arteriosclerosis, Thrombosis, and Vascular Biology, 1996, 16, 304-309.	2.4	68
43	Genetic Polymorphism of 5,10-MTHFR Reductase Gene in Offspring of Patients with Myocardial Infarction. Thrombosis and Haemostasis, 1999, 82, 19-23.	3.4	65
44	Population genetics of venous thromboembolism. Thrombosis and Haemostasis, 2011, 105, 221-231.	3.4	64
45	Validation of the ISTH/SSC bleeding assessment tool for inherited platelet disorders: A communication from the Platelet Physiology SSC. Journal of Thrombosis and Haemostasis, 2020, 18, 732-739.	3.8	64
46	Haplotype M2 in the annexin A5 (ANXA5) gene and the occurrence of obstetric complications. Thrombosis and Haemostasis, 2009, 102, 309-313.	3.4	63
47	Glanzmannâ€™s Thrombasthenia: Identification of 19 New Mutations in 30 Patients. Thrombosis and Haemostasis, 2002, 87, 1034-1042.	3.4	62
48	Maternal thrombophilia and the risk of recurrence of preeclampsia. American Journal of Obstetrics and Gynecology, 2009, 200, 46.e1-46.e5.	1.3	62
49	Role of cytochrome P4502D6 functional polymorphisms in the efficacy of donepezil in patients with Alzheimer's disease. Pharmacogenetics and Genomics, 2011, 21, 225-230.	1.5	62
50	Effect of Additional Treatments Combined with Conventional Therapies in Pregnant Patients with High-Risk Antiphospholipid Syndrome: A Multicentre Study. Thrombosis and Haemostasis, 2018, 47, 639-646.	3.4	62
51	A novel CISD2 intragenic deletion, optic neuropathy and platelet aggregation defect in Wolfram syndrome type 2. BMC Medical Genetics, 2014, 15, 88.	2.1	59
52	Hemostatic balance in patients with liver cirrhosis: Report of a consensus conference. Digestive and Liver Disease, 2016, 48, 455-467.	0.9	57
53	Hyperhomocysteinaemia in chronic liver diseases: role of disease stage, vitamin status and methylenetetrahydrofolate reductase genetics. Liver International, 2005, 25, 49-56.	3.9	55
54	Development of a Risk Prediction Score for Occult Cancer in Patients With VTE. Chest, 2017, 151, 564-571.	0.8	51

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55	C-Reactive Protein in Offspring Is Associated With the Occurrence of Myocardial Infarction in First-Degree Relatives. Arteriosclerosis, Thrombosis, and Vascular Biology, 2000, 20, 198-203.	2.4	49
56	A new JAK2 gene mutation in patients with polycythemia vera and splanchnic vein thrombosis. Blood, 2007, 110, 2768-2768.	1.4	49
57	Screening for thrombophilia and antithrombotic prophylaxis in pregnancy: Guidelines of the Italian Society for Haemostasis and Thrombosis (SISET). Thrombosis Research, 2009, 124, e19-e25.	1.7	48
58	Venous Thromboembolism in Women Undergoing Assisted Reproductive Technologies: Data from the RIETE Registry. Thrombosis and Haemostasis, 2018, 118, 1962-1968.	3.4	48
59	Determining sulfur-containing amino acids by capillary electrophoresis: A fast novel method for total homocyst(e)ine human plasma. Electrophoresis, 1999, 20, 569-574.	2.4	46
60	Annexin V expression in human placenta is influenced by the carriership of the common haplotype M2. Fertility and Sterility, 2009, 91, 940-942.	1.0	45
61	Lowâ€dose aspirin for in vitro fertilization or intracytoplasmic sperm injection: a systematic review and a metaâ€analysis of the literature. Journal of Thrombosis and Haemostasis, 2012, 10, 2075-2085.	3.8	45
62	Treatment of Right Heart Thrombi Associated with Acute Pulmonary Embolism. American Journal of Medicine, 2017, 130, 588-595.	1.5	45
63	Gain-of-function gene mutations and venous thromboembolism: distinct roles in different clinical settings. Journal of Medical Genetics, 2007, 44, 412-416.	3.2	44
64	Identification of 217 unreported mutations in the F8 gene in a group of 1,410 unselected Italian patients with hemophilia A. Journal of Human Genetics, 2008, 53, 275-284.	2.3	44
65	ABCB1 SNP rs4148738 modulation of apixaban interindividual variability. Thrombosis Research, 2016, 145, 24-26.	1.7	41
66	Low protein Z levels and risk of occurrence of deep vein thrombosis. Journal of Thrombosis and Haemostasis, 2006, 4, 2417-2422.	3.8	40
67	Adverse pregnancy outcomes are associated with multiple maternal thrombophilic factors. European Journal of Obstetrics, Gynecology and Reproductive Biology, 2004, 117, 144-147.	1.1	39
68	Reduced allele specific annexin A5 mRNA levels in placentas carrying the M2/ANXA5 allele. Placenta, 2010, 31, 937-940.	1.5	39
69	Antithrombotic Treatment of Splanchnic Vein Thrombosis: Results of an International Registry. Seminars in Thrombosis and Hemostasis, 2014, 40, 099-105.	2.7	39
70	The risk of first venous thromboembolism during pregnancy and puerperium in double heterozygotes for factor V Leiden and prothrombin G20210A. Journal of Thrombosis and Haemostasis, 2008, 6, 494-498.	3.8	38
71	Iron-dependent erythropoiesis in women with excessive menstrual blood losses and women with normal menses. Annals of Hematology, 2014, 93, 557-563.	1.8	38
72	Pulmonary Thrombosis: A Clinical Pathological Entity Distinct from Pulmonary Embolism?. Seminars in Thrombosis and Hemostasis, 2019, 45, 778-783.	2.7	36

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73	Polymorphisms in factor II and factor VII genes modulate oral anticoagulation with warfarin. Haematologica, 2004, 89, 1510-6.	3.5	36
74	FV HR2 Haplotype as Additional Inherited Risk Factor for Deep Vein Thrombosis in Individuals with a High-Risk Profile. Thrombosis and Haemostasis, 2002, 87, 32-36.	3.4	35
75	Outcomes during anticoagulation in patients with symptomatic vs. incidental splanchnic vein thrombosis. Thrombosis Research, 2018, 164, 69-74.	1.7	35
76	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.	1.7	34
77	Liquid chromatographyâ€“tandem mass spectrometry method as the golden standard for therapeutic drug monitoring in renal transplant. Journal of Pharmaceutical and Biomedical Analysis, 2013, 86, 123-126.	2.8	34
78	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.	1.4	33
79	Homocysteine metabolism in families from southern Italy with neural tube defects: role of genetic and nutritional determinants. Prenatal Diagnosis, 2006, 26, 1-5.	2.3	32
80	Identification of fetal gender in maternal blood is a helpful tool in the prenatal diagnosis of haemophilia. Haemophilia, 2006, 12, 417-422.	2.1	32
81	Reduction of ADAMTS13 Levels Predicts Mortality in SARS-CoV-2 Patients. TH Open, 2020, 04, e203-e206.	1.4	32
82	Early Use of Echocardiography in Patients With Acute Pulmonary Embolism: Findings From the RIETE Registry. Journal of the American Heart Association, 2018, 7, e009042.	3.7	31
83	Plasma Lipoprotein(a) Levels in Subjects Attending a Metabolic Ward. Arteriosclerosis, Thrombosis, and Vascular Biology, 1996, 16, 120-128.	2.4	30
84	Inherited abnormalities of fibrinogen: 10-year clinical experience of an Italian group. Blood Coagulation and Fibrinolysis, 2006, 17, 235-240.	1.0	29
85	Symptomatic Venous Thromboembolism and Thrombophilic Status in Adult Acute Leukemia: A Single-Center Experience of 114 Patients at Diagnosis. Acta Haematologica, 2007, 117, 215-220.	1.4	29
86	Clinical Relevance of Polymorphic Markers of Arterial Thrombosis. Thrombosis and Haemostasis, 1997, 78, 462-466.	3.4	29
87	Age and homocysteine plasma levels are risk factors for thrombotic complications after ovarian stimulation. Human Reproduction, 2004, 19, 1796-1799.	0.9	28
88	Paediatric arterial ischaemic stroke and cerebral sinovenous thrombosis. Thrombosis and Haemostasis, 2015, 113, 1270-1277.	3.4	28
89	Antithrombotic prophylaxis during pregnancy in women with deficiency of natural anticoagulants. Blood Coagulation and Fibrinolysis, 2008, 19, 226-230.	1.0	27
90	Blood supply and transfusion support in southern Italy: findings during the first four weeks of the SARS-CoV-2 pandemic. Blood Transfusion, 2020, 18, 230-232.	0.4	26

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91	Lower birth-weight in neonates of mothers carrying factor V G1691A and factor II A(20210) mutations. Haematologica, 2002, 87, 177-81.	3.5	26
92	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, 477-484.	3.4	25
93	Low molecular weight heparin use during pregnancy and risk of postpartum hemorrhage: a systematic review and meta-analysis. Journal of Maternal-Fetal and Neonatal Medicine, 2019, 32, 1893-1900.	1.5	25
94	Occurrence of factor V Leiden mutation (Arg506Gln) and anticardiolipin antibodies in migraine patients. Neurological Sciences, 2002, 22, 455-458.	1.9	24
95	PAI-1 4G/5G and ACE I/D gene polymorphisms and the occurrence of myocardial infarction in patients on intermittent dialysis. Nephrology Dialysis Transplantation, 2003, 18, 1142-1146.	0.7	24
96	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, 507-514.	2.3	24
97	Efficacy of low molecular weight heparin in patients undergoing inÂvitro fertilization or intracytoplasmic sperm injection. Journal of Thrombosis and Haemostasis, 2011, 9, 2503-2506.	3.8	24
98	Pregnancy-related venous thrombosis: comparison between spontaneous and ART conception in an Italian cohort. BMJ Open, 2015, 5, e008213.	1.9	24
99	The prevention and treatment of venous thromboembolism in pregnancy. Expert Review of Cardiovascular Therapy, 2017, 15, 397-402.	1.5	24
100	Protein Z levels and unexplained fetal losses. Fertility and Sterility, 2004, 82, 982-983.	1.0	23
101	The use of LMWH in pregnancies at risk: new evidence or perception?. Journal of Thrombosis and Haemostasis, 2005, 3, 778-779.	3.8	23
102	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.e1-461.e5.	1.3	23
103	Towards the genetic basis of cerebral venous thrombosisâ€"the BEAST Consortium: a study protocol: TableÂ1. BMJ Open, 2016, 6, e012351.	1.9	23
104	DVT Management andÂOutcome Trends, 2001 to 2014. Chest, 2016, 150, 374-383.	0.8	23
105	Realâ€life Use of Anticoagulants in Venous Thromboembolism With a Focus on Patients With Exclusion Criteria for Direct Oral Anticoagulants. Clinical Pharmacology and Therapeutics, 2018, 103, 684-691.	4.7	23
106	Thrombocytopenia and Mortality Risk in Patients With Atrial Fibrillation: An Analysis From the START Registry. Journal of the American Heart Association, 2019, 8, e012596.	3.7	23
107	Managing anticoagulation in the COVID-19 era between lockdown and reopening phases. Internal and Emergency Medicine, 2020, 15, 783-786.	2.0	23
108	Compound Heterozygosity (554-589 del, C515-T Transition) in the Platelet Glycoprotein IbÎ± Gene in a Patient with a Severe Bleeding Tendency. Thrombosis and Haemostasis, 1999, 81, 486-492.	3.4	22

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109	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, 382-386.	0.7	22
110	The anti-CD20 monoclonal antibody rituximab to treat acquired haemophilia A. Blood Transfusion, 2016, 14, 255-61.	0.4	21
111	Genetic modulation of plasma fibrinogen concentrations: Possible importance of interleukin-6. Journal of Thrombosis and Thrombolysis, 1996, 3, 51-56.	2.1	20
112	Occurrence of the JAK2 V617F mutation in the Buddâ€“Chiari syndrome. Blood Coagulation and Fibrinolysis, 2008, 19, 459-462.	1.0	20
113	Predictors of active cancer thromboembolic outcomes. Thrombosis and Haemostasis, 2017, 117, 1192-1198.	3.4	20
114	Lopinavir/Ritonavir and Darunavir/Cobicistat in Hospitalized COVID-19 Patients: Findings From the Multicenter Italian CORIST Study. Frontiers in Medicine, 2021, 8, 639970.	2.6	20
115	Methylene Tetrahydrofolate Reductase (MTHFR) 677Tâ†'C Mutation and Unexplained Early Pregnancy Loss. Thrombosis and Haemostasis, 1998, 79, 1056-1057.	3.4	20
116	The JAK2 rs12343867 CC genotype frequently occurs in patients with splanchnic venous thrombosis without the JAK2V617F mutation: a retrospective study. Journal of Thrombosis and Haemostasis, 2010, 8, 413-416.	3.8	19
117	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, 104-113.	2.8	19
118	Risk of obstetric and thromboembolic complications in family members of women with previous adverse obstetric outcomes carrying common inherited thombophilias. Journal of Thrombosis and Haemostasis, 2012, 10, 223-228.	3.8	19
119	Validation of PLASMIC score and follow-up data in a cohort of patients with suspected microangiopathies from Southern Italy. Journal of Thrombosis and Thrombolysis, 2018, 46, 174-179.	2.1	19
120	The ISTH bleeding assessment tool as predictor of bleeding events in inherited platelet disorders: Communication from the ISTH SSC Subcommittee on Platelet Physiology. Journal of Thrombosis and Haemostasis, 2021, 19, 1364-1371.	3.8	19
121	Rivaroxaban for the treatment of noncirrhotic splanchnic vein thrombosis: an interventional prospective cohort study. Blood Advances, 2022, 6, 3569-3578.	5.2	19
122	Sex modulation of the occurrence of jak2 v617f mutation in patients with splanchnic venous thrombosis. Thrombosis Research, 2011, 128, 233-236.	1.7	18
123	A novel congenital dysprothrombinemia leading to defective prothrombin maturation. Thrombosis Research, 2014, 134, 1135-1141.	1.7	18
124	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.	2.1	18
125	Detection of the Factor V Leiden Using SSCP. Thrombosis and Haemostasis, 1996, 76, 814-815.	3.4	18
126	Prediction of early mortality in patients with cancer-associated thrombosis in the RIETE Database. International Angiology, 2019, 38, 173-184.	0.9	17

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127	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.	0.9	17
128	A Frameshift Mutation in the Human Fibrinogen AÎ±-chain Gene (AÎ± [499] Ala Frameshift Stop) Leading to Dysfibrinogen San Giovanni Rotondo. Thrombosis and Haemostasis, 2001, 86, 1483-1488.	3.4	16
129	The M2 haplotype in the ANXA5 gene is an independent risk factor for idiopathic small-for-gestational age newborns. Molecular Human Reproduction, 2012, 18, 510-513.	2.8	16
130	Role of Thrombophilia in Adverse Obstetric Outcomes and Their Prevention Using Antithrombotic Therapy. Seminars in Thrombosis and Hemostasis, 2009, 35, 630-643.	2.7	15
131	Stable-isotope dilution LCâ€“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, 3292-3299.	2.3	15
132	Analysis of noncatheter-associated upper extremity deep venous thrombosis from the RIETE registry. Journal of Vascular Surgery: Venous and Lymphatic Disorders, 2017, 5, 18-24.e1.	1.6	15
133	Vitamin K Antagonists After 6 Months of Low-Molecular-Weight Heparin in Cancer Patients with Venous Thromboembolism. American Journal of Medicine, 2018, 131, 430-437.	1.5	15
134	Polymorphism of the Angiotensin-Converting Enzyme Gene in End-Stage Renal Failure Patients. Nephron, 2000, 85, 54-59.	1.8	14
135	Correlation between factors involved in the local haemostasis and angiogenesis in full term human placenta. Thrombosis Research, 2008, 122, 376-382.	1.7	14
136	New TET2 gene mutations in patients with myeloproliferative neoplasms and splanchnic vein thrombosis. Journal of Thrombosis and Haemostasis, 2010, 8, 1142-1144.	3.8	14
137	Management of Patients Taking Oral Anticoagulants Who Need Urgent Surgery for Hip Fracture. Seminars in Thrombosis and Hemostasis, 2019, 45, 164-170.	2.7	14
138	Lack of association between genetic variants in the mannose-binding lectin 2 (MBL2) gene and HPV infection. European Journal of Epidemiology, 2007, 22, 159-162.	5.7	13
139	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, 148-156.	3.8	13
140	Impact of common thrombophilias and JAK2 V617F on pregnancy outcomes in unselected Italian women. Journal of Thrombosis and Haemostasis, 2011, 9, 496-501.	3.8	13
141	Clinical utility of screening for CALR gene exon 9 mutations in patients with splanchnic venous thrombosis. Thrombosis and Haemostasis, 2015, 113, 1381-1382.	3.4	13
142	Very elderly patients with venous thromboembolism on oral anticoagulation with VKAs or DOACs: Results from the prospective multicenter START2-Register Study. Thrombosis Research, 2019, 183, 28-32.	1.7	13
143	Thrombotic microangiopathy, DIC-syndrome and COVID-19: link with pregnancy prothrombotic state. Journal of Maternal-Fetal and Neonatal Medicine, 2022, 35, 2536-2544.	1.5	13
144	Homocysteine levels in amniotic fluid. Thrombosis and Haemostasis, 2006, 95, 625-628.	3.4	12

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145	Pregnancy in a woman with a history of Budd-Chiari syndrome treated by porto-systemic shunt, protein C deficiency and bicornuate uterus. Thrombosis and Haemostasis, 2006, 95, 1033-1034.	3.4	12
146	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, 64-69.	2.3	12
147	The G20210A prothrombin variant and the risk of venous thromboembolism or fetal loss in pregnant women: a family study. Journal of Thrombosis and Haemostasis, 2007, 5, 2193-2196.	3.8	12
148	Venous thrombosis in oral contraceptive users and the presence of the JAK2 V617F mutation. Thrombosis and Haemostasis, 2008, 99, 640-642.	3.4	12
149	Setting up a 2Dâ€LC/MS/MS method for the rapid quantitation of the prostanoid metabolites 6â€oxoâ€PGF_{1Î±} and TXB₂ as markers for hemostasis assessment. Journal of Mass Spectrometry, 2009, 44, 346-352.	1.6	12
150	Clinical histories and molecular characterization of two afibrinogenemic patients: insights into clinical management. Haemophilia, 2012, 18, e16-8.	2.1	12
151	Comparative clinical prognosis of massive and nonâ€massive pulmonary embolism: A registryâ€based cohort study. Journal of Thrombosis and Haemostasis, 2021, 19, 408-416.	3.8	12
152	Identification of Three Novel Mutations in Hereditary Protein S Deficiency. Thrombosis and Haemostasis, 1997, 77, 021-025.	3.4	12
153	Inherited thrombophilia and gestational vascular complications. Best Practice and Research in Clinical Haematology, 2003, 16, 321-332.	1.7	11
154	Markers of haemostasis and angiogenesis in placentae from gestational vascular complications: Impairment of mechanisms involved in maintaining intervillous blood flow. Thrombosis Research, 2010, 125, 267-271.	1.7	11
155	Low-molecular â€“weight heparin in pregnancies after ART -A retrospective study-. Thrombosis Research, 2014, 134, 336-339.	1.7	11
156	Venous thrombosis in afibrinogenemia: a successful use of rivaroxaban. Haemophilia, 2015, 21, e431-3.	2.1	11
157	Assisted reproductive technologies and thrombosis. Thrombosis Research, 2015, 135, S44-S45.	1.7	11
158	Aspirin and heparin in pregnancy. Expert Opinion on Pharmacotherapy, 2015, 16, 1793-1803.	1.8	11
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