Tiscia Giovanni, L

List of Publications by Year
in descending order

56 papers	1,103 citations	430754 18 h-index	434063 31 g-index
57 all docs	57 docs citations	57 times ranked	1378 citing authors

#	Article	IF	Citations
1	Obstetric outcomes in pregnant COVID-19 women: the imbalance of von Willebrand factor and ADAMTS13 axis. BMC Pregnancy and Childbirth, 2022, 22, 142.	0.9	6
2	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.	0.4	17
3	The Prognostic Value of ADAMTS-13 and von Willebrand Factor in COVID-19 Patients: Prospective Evaluation by Care Setting. Diagnostics, 2021, 11, 1648.	1.3	6
4	Focus on Key Issues in Immune Thrombotic Thrombocytopenic Purpura: Italian Experience of Six Centers. Journal of Clinical Medicine, 2021, 10, 5702.	1.0	0
5	Recurrent thrombocytopenia in pregnancy: is it always an obstetric complication?. Internal and Emergency Medicine, 2020, 15, 1271-1274.	1.0	0
6	<scp>PLASMIC</scp> score: Not intended to replace but rather to prompt the <scp>ADAMTS13</scp> testing. Transfusion, 2020, 60, 3070-3072.	0.8	1
7	Thromboelastography Parameters in Italian Pregnant Women: Do Antithrombotic Drugs Change Reference Values?. Journal of Investigative Medicine, 2020, 68, 902-905.	0.7	3
8	Management of Patients Taking Oral Anticoagulants Who Need Urgent Surgery for Hip Fracture. Seminars in Thrombosis and Hemostasis, 2019, 45, 164-170.	1.5	14
9	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.	0.9	0
10	Human Fibrinogen: Molecular and Genetic Aspects of Congenital Disorders. International Journal of Molecular Sciences, 2018, 19, 1597.	1.8	51
11	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.	1.0	19
12	Factor XI gene variants in factor XI-deficient patients of Southern Italy: identification of a novel mutation and genotypeâ€“phenotype relationship. Human Genome Variation, 2017, 4, 17043.	0.4	6
13	Factor VII deficiency: a novel missense variant and genotypeâ€“phenotype correlation in patients from Southern Italy. Human Genome Variation, 2017, 4, 17048.	0.4	6
14	Postpartum haemorrhage in a woman with essential thrombocythemia carrying calreticulin mutation. Blood Coagulation and Fibrinolysis, 2016, 27, 727-728.	0.5	1
15	ABCB1 SNP rs4148738 modulation of apixaban interindividual variability. Thrombosis Research, 2016, 145, 24-26.	0.8	41
16	Modulation of factors involved in placental haemostasis and angiogenesis by low-molecular-weight-heparins. Archives of Gynecology and Obstetrics, 2016, 294, 1323-1329.	0.8	2
17	Pharmacogenetics of dabigatran etexilate interindividual variability. Thrombosis Research, 2016, 144, 1-5.	0.8	69
18	Functional characterization of annexin A5 gene promoter allelic variants. Thrombosis Research, 2016, 144, 93-99.	0.8	7

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19	Clinical utility of screening for CALR gene exon 9 mutations in patients with splanchnic venous thrombosis. Thrombosis and Haemostasis, 2015, 113, 1381-1382.	1.8	13
20	Pregnancy-related venous thrombosis: comparison between spontaneous and ART conception in an Italian cohort. BMJ Open, 2015, 5, e008213.	0.8	24
21	Genetic variations in the annexin A5 gene and the risk ofÂpregnancyâ€related venous thrombosis. Journal of Thrombosis and Haemostasis, 2015, 13, 409-413.	1.9	10
22	Aspirin and heparin in pregnancy. Expert Opinion on Pharmacotherapy, 2015, 16, 1793-1803.	0.9	11
23	Clinical Utility of Antithrombotic Prophylaxis in ART Procedures: An Italian Experience. PLoS ONE, 2014, 9, e97604.	1.1	6
24	Clinical Pregnancies and Live Births in women approaching ART: A follow-up analysis of 157 women after thrombophilia screening. Thrombosis Research, 2014, 133, 168-172.	0.8	9
25	A novel congenital dysprothrombinemia leading to defective prothrombin maturation. Thrombosis Research, 2014, 134, 1135-1141.	0.8	18
26	Low-molecular â€“weight heparin in pregnancies after ART -A retrospective study-. Thrombosis Research, 2014, 134, 336-339.	0.8	11
27	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.	0.8	34
28	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.	1.4	34
29	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.	1.5	28
30	TET2 Mutations in Ph-Negative Myeloproliferative Neoplasms: Identification of Three Novel Mutations and Relationship with Clinical and Laboratory Findings. BioMed Research International, 2013, 2013, 1-5.	0.9	10
31	Structural analysis of protein Z gene variants in patients with foetal losses. Thrombosis and Haemostasis, 2013, 110, 534-542.	1.8	3
32	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.	1.3	16
33	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, 988-993.	1.2	8
34	C0267 Genotype-phenotype relationship in Italian subjects with congenital FXI deficiency. Thrombosis Research, 2012, 130, S140.	0.8	0
35	C0264 Protein Z mutations in patients with fetal losses: Structural analysis. Thrombosis Research, 2012, 130, S146.	0.8	0
36	C0268 Higher risk of idiopathic small for gestational age newborns in Italian women carrying the annexin A5 M2 haplotype. Thrombosis Research, 2012, 130, S195.	0.8	0

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37	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.	1.8	25
38	Clinical histories and molecular characterization of two afibrinogenemic patients: insights into clinical management. Haemophilia, 2012, 18, e16-8.	1.0	12
39	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.	1.9	19
40	Sex modulation of the occurrence of jak2 v617f mutation in patients with splanchnic venous thrombosis. Thrombosis Research, 2011, 128, 233-236.	0.8	18
41	Impact of common thrombophilias and JAK2 V617F on pregnancy outcomes in unselected Italian women. Journal of Thrombosis and Haemostasis, 2011, 9, 496-501.	1.9	13
42	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.	0.7	23
43	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.	1.9	19
44	New TET2 gene mutations in patients with myeloproliferative neoplasms and splanchnic vein thrombosis. Journal of Thrombosis and Haemostasis, 2010, 8, 1142-1144.	1.9	14
45	Factor XI deficiency: two novel mutations in asymptomatic Italian patients. Haemophilia, 2010, 16, 767-770.	1.0	4
46	The haplotype M2 within the ANXA5 gene is independently associated with the occurrence of deep venous thrombosis. Thrombosis and Haemostasis, 2010, 103, 1102-1103.	1.8	8
47	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.	0.8	11
48	Maternal thrombophilia and the risk of recurrence of preeclampsia. American Journal of Obstetrics and Gynecology, 2009, 200, 46.e1-46.e5.	0.7	62
49	Annexin V expression in human placenta is influenced by the carriership of the common haplotype M2. Fertility and Sterility, 2009, 91, 940-942.	0.5	45
50	Haplotype M2 in the annexin A5 (ANXA5) gene and the occurrence of obstetric complications. Thrombosis and Haemostasis, 2009, 102, 309-313.	1.8	63
51	Occurrence of the JAK2 V617F mutation in the Buddâ€“Chiari syndrome. Blood Coagulation and Fibrinolysis, 2008, 19, 459-462.	0.5	20
52	Venous thrombosis in oral contraceptive users and the presence of the JAK2 V617F mutation. Thrombosis and Haemostasis, 2008, 99, 640-642.	1.8	12
53	A new JAK2 gene mutation in patients with polycythemia vera and splanchnic vein thrombosis. Blood, 2007, 110, 2768-2768.	0.6	49
54	The JAK2 V617F mutation frequently occurs in patients with portal and mesenteric venous thrombosis. Journal of Thrombosis and Haemostasis, 2007, 5, 55-61.	1.9	150

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55	The COX-2 G/C â^'765 polymorphism may modulate the occurrence of cerebrovascular ischemia. Blood Coagulation and Fibrinolysis, 2006, 17, 93-96.	0.5	44
56	More on: factor V Leiden and prothrombin G20210A polymorphisms as risk factors for miscarriage during a first-intended pregnancy: the matched case-control 'NOHA first' study. Journal of Thrombosis and Haemostasis, 2006, 4, 709-710.	1.9	6